

# **Biodiversity of leaf beetles (Coleoptera: Chrysomelidae) in a tropical montane rainforest ecosystem assessed with DNA barcoding**



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**Birthe Thormann**  
aus Prüm

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Erstgutachter: Prof. Dr. Johann Wolfgang Wägele

Zweitgutachter: Prof. Dr. Thomas Wagner

Kommissionsmitglied (fachnah): Prof. Dr. Maximilian Weigend

Kommissionsmitglied (fachfremd): Prof. Dr. Christian Borgemeister

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*When the distinguished British biologist J.B.S. Haldane, who found himself in the company of a group of theologians, was asked what one could conclude as to the nature of the Creator from a study of his creation, Haldane is said to have answered "An inordinate fondness for beetles".*

Anecdote



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## Summary

The aim of the present study was the assessment of an unknown tropical insect fauna without traditional taxonomy. For this purpose, the diversity of leaf beetles (Coleoptera: Chrysomelidae) in the montane rainforest of the Reserva Biológica San Francisco (RBSF) and parts of the Podocarpus National Park in southern Ecuador was investigated. Beetles were sampled at three different elevations, '1000 m' (Bombuscaro; 1020–1075 m a.s.l.), '2000 m' (Estación Científica San Francisco – ECSF; 1913–2089 m a.s.l.), and '3000 m' (Cajanuma; 2805–2891 m a.s.l.) with a set of different collection methods. Beetles were mainly sampled with sweep netting, beating, and hand-collection from the lower understorey vegetation of 36 sampling plots (12 per elevation, six of them in the valleys, six on the ridges) following a standardized sampling protocol. A total of 4286 leaf beetles have been collected, 1775 of these (usually one of each morphospecies per sample) were sorted into 515 different morphospecies, DNA barcoded, and assigned to molecular operational taxonomic units (MOTUs).

The study covers aspects of community structure and its changes with increasing elevation. Methodological aspects of rapid biodiversity assessment are evaluated: Different collection methods and morphological and sequence-based methods for species delimitation are compared.

### **General leaf beetle diversity patterns in an Andean mountain forest**

Leaf beetle assemblages showed patterns typical for tropical arthropods: They were species-rich, with few common species but a high percentage of rare species. 1583 specimens were sorted into 473 morphospecies, and for 1334 of them a DNA barcode could be obtained. They belong to 416 morphospecies and were grouped into 459 MOTUs. Species accumulation curves showed no saturation indicating a further increase in species numbers with additional sampling. Species number estimates ranged up to 916 morphospecies (chao2) for the 1583 analysed individuals, and 705 morphospecies, respectively 805 MOTUs for the 1334 barcoded individuals. The higher MOTU number compared to morphospecies number suggests a high level of potential cryptic diversity that was not recognized by the morphospecies approach alone. The leaf beetle community showed an uneven distribution of incidence and abundance with very few common morphospecies (5% found in more than ten samples, 10% represented by more than ten individuals) and a high percentage of uniques (morphospecies found in one single sample; 50% of all morphospecies), respectively singletons (one single individual found; 45% of all morphospecies). The singleton curve did not reach saturation. Most morphospecies were restricted to one single elevational level (91%), indicating a high turnover of communities with elevation. This pattern was even more apparent for MOTUs (96%) and haplotypes (99%). More than half of the morphospecies belonged to Alticinae (53%), 21% were Galerucinae, 14% Eumolpinae, 5% Hispinae, and 4% Cassidinae. Criocerinae, Chrysomelinae, Lamprosomatinae, and Cryptocephalinae together accounted for 3% of all morpho-

species. Rank order remained the same when number of individuals was considered. Composition of the subgroups changed slightly with elevation.

### **Diversity patterns along an elevational gradient inferred with DNA barcode data**

Leaf beetle assemblages from the 36 study plots were sampled and differences between the three elevations and the two microhabitats (forest on ridges and in valleys) were analysed based on DNA barcode data. The importance of small-scale topography for elevational diversity patterns was evaluated: It was tested whether elevational diversity differs between ridge and valley forests and if the species turnover between and within habitats varies with elevation and changes patterns of elevational diversity when scaling up from the local (sampling plot) to the regional (elevational belt) level. MOTUs were determined using PTP modelling and data was analysed using permutational MANOVA analysis and ordinary linear models.

When study sites of both habitats were pooled, local leaf beetle diversity showed a clear mid-elevational peak pattern. However, only leaf beetle diversity in ridge forests peaked at mid-elevations, while the diversity in valley forests was similarly high at 1000 and 2000 m a.s.l. and declined at highest elevations. When scaling up to the regional scale, levels of diversity were generally similar at the two lower elevations and declined at 3000 m a.s.l. The scale-dependent shift in diversity patterns was caused by a higher turnover of species communities between and within habitats at lower than at mid-elevations, suggesting more specialized herbivore communities in the more productive lower elevations. The study underscores the importance of topography and spatial scale for the inference of diversity patterns. Changes in ecosystem productivity but also area and temperature with elevation might also influence the genetic diversity within species, however, levels of genetic diversity (haplotype diversity per MOTU) did not differ among elevational levels. Biodiversity patterns along the elevational gradient were revealed by MOTUs and morphospecies in the same way.

### **Comparison of morphospecies sorting and DNA barcoding**

1475 barcoded individuals were assigned to MOTUs and the results were compared with the morphospecies sorting. The barcode approach estimated 10% higher species numbers (448 morphospecies, 493 MOTUs). This was caused by a higher number of splittings than lumpings of morphospecies. The similar numbers of morphospecies and MOTUs arose partly due to the fact that splittings and lumpings compensated one another. However, the number of perfect matches was comparatively low: 63% of all morphospecies corresponded exactly with one MOTU. Most lumpings united individuals of two morphospecies in one MOTU (76%), in some cases, individuals of up to five morphospecies (4%) were lumped. Similarly, most splittings divided a morphospecies in two networks (69%), only once a morphospecies was split into six MOTUs (1%). The subgroups most challenging for morphospecies sorting were Galerucinae and especially Alticinae. Difficulties most probably arose due to the large number of specimens and species.

DNA barcoding showed to be a valuable tool in cases where morphospecies sorting is exacerbated by pronounced intraspecific variation in colour, shape, or size, and may reveal cryptic diversity. Especially in species that are small and/or lack conspicuous external characters barcoding is a useful tool to complement morphospecies sorting. Particularly in large, specimen- and species-rich data sets DNA barcoding can facilitate morphospecies sorting and can result into a more accurate species delimitation.

### **Influence of different species delimitation methods on species richness estimates**

For a subset of 674 barcoded specimens, a set of four different DNA-based species delimitation methods and their influence on species richness estimates were compared. Distance-based clustering, statistical parsimony analysis, GMYC-, and PTP modelling led to highly similar results. The reason probably lies within the structure of the underlying data set: It is geographically restricted and undersampled with a high proportion of singletons what turns it insensitive against differences in species delimitation methods. Several cases of splittings and lumpings led to discrepancies between morphospecies and MOTU assignment and generally MOTU numbers were ~8% higher than morphospecies numbers.

Morphospecies sorting and DNA barcoding allow similar conclusions on leaf beetle diversity: The leaf beetle fauna is species-rich with a strong turnover among elevations. Most morphospecies were found only at a single elevational level, also when singletons and doubletons have been excluded. This pattern was even more visible for MOTUs and haplotypes. The high turnover between leaf beetle communities at the different elevations is also visible in the species accumulation curves: If to the specimens of one elevation the specimens of a second elevation were added, the curves showed once more a further increase.

### **Comparison of sampling methods**

Within the present study a total of 1174 samples were taken. They varied considerably in size and effort as different sampling methods were used. The focus was on standardized sampling with sweep netting, beating, and hand-collection on the sampling plots. Malaise trapping, light trapping, and additional hand-collection completed the sampling.

In sweep netting-, beating-, hand-collection-, and light trap samples on average only few individuals and morphospecies were caught per single sample (less than five). In contrast, the Malaise traps were highly efficient on a per sample basis: They yielded a mean of 31 individuals and 15 morphospecies per sample. Collection efficiency for certain subgroups slightly differed between the different methods. Even after 298.5 sampling hours the species accumulation curve of the standardized plot samples showed no saturation indicating that a further increase of morphospecies number is expected with further sampling.



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# General introduction

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## 1.1 Background of the study

We are right in the middle of an anthropogenic mass extinction with rates of decline in biodiversity comparable to previous mass extinction events in the fossil record (Barnosky et al., 2011; Dirzo et al., 2014; Pimm et al., 1995). This rapid loss in biological diversity has been termed the global biodiversity crisis, and at least since the signature of the Convention on Biological Diversity (CBD) at the United Nations Conference on Environment and Development in Rio de Janeiro in 1992, the problem has been acknowledged by politics and entered public awareness (<https://www.iucn.org>). Biodiversity is threatened mainly by habitat loss and degradation, but also invasive alien species, over-exploitation of natural resources, pollution and diseases, and climate change (Bradshaw et al. (2009); Primack (2014); <https://www.iucn.org>). Human activity is responsible for most of those perils.

Most biodiversity concentrates in tropical habitats (Bradshaw et al., 2009; Dirzo and Raven, 2003). Especially tropical rainforests are among the most species-rich and ecologically complex ecosystems: Although they cover only seven percent of the Earth's surface, it is estimated that they harbour more than half of all existing species on the planet (Bradshaw et al., 2009; Primack, 2014; Wilson, 1988). Threats to tropical forests are manifold. The rapidly progressing deforestation with fragmentation and overexploitation entails a string of adverse consequences that interact and create negative synergisms (Laurance, 1999; Laurance and Peres, 2006).

Whereas the gravity of habitat destruction and biodiversity loss as well as the urgent need for action are unmistakable, we are still not even able to specify the global number of species. It lies probably within the range of five to 15 million (Chapman, 2009; May, 2000, 2010; Mora et al., 2011; Wilson, 2003). The best known and most intensively studied components of tropical rainforests are mammals, birds, and higher plants that make up only a small fraction, probably less than one percent of the total number of species (Corlett and Primack, 2011). In contrast, most of those groups that account for the majority of biodiversity remain clearly understudied. Invertebrates, and especially insects, are the dominant animals of the rainforest contributing the majority of species, individuals, and biomass (Corlett and Primack, 2011; Primack, 2014). Unfortunately, the so-called taxonomic impediment is especially severe in those groups: Most species cannot be identified and millions are still undescribed due to a shortage of trained taxonomists and curators and a lack of simple-to-use identification guides (<http://www.cbd.int>).

To alleviate the problem of rapid biodiversity-loss with the concurrent gaps in our

taxonomic knowledge, an acceleration of biodiversity assessment is necessary. One possible way is the integration of DNA barcoding for exploring unknown biodiversity. On the one hand it can help to discover and describe species, a task that has never been more urgent (Frézal and Leblois, 2008; Hebert et al., 2003a; Scheffers et al., 2012). On the other hand it can reveal patterns of biodiversity and community ecology while the long lasting process of thorough taxonomic identification and formal description of new species is postponed (Smith et al., 2005; Tänzler et al., 2012). Initially developed as a global species identification system, during the last decade it has proven to be very useful in various fields of research and has also been used successfully in beetle communities (Baselga et al., 2013; Papadopoulou et al., 2013; Tänzler et al., 2012).

In the present study, DNA barcoding is used to investigate the unknown leaf beetle fauna (Coleoptera: Chrysomelidae) in a tropical montane rainforest in Ecuador. DNA barcode data is used along with a morphospecies approach. It is tested, how the methods agree and which conclusions they allow about the diversity and change of leaf beetle communities along an elevational gradient. For that purpose Neotropical Chrysomelidae are particularly attractive study organisms as they are megadiverse and hitherto taxonomically considerably understudied. As an integral component of the herbivorous insect fauna in rainforests they have important functions within ecosystems and are of great relevance for ecosystems' diversity (Andrew and Hughes, 2004; Basset, 2001; Coley and Barone, 1996; Janzen, 1970; Price, 2002; Wagner, 2000).

The investigated region is located in southern Ecuador in the Tropical Andes, a designated biodiversity hotspot for various taxa (Brummitt and Lughadha, 2003; Myers et al., 2000). In tropical mountains, the peaking species richness at low latitudes is enhanced by a high species turnover along elevational gradients (Brühl et al., 1999; Gaston, 2000; Smith et al., 2014). This leads to the exceptional species numbers of tropical montane rainforests that can even exceed those of lowland rainforests (Beck and Kottke, 2008; Rodriguez-Castaneda et al., 2010). With some exceptions (e.g. Brehm and Fiedler (2003, 2004); Brehm et al. (2003a,b); Escobar et al. (2007, 2005, 2006); Hilt et al. (2006, 2007); Janzen et al. (1976); Moret (2009); Olson (1994)), the insect diversity of the Tropical Andes is still understudied and comparatively little is known about diversity patterns in Andean montane forests (for an overview see Larsen et al. (2011)). Montane rainforests face many of the same threats as other tropical forests, however, especially cloud forests are particularly susceptible to climate change due to their unique ecology and their location on mountain slopes (Bubb et al., 2004; Hamilton et al., 1995). Ecuador is a megadiverse country: On a comparatively small area, it harbours an outstanding variety of habitats along pronounced elevational and wet–dry gradients. High beta-diversity along these gradients favours an enormous biological diversity (Brehm et al., 2008a; Dangles, 2009). It shelters one of the most species-rich but also most endangered insect faunas on Earth (Dangles, 2009). Ecuador's dense population (55 inhabitants/km<sup>2</sup>) puts strong pressure on its natural ecosystems (Dangles, 2009). The country suffers the highest annual deforestation rate (-1.9%) in South America (FAO, 2010), mainly

caused by conversion of forest into agropastoral land (Mosandl et al., 2008).

## 1.2 DNA barcoding

DNA barcoding as a global identification system based on a standard molecular method was proposed by Hebert et al. in 2003 in order to accelerate species discovery and identification, and to overcome the limitations of morphological identification (Hebert et al., 2003a). However, the term 'DNA barcodes' was already used by Arnot et al. (1993) and use of DNA sequence differences for identification and discrimination of species has been established for many years for morphologically scarcely identifiable groups such as viruses, bacteria, protists, or fungi (Allander et al., 2001; Bruns et al., 1991; Hamels et al., 2001; Nanney, 1982; Pace, 1997).

Also for higher organisms a DNA based practice approach to taxa recognition is highly expedient and beneficial (Savolainen et al., 2005). It can help to lighten the taxonomic impediment, the lack of taxonomic expertise to effectively identify and describe the remaining biodiversity on Earth. This problem is especially pressing in the light of the rapid biodiversity loss and notably severe with highly diverse arthropod taxa (Cardoso et al. (2011); Evenhuis (2007); CBD Guide to the Global Taxonomy Initiative, <http://www.cbd.int>). The traditional means of studying biodiversity depend on expert knowledge from taxonomists with years of education and training. This knowledge is limited to certain taxa and furthermore, the identification of species is time intensive (Harris and Bellino, 2013). The focus of taxonomic expertise is biased towards vertebrates, flowering plants, or certain insect taxa, whereas in contrast many important groups as e.g. nematodes, mites, or diatoms, are neglected (Tautz et al., 2003). Several authors claimed that traditional taxonomy will not be able to cover all requested identification of biodiversity, but that new approaches are needed to modernize taxonomy (Frézal and Leblois, 2008; Godfray, 2002; Hebert et al., 2003a; Stoeckle, 2003).

DNA barcoding represents the idea of a unique DNA sequence for each species in analogy to industrial 'barcodes', universal product codes which can be used to identify retail products (Hebert et al., 2003a,b; Savolainen et al., 2005). A fragment of the mitochondrial gene cytochrome *c* oxidase I (COI) established as a standard marker for animal species identification (more information about COI as barcode marker is given in Chapter 2.4). This fragment can be amplified with universal markers across a broad range of species (Folmer et al., 1994; Hebert et al., 2003a). The principle of DNA barcoding has been extended to other organisms like fungi (Begerow et al., 2010; Schoch et al., 2012; Seifert, 2009; Seifert et al., 2007) and plants (Chase et al., 2007; Dunning and Savolainen, 2010; Kress and Erickson, 2007; Kress et al., 2005; Pennisi, 2007; Rubinoff et al., 2006) where the search for a universal barcode marker turned out to be difficult.

In 2004, the Consortium for the Barcode of Life (CBOL, <http://www.barcodeoflife.org>) was founded as an international initiative for promoting the development of DNA barcoding as a global standard for species identification with numerous

member organizations such as natural history museums, zoos, herbaria, botanical gardens, university departments, as well as private companies and governmental organizations. Major CBOL projects are e.g. the All Birds Barcoding Initiative (ABBI), the Bee Barcode of Life Initiative (Bee-BOL), the Mosquito Barcode Initiative (MBI), or the International Network for Barcoding Invasive and Pest Species (INBIPS).

The objective of the international Barcode of Life project (iBOL, <http://www.barcodeoflife.org>) is the creation of large numbers of barcodes with a construction of a barcode reference library and the development of instruments and informatics tools for application. The library, Barcode of Life Data Systems (BOLD, <http://www.boldsystems.org>), is also a public workbench for researchers who can assemble, test, and analyse their data in BOLD. iBOL has members in 25 nations and different working groups are devoted to certain taxonomic groups or habitat types (e.g. vertebrates, land plants, fungi, marine bio-surveillance, polar life).

DNA barcoding claims being a rapid and cost-efficient method that moreover is potentially applicable by everyone irrespective of their background training (Hebert and Gregory, 2005; Stoeckle, 2003; Stoeckle et al., 2003). It is also supposed to help in cases where phenotypic plasticity or intraspecific variability impede morphological identification and to facilitate discovery of cryptic diversity (Hebert et al., 2003a). It is applicable to all life forms (whereas keys are often only for one particular life stage or gender) as well as processed or parts of organisms (Hebert et al., 2003a; Stoeckle, 2003; Stoeckle et al., 2003).

Indeed, a vast number of studies within the last decade affirmed the value of DNA barcoding across a broad range of possible applications: It has been shown to be especially useful in difficult groups where morphological traits do not clearly discriminate species. These include very small organisms where body-size precludes visual identification (meio- and micro-fauna, zooplankton), species only distinguishable by subtle or geographically variable morphological characters, as well as species with polymorphic life cycles and/or pronounced phenotypic plasticity (Blaxter et al., 2004; Bucklin et al., 2007; Decaëns et al., 2013; Frézal and Leblois, 2008; Plaisance et al., 2009; Vences et al., 2005). It has also helped in studying cryptic diversity (Hebert et al., 2004; Smith et al., 2006) and has been successfully used for identifying immature stages (e.g. eggs, larvae, seedlings) and assort them to adults (Ahrens et al., 2007; Janzen et al., 2005; Vences et al., 2005). Barcoding of gut contents can give information about an organism's diet (Blankenship and Yayanos, 2005; Zeale et al., 2010). This variety of applications makes barcoding a useful tool in ecology, forensics, and biosecurity (Armstrong and Ball, 2005; Besansky et al., 2003; Chen et al., 2004; Joly et al., 2014; Valentini et al., 2008; Wells and Sperling, 2001). The identification of damaged or processed organisms or fragments is important for food safety and consumer protection as well as for conservation issues as it can help preventing poaching and illegal trade of endangered species (Ardura et al., 2010; Dalton and Kotze, 2011; Eaton et al., 2010; Galimberti et al., 2013; Wong and Hanner, 2008; Yan et al., 2013). It has successfully supported biodiversity inventories and can substitute or complement taxonomically valid species or morphospecies in

community ecology studies of unknown faunas (Janzen et al., 2005; Tänzler et al., 2012).

Despite a broad acceptance and utilization of DNA barcoding, since its beginnings it has aroused criticism as well (DeSalle et al., 2005; Moritz and Cicero, 2004; Will et al., 2005; Will and Rubinoff, 2004). On the one hand there is criticism on the part of taxonomists that are apprehensive of competition or being booted out by barcoding (Ebach and Holdrege, 2005; Lipscomb et al., 2003; Wheeler, 2004). Another point where criticism tackles is the premise that genetic variation among species is normally lower than between species (Hebert and Gregory, 2005; Hebert et al., 2003b). That phenomenon is called the 'barcoding gap', however, in practice there may be overlap between inter- and intraspecific distances, or the barcoding gap may be artificially exaggerated by inappropriate sampling: It has been argued that DNA barcoding fails when a comprehensive sampling exists, i.e. when the whole genetic variance of a species across a broad geographic range is assessed and many closely related species are included. Intra- and interspecific distance can overlap, on the one hand because the intraspecific distances are larger than when only analysing a narrow cut-out of all intraspecific distances, and on the other hand increasing geographical scale of sampling decreases the interspecific divergence due to encountering more closely related, allopatrically distributed species in a geographically expanding data set (Bergsten et al., 2012; Meyer and Paulay, 2005; Wiemers and Fiedler, 2007). In contrast, the barcoding gap is more pronounced on a local scale and for data sets lacking large numbers of closely related species (Moritz and Cicero, 2004).

Certain aspects concerning the use of COI as single marker that require cautiousness are explained in Chapter 2.4.

Methodological approaches that are demarcated from DNA barcoding *sensu strictu*, but still closely tied to the concept of DNA barcoding and partly overlapping are DNA taxonomy, reverse taxonomy, and integrative taxonomy:

*DNA taxonomy* sets the DNA based identification in the focus of taxonomy with DNA being the scaffold of a taxonomic reference system (Tautz et al., 2003). In contrast to DNA barcoding that can be understood as a means of identifying *a priori* entities by sequence similarity, DNA taxonomy concerns the circumscription and delineation of species using evolutionary species concepts (Vogler and Monaghan, 2006).

*Reverse taxonomy* is a sequence-based approach to access unknown diversity. Taxa are at first only identified via their signature sequences that can be re-identified unequivocally in future collections, but are not yet morphologically analysed and formally described (Markmann and Tautz, 2005). Reverse taxonomy can be based on COI sequences but often other markers have been used (Markmann and Tautz, 2005; Randrianiaina et al., 2011).

*Integrative taxonomy* aims at delimiting species boundaries from multiple and complementary perspectives. Traditional morphology-based taxonomy is combined with e.g. molecular, behavioural, developmental, or ecological data (Dayrat, 2005). In

many studies COI is included, often in combination with other markers (Damm et al., 2010; Gibbs, 2009; Heethoff et al., 2011; Mengual et al., 2006; Roe and Sperling, 2007).

In the context of DNA barcoding and DNA taxonomy, often the *MOTU concept* appears (Blaxter, 2004; Floyd et al., 2002). MOTU signifies 'molecular operational taxonomic unit', i.e. a group of specimens defined by sequence identity: If two specimens yield sequences that are identical within some defined cut-off, they are assigned to the same MOTU (Blaxter, 2004). In this study, the term MOTU is used in a broader sense meaning a group of specimens that is delimited by any molecular species delimitation method (e.g. a GMYC-, PTP-, distance-cluster, or a haplotype network). Different methods of molecular species delimitation are explained in Chapters 2.5 and 6.

## 1.3 Chrysomelidae LATREILLE, 1802

### 1.3.1 Biology and ecology

#### General biology and ecology

Chrysomelidae (leaf beetles; Coleoptera: Polyphaga: Cucujiformia: Chrysomeloidea) belong with Cerambycidae (longhorn beetles) to Chrysomeloidea that together with the Curculionoidea (weevils) make up the megadiverse lineage of 'Phytophaga' that constitute about 40% of all known beetle species (Farrell, 1998; Gómez-Zurita et al., 2007; Riley et al., 2002). With over 37,000 described species and more than 2,000 genera Chrysomelidae are one of the largest beetle families (Jolivet et al., 1988). The total number of existing leaf beetle species is probably 60,000 or higher (Jolivet, 1988; Reid, 1995). Chrysomelidae have a worldwide distribution (except arctic regions) with the by far greatest diversity found in the tropics (Riley et al., 2002).

Leaf beetles have a highly variable body shape from elongate-cylindric to oval-convex or depressed (Riley et al., 2002). Size varies from less than one to ~27 mm (Jolivet and Petitpierre, 1981; Jolivet et al., 1988). They show various colours, commonly bright or metallic, often dorsally bicoloured and formed into distinctive patterns (Riley et al., 2002).

Chrysomelidae bear five tarsomeres and are characterized by a bilobed third tarsomere that hides the reduced fourth tarsomere (pseudotetramerous condition). Also typical are large ventral tarsal pads with adhesive hairs that likely aid attachment and locomotion on plants (Fig. 1.1). Antennae are generally short to medium-length with usually eleven antennomeres (Jolivet et al., 1988; Riley et al., 2002).

Chrysomelidae are phytophagous during larval and adult stage (Jolivet and Petitpierre, 1981). Adult chrysomelids usually feed on green parts of plants. Some groups secondarily feed on pollen, flowers, roots, seeds, and ant nests debris. Also leaf beetle larvae have a variety of feeding habits: Besides feeding on leaves or subterranean parts of plants there are also leaf-miners or consumers of dead plant material (Riley et al., 2002).

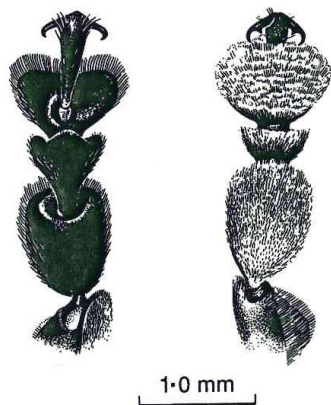


Figure 1.1: Dorsal and ventral view of the tarsus of a typical leaf beetle (Chrysomelinae) with pilose tarsal pads (Lawrence and Britton, 1994).

Traditionally, a restriction of tropical rainforest herbivorous insects to a narrow set of host plants is assumed (Coley and Barone, 1996; Erwin, 1982) and indeed a lot of species seem to be specialists for a certain species, genus, or family (Barone, 1998; Janzen, 1988). However, there is growing evidence that many tropical insects are less specialized than was previously thought (Basset, 1992; Descampe et al., 2008; Novotny et al., 2002b). Assumptions about host plant range that are derived from field observations are often skewed towards specialization as often only a fraction of a (geographical) broader range of host plants is observed (Descampe et al., 2008). In contrast, insects often only sit on plants for resting, shelter, sun-basking, or sexual display, but without feeding on them (Descampe et al., 2008; Moran and Southwood, 1982; Novotny and Basset, 2000).

For leaf beetles, food plant records are estimated to exist for ~30% of described species, especially well studied and important chrysomeline, alticine, and galerucine genera (Jolivet, 1988; Jolivet and Hawkeswood, 1995). A broad variety of plant families are selected by Chrysomelidae (several are discussed in detail by Jolivet and Hawkeswood (1995)). They mainly belong to angiosperms, both monocotyledons ( $\geq 28$  families) and dicotyledons ( $\geq 120$  families) with most leaf beetle subfamilies preferring dicotyledons. However, also gymnosperms and even pteridophytes are chosen, mostly by Alticinae (Jolivet and Hawkeswood, 1995; Riley et al., 2002). There is scarce evidence for leaf beetles (only alticines) feeding on bryophytes (Jolivet and Hawkeswood, 1995; Konstantinov et al., 2013; Konstantinov and Chamorro-Lacayo, 2006). Among the dicotyledon families that are most often recorded in the literature as host plants are Asteraceae, Convolvulaceae, Brassicaceae, Cucurbitaceae, Lamiaceae and Verbenaceae, Fabaceae (and other legumes), Rosaceae, and Solanaceae (Jolivet and Hawkeswood, 1995).

Some Chrysomelidae feed on myrmecophilous plants and developed protective adaptations against the ants e.g. avoiding contact with the ants, mining into the leaf, toxicity, reflex bleeding, stomach discharges, or larval cycloalexia (circular defence; Jolivet et al. (1990)) (Jolivet and Hawkeswood, 1995). Generally, many defensive strategies have evolved within Chrysomelidae, probably in adaptation to their life on the leaf surface where they are prone to numerous biotic and abiotic perils

(predation, desiccation, plant chemical and physical defences; Vencl et al. (2004)). These defensive mechanisms range from crypsis (e.g. cassidines), mechanical devices (e.g. spines of hispines), and sudden escape (Alticinae) to chemical defence in many brightly coloured, aposematic species (Pasteels et al., 1988). An interesting behaviour in leaf beetles is the defensive use of faecal material. Adults and immature stages of species in several subfamilies developed elaborate faecal constructions that are used as camouflage, clubs, and protective covers (Chaboo, 2007; Chaboo et al., 2008; Furth, 1982a, 2004; Müller and Hilker, 2004; Riley et al., 2002). Larval egg cases from faecal material may also serve as protection against desiccation (Furth, 1982a, 2004).

Chrysomelidae have a broad range of reproduction ways. Whereas laying of eggs is most common there is also viviparity and intermediate ways with laying eggs containing more or less developed embryos (Jolivet and Petitpierre, 1981). In some species of Cassidinae and Chrysomelinae a kind of parental care has been observed where females sit on top of the eggs, larvae, and pupae in order to protect them (Buzzi, 1988; Chaboo, 2007; Windsor and Choe, 1994).

### **Significance for humans: Chrysomelidae as pests and biological control agents**

Many leaf beetles are serious pests of agricultural crops and forests, especially in the subfamilies of Criocerinae, Eumolpinae, Galerucinae, Alticinae, Hispinae, and Chrysomelinae (Jolivet et al., 1988). One of the most well-known and destructive agricultural insect pests worldwide is the Colorado potato beetle (*Leptinotarsa decemlineata* SAY, Chrysomelinae) (Alyokhin, 2009; Bishop and Grafius, 1996; Hare, 1990; Weber, 2003). Originating in Mexico, populations quickly spread throughout North America from the 1800's and throughout Eurasia since 1922 (Alyokhin, 2009; Bishop and Grafius, 1996). Adults and larvae of the Colorado potato beetle severely damage potato crops by defoliation (Alyokhin, 2009; Bishop and Grafius, 1996). The species has an impressive ability to evolve insecticide resistance (Alyokhin et al., 2008; Forgash, 1985).

Many significant pests of North American agriculture belong to the diabroticine group of Galerucinae. *Diabrotica virgifera virgifera* LÉCONTE (western corn rootworm), *Diabrotica barberi* SMITH & LAWRENCE (northern corn rootworm), and *Diabrotica undecimpunctata howardi* BARBER (southern corn rootworm) are major pests of cultivated corn, *Zea mays* L., with their larvae feeding on the roots (Ciosi et al., 2008; Roehrdanz et al., 2003). Corn rootworms also attack additional crops as cucurbits and legumes (Krysan, 1986; Metcalf, 1986). They are native to North America but the highly invasive *D. virgifera* is also a serious threat of European agriculture (Ciosi et al., 2008; Gray et al., 2009; Miller et al., 2005; Moeser and Vidal, 2004). Other diabroticine pests are the Mexican corn rootworm (*D. virgifera zeae* KRYSAN & SMITH), the banded cucumber beetle (*D. adelpha* HAROLD) and the western spotted cucumber beetle (*D. undecimpunctata undecimpunctata* MANNERHEIM), and the bean leaf beetle (*Cerotoma trifurcata* FORSTER and other *Cerotoma* species), feeding on a variety of leguminous host plants, especially soybean (Kogan et al., 1980;



Krysan, 1986; Lam and Pedigo, 2004). Among Alticinae there are to name several pests in the genus *Epitrix*: the potato or tuber flea beetles (*E. cucumeris* HARRIS, *E. similaris* GENTNER, *E. tuberis* GENTNER) that attack potato tubers and foliage (Gentner 1944), and the tobacco flea beetle *E. hirtipennis* MELSHEIMER. Other serious pests, especially of crucifer field crops, are found within the genus *Phyllotreta*, e.g. the cabbage or crucifer flea beetle *P. cruciferae* GOEZE, and the striped flea beetle *P. striolata* FABRICIUS. Especially among Hispinae, there are found important pests of palm trees (Mariau, 2004). Numerous species live off oil and coconut palms (Mariau, 2004). The coconut hispine beetle or coconut leaf beetle (*Bron-tispa longissima* GESTRO) is a serious pest of palms, especially *Cocos nucifera* L. It has enormously expanded and is listed in the Global Invasive Species Database (2010) (Takano et al., 2011, 2012). The hispine *Coelaenomenodera lameensis* BERTI & MARIAU, the most serious oil palm pest throughout West Africa has caused severe defoliation over wide areas of oil palm distribution (Mariau, 2004). The tortoise beetle *Paropsis atomaria* OLIVIER represents an emergent pest of *Eucalyptus* plantations in Australia (Schutze et al., 2006). Economically important Criocerinae that damage cereals are the cereal leaf beetle *Oulema melanopus* L., the rice leaf beetle *O. oryzae* KUWAYAMA, and the cereal pest *O. gallaeciana* VON HEYDEN (Wellso and Hoxie, 1988). Among Eumolpinae, there are some cacao pests in Brasil (Ferronato, 1988) and several species attacking sweetpotato, e.g. *Typophorus nigritus viridicyaneus* CROTCH, *Colasposoma dauricum* MANNERHEIM, and *C. sellatum* BALY (Alaijos and Lee, 2005; Jackson et al., 2003; Reid and Storey, 1993). Eumolpinae of the genus *Eucolaspis* cause economic loss on apple orchards (Doddala et al., 2013) and the eumolpine *Trichiona nigra* JACOBY has been recently reported to cause severe feeding damage on pomegranate in India (Jayanthi and Verghese, 2014).

On the other hand, benefit can be derived from the chrysomelids' herbivory and host-specificity: Several leaf beetle species are used for biological control of imported noxious weeds that can cause enormous ecological and economical damage (Jolivet et al., 1988).

Several invasive species of Asian saltcedars (*Tamarix* sp.) cause great damage of riparian ecosystems in the western United States. The galerucine *Diorhabda elongata* BRULLÉ *deserticola* CHEN from Asia has been introduced as biological control agent (DeLoach et al., 2003; Lewis et al., 2003). The common ragweed (*Ambrosia artemisiifolia* L., Asteraceae) is a harmful agricultural weed that is native in North America. Its pollen are highly allergenic. It has invaded Europe as contaminant of agricultural products and spread first slowly but booming since the 1990's facilitated by socio-economic factors (Kiss, 2007). *Ophraella communis* LESAGE, a galerucine from North America, is the most promising biocontrol agent of ragweed (Kiss, 2007). Several species of European *Aphthona* flea-beetles (Alticinae) have been introduced into North America to control leafy spurge (*Euphorbia esula* L.), a weed introduced from Eurasia that is very persistent and invades a variety of habitats (Gassmann et al., 1996; Kirby et al., 2000; Lym and Nelson, 2000). In an attempt to control *Lantana camara* L. (Verbenaceae) (and allied *Lantana* species), an aggressive, vig-

orously growing weed that has become a plague over most of the tropics (Sharma et al., 2005), Hispinae (e.g. *Octotoma scabripennis* GUÉRIN-MÉNEVILLE, *Uroplata girardi* PIC) have been introduced into several regions of the world (Broughton, 2001; Cilliers and Naser, 1991; Harley, 1969). *Chrysolina quadrigemina* SUFFRIAN (Chrysomelinae; released in California to control Klamath weed *Hypericum perforatum* L., Clusiaceae), *Uroplata girardi* PIC (a hispine leaf-miner supposed to control *Lantana camara* L. in Australia), and *Zygogramma bicolorata* PALLISTER (released in India for control of the parthenium weed *Parthenium hysterophorus* L., Astera-ceae) belong to the very few recorded examples of biocontrol agents attacking also non-target plant species (McFadyen, 1998).

### Evolution and fossil history

Despite an abundance of available material, the fossil history of Chrysomelidae is relatively poorly documented (Chaboo and Engel, 2009; Santiago-Blay, 1994). The great species diversity of leaf beetles and other phytophagous insects is commonly ascribed to their co-evolution with the rapidly radiating land plants in the Tertiary (Ehrlich and Raven, 1964; Farrell, 1998; Farrell et al., 1992). The phylogeny of Chrysomelidae is thought to reflect that of major lineages of angiosperms i.e. the available host plant lineages at that time (contemporaneous lineage diversification). The most basal lineages of Chrysomelidae are supposed to be associated with primitive cycads and conifers followed by a large diversification of lineages on di- and monocotyledonous angiosperms (Farrell, 1998; Farrell and Sequeira, 2004; McKenna and Farrell, 2006). Based on these assumptions, the origin of Chrysomelidae seems to be early- to mid-cretaceous leading to a discrepancy between the molecular calibrations and the fossil record (Gómez-Zurita et al. (2007) and references therein). The attribution of Jurassic fossils to Chrysomelidae (Santiago-Blay, 1994) is doubtful, and also in the Cretaceous chrysomelid fossils are essentially absent (Chaboo, 2007; Gómez-Zurita et al., 2007). Most appear only in the Eocene (34–56 Ma), representing most major subfamilies. An exception is a Canadian Mesozoic fossil dated to 72 Ma and identified as a primitive chrysomelid probably representing an early lineage which pre-dates the diversification of major extant subfamilies. The oldest clearly identifiable record is *Donacia wightoni* ASKEVOLD from the Canadian Palaeocene (~58 Ma; Askevold (1990)) (for an overview see Gómez-Zurita et al. (2007) and references therein). Feeding damage on fossil leaves that has been ascribed to hispines is dated  $\geq 60$  Ma and marks the beginning of the hispine/Zingiberales association (Wilf et al., 2000).

An alternative to the co-evolution hypothesis is a time-displaced diversification of the herbivores with radiation of herbivores being based on a pre-existing diversity of host plants (sequential evolution; Jermy (1976)). This scenario is supported by a study proposing a later origin of Chrysomelidae (end of the Cretaceous, 74–79 Ma) than the previous studies suggest and consequently a basal chrysomelid diversification substantially younger than the radiation of their hosts (Gómez-Zurita et al., 2007). A time lag between host radiation and the colonization by herbivores has been shown for several insects (Lopez-Vaamonde et al., 2006; McKenna et al., 2009).

### Leaf beetle biology and ecology – State of the art

The qualities that distinguish Chrysomelidae as interesting study organisms are their species richness as well as their herbivorous mode of life. Herbivorous insects are an extremely species-rich feeding guild with important functions in ecosystems and great relevance for ecosystems' diversity (Coley and Barone, 1996; Janzen, 1970; Metcalfe et al., 2014; Price, 2002; Rinker and Lowman, 2004). According to the Janzen-Connell hypothesis host-specific herbivores maintain the high plant diversity of tropical forests (Clark and Clark, 1984; Connell, 1971; Janzen, 1970; Wright, 2002). Herbivorous insects are major consumers of plant material and in turn an important resource as prey or host for predators and parasitoids (Coley and Barone, 1996; Janzen, 1987; Mattson and Addy, 1975; Price, 2002). Especially herbivorous beetles, particularly Chrysomelidae, and their degree of host-specificity have played a fundamental role in species number estimates (Erwin, 1982; Novotny et al., 2002b; Ødegaard, 2000). In many habitats (e.g. tropical rainforest canopy) leaf beetles represent a large part of the herbivorous insect fauna (Andrew and Hughes, 2004; Basset, 2001; Wagner, 2000) and are essential for a true understanding of insect communities or plant-herbivore-interactions (Flowers and Hanson, 2003). An advantage is that they are easily collected and readily noticed even by non-specialists (Flowers and Hanson, 2003). Therefore, beside a multitude of studies on leaf beetle morphology and biology (e.g. Jolivet (1994); Jolivet et al. (1988); Schmitt (1994); Suzuki (1994)), a focus in Chrysomelidae research lies on their plant-herbivore-interactions (e.g. Adati and Matsuda (1993); Descampe et al. (2008); Flowers and Janzen (1997); García-Robledo et al. (2013a); Hawkeswood (1986); Jolivet (1999); McKenna and Farrell (2005); Meskens et al. (2008)).

Studies on diversity of leaf beetle communities usually address biodiversity of a certain region and often analyse the turnover along environmental gradients (e.g. Andrew and Hughes (2004); Aslan and Ayvaz (2009); Baselga and Novoa (2007); Şen and Gök (2009); Furth (2013); Gavrilović and Čurčić (2013); Lesage et al. (2008); Linzmeier et al. (2006); Ohsawa and Nagaike (2006); Sánchez-Reyes et al. (2014)).

So far there have been comparatively few studies of leaf beetle diversity in Neotropical ecosystems (Flowers and Hanson (2003); e.g. Charles and Bassett (2005); Farrell and Erwin (1988); Furth (2013); Linzmeier et al. (2006); Linzmeier and Ribeiro-Costa (2009); Sánchez-Reyes et al. (2014)). As for Neotropical Chrysomelidae the poor taxonomic situation impedes species-level identification (see Chapter 1.4) methods postponing species-level identification and using morphospecies or MOTUs instead are standing to reason. Recently, DNA barcoding approaches have been used for studying leaf beetle diversity and ecology (e.g. García-Robledo et al. (2013a,b, 2015); Germain et al. (2013); Jurado-Rivera et al. (2009); Kubisz et al. (2012); Papadopoulou et al. (2013)).

### 1.3.2 Systematics and taxonomy

Chrysomelidae are considered monophyletic (Duckett et al., 2004; Gómez-Zurita et al., 2007; Reid, 1995), but basal relationships within Chrysomelidae are not yet ultimately agreed on (Gómez-Zurita et al., 2008). For recent phylogenies see e.g. Reid (1995), Farrell (1998), Duckett et al. (2004), Farrell and Sequeira (2004), and Gómez-Zurita et al. (2008). Orsodacnidae (Orsodacninae and Aulacoscelidinae) and Megalopodidae (Megalopodinae, Zeugophorinae and Palophaginae) that have been included in Chrysomelidae by several authors are currently considered to be basal Chrysomeloidea (Duckett et al. (2004); Reid (1995); overview in Gómez-Zurita et al. (2008)). In contrast to former classifications into up to 16 subfamilies (Seeno and Wilcox, 1982), there are currently 12 well delineated taxonomic groups: Bruchinae, Cassidinae (including hispines), Chrysomelinae, Criocerinae, Cryptocephalinae, Donaciinae, Eumolpinae, Galerucinae (including alticines), Lamprosomatinae, Sagrinae, Spilopyrinae, and Synetinae (Bouchard et al., 2011; Gómez-Zurita et al., 2007). Seed beetles (Bruchinae) have traditionally been treated as a separate family (Riley et al., 2002) and are not included in the present study. Protoscelidinae is an extinct subfamily (Bouchard et al., 2011).

The following taxa are relevant for the present study and therefore briefly described:

#### **Galerucinae LATREILLE, 1802 and Alticinae SPINOLA, 1844**

Galerucinae s.l. (= Galerucinae sensu Reid (1995), or 'Trichostoma') are morphologically diverse. Their monophyly is generally acknowledged and they are typically treated as two groups, Alticinae/Alticini and Galerucinae s.str./Galerucini (Duckett et al. (2004); Lingafelter and Konstantinov (1999); and references therein).

Alticinae (flea beetles; Fig. 1.2A) comprise around 8,000 species (Furth et al., 2003). Their body shape is compactly ovate and convex (Reid and Beatson, 2013). They are easily recognized by their thickened hind femora which contain the metafemoral spring (Furth, 1982b, 1988), an internal structure allowing the beetle to perform huge jumps to escape from predators (Maulik, 1929). It has been widely used as distinguishing character between Alticinae and Galerucinae (Furth, 1988). In contrast, Galerucinae s.str. (Fig. 1.2B) with ~6000 species (Ge et al., 2012; Jolivet, 1988) are more loosely elongate and depressed and lack the metafemoral spring (Reid and Beatson, 2013).

Galerucinae usually feed on dicotyledons (Mariau, 2004). They are basically Cucurbitaceae, Leguminosae, or Verbenaceae feeders and adapted to many plant families; host plants from almost 100 families have been recorded (Jolivet, 1988). A very large New World genus that includes several significant agricultural pests is *Diabrotica* with 300 (s.str.), respectively 600 (s.l.) species (Hammack and French, 2007; Jolivet, 1988). Alticinae have an especially complex food selection (Jolivet, 1988). Most are specialized and well-adapted to their host plant (Jolivet, 1988). It has been observed that Alticinae chew completely different plants at the end of the season (Jolivet, 1988).

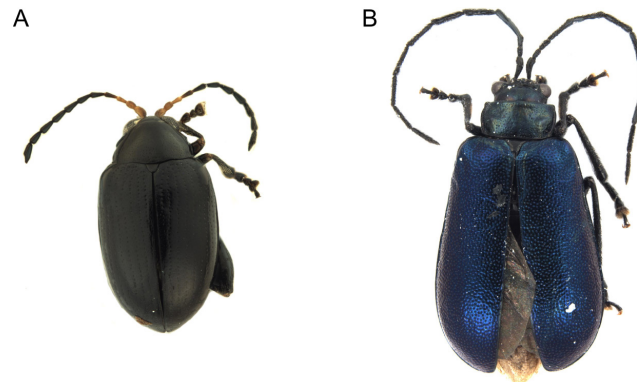


Figure 1.2: **Alticinae (A)**, **Galerucinae (B)**. Specimens 2050\_Alticinae\_sp\_123 and 3438\_Galerucinae\_sp\_031.

Relationships between the closely related Galerucinae s.str. and Alticinae are controversial (Duckett et al., 2004): Alticinae have been treated as a separate subfamily or as a tribe (Alticini) within Galerucinae (Furth and Suzuki, 1994; Lingafelter and Konstantinov, 1999; Reid, 1995; Seeno and Wilcox, 1982). The monophyly of either Alticinae and/or Galerucinae (i. Alticinae nested within Galerucinae: Lingafelter and Konstantinov (1999), ii. reciprocal monophyly of the two groups: Gómez-Zurita et al. (2008), or iii. monophyletic Galerucinae within flea beetles: Duckett et al. (2004); Kim et al. (2003); Reid (1995)) has been challenged by Ge et al. (2011, 2012) who included several problematic taxa considered 'incertae sedis' and propagated multiple origins of the complex jumping mechanism.

For a better understanding in this study the terms Galerucinae and Alticinae are retained (according e.g. Furth and Suzuki (1994); Jolivet and Petitpierre (1981); Seeno and Wilcox (1982)) keeping in mind that their status as subfamilies of equal rank and also their respective monophyly is in question (e.g. Crowson and Crowson (1996); Lingafelter and Konstantinov (1999); Reid (1995); for an overview see Lingafelter and Konstantinov (1999)).

### **Cassidinae GYLLENHAL, 1813 and Hispinae GYLLENHAL, 1813**

Cassidinae s.l. (Cassidinae s.str. + Hispinae, or 'Cryptostoma') are noteworthy for their specialized plant associations with monocots and eudicots, diverse morphologies in immatures and adults, and a range of social behaviours from solitary to subsocial (Chaboo and Engel, 2009). They are cosmopolitan but primarily tropical, and most species are found in the Neotropics (Chaboo, 2007). New and Old World fauna show little overlap (Chaboo, 2007). They have a broad variation in host plant selection, from polyphagous to oligophagous or monophagous to plant species (Chaboo, 2007).

Until recently, two groups of Cassidinae s.l. have been treated as two subfamilies by most authors (e.g. Farrell (1998); Seeno and Wilcox (1982); Verma (1996));

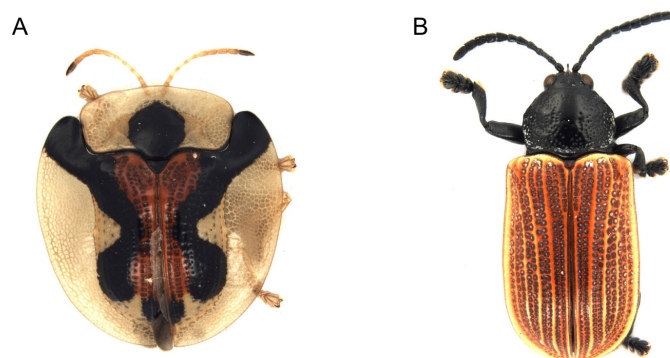


Figure 1.3: **Cassidinae (A), Hispinae (B)**. Specimens 3861\_Cassidinae\_sp\_015 and 4783\_Hispinae\_sp\_016.

Hispinae GYLLENHAL s.str. (leaf-mining beetles) and Cassidinae GYLLENHAL s.str. (tortoise beetles). A detailed history of the classification of the two groups is given by Borowiec (1995) and Staines (2002) and research on their relationships is reviewed in Chaboo (2007). Morphologically and biologically there is no valid reason for retaining Hispinae and Cassidinae as separate subfamilies; intermediate forms (e.g. *Cephaloleiini*, *Imatidiini*) bridge the subfamilies (Borowiec, 1995; Staines, 2002). Currently, Cassidinae is the correct name for the clade Hispinae + Cassidinae with the hispine or hispiform genera being considered as a basal grade of Cassidinae (Borowiec, 1995; Chaboo, 2007; Chaboo and Engel, 2009; Staines, 2002). However, as the two groups reflect two characteristic, complex morphologies and ecological-behavioural forms and for a better understanding in this study the terms Hispinae and Cassidinae (meaning Cassidinae s.str.) are used.

There are ca. 3000 species (Jolivet, 1988) of Cassidinae s.str. (tortoise beetles). They are one of the most specialized chrysomelid subfamilies (Jolivet, 1988). They have a characteristic tortoise-like form induced by broadly expanded elytral and pronotal margins that frequently shield the heads (Chaboo and Engel (2009); Fig. 1.3A). They are reluctant flyers and there are brachypterous and wingless cassidine species (Chaboo, 2007). Tortoise beetles show an extremely diverse colouration, including metallic, iridescent, transparent, silver, and golden (Chaboo, 2007). Colour change and colour polymorphism occur frequently (Buzzi, 1988; Chaboo, 2007).

Although Cassidinae feed on 32 plant families, Convolvulaceae and Asteraceae are preferred (Borowiec and Świętojańska, 2014; Chaboo, 2007; Jolivet, 1988). Especially many tropical species feed on Convolvulaceae (Jolivet, 1988). Genera close to Hispinae (e.g. *Imatidium*) feed on palm trees (Jolivet, 1988).

Cassidinae show an interesting behavioural repertoire, especially concerning reproduction: Courtship behaviour has been described for several species and in some species post-copulatory attendance has been observed (Chaboo, 2007). Mating can last more than 24 hours (Chaboo, 2007). Some cassidine larvae construct a shield

from faecal material and exuviae that is carried over the dorsum and is retained by some pupae (Chaboo, 2007; Chaboo and Engel, 2009). Those shields can be very sophisticated and show a remarkable variety of architectures (Chaboo, 2007). They protect the immature cassidines from predation and desiccation (Chaboo and Engel, 2009). In many cassidines larval gregariousness is common (Chaboo, 2007). For 17 cassidine species maternal care has been observed, a behaviour that is very rare in beetles and insects in general (Chaboo, 2007). Females have been observed sitting on top of the eggs, larvae, and pupae in order to protect them (Buzzi, 1988; Chaboo, 2007). They guard the immatures until the young adults emerge and attack threatening predators such as ants or reduviids (Chaboo, 2007). Females herd their larvae and, in the face of threats, even guide them to new leaves by prodding and pushing them (Chaboo, 2007).

Cassidinae are almost worldwide distributed, although they have a much greater diversity in the tropics, especially in tropical South America (Borowiec and Świętojańska, 2014). According to Blackwelder (1947) there are ~2000 Neotropical cassidine species. Ecuador's Cassidinae (s.str.) seem quite well studied compared to the other subfamilies (e.g. Borowiec (1998, 2000a,b); Flowers and Chaboo (2009); Sekerka and Windsor (2012)). A checklist of 200 species of Cassidinae recorded for Ecuador has been provided by Borowiec (1998) who estimated the total number of species living in Ecuador at ca. 250. The only chrysomelid type specimens deposited at the Invertebrate Section of the Museum of Zoology at the Pontifical Catholic University of Ecuador, Quito, are all Cassidinae (Donoso et al., 2009).

The approximately 3000 species of Hispinae (leaf-mining beetles) contain typically spiny or strongly sculptured beetles (Chaboo and Engel (2009); Jolivet (1988); Fig. 1.3B). Their immatures are broadly characterized as leaf-miners, although their biology ranges from leaf-tube scrapers to stem-miners, and even to open-leaf feeders (Chaboo, 2007; Chaboo and Engel, 2009). Most Hispinae feed on Monocotyledons, however others on Dicotyledons, in at least 37 plant families (Jolivet, 1988). Many Hispinae feed on palm trees (Jolivet, 1988). Noteworthy are the Neotropical 'rolled-leaf' hispine beetles (or 'hispid Cassidinae', principally the tribe Cephaloleiini, >380 Neotropical species; Descampe et al. (2008)). They are found in tightly rolled apical leaves of monocots, mostly closely associated with Zingiberales, some with Arecaceae (Descampe et al., 2008; McKenna and Farrell, 2005). The close association of Hispinae with Zingiberales probably exists for >60 ma (Wilf et al., 2000). A review of the hispine/Zingiberales interaction was published by Staines (2004).

Seeno and Wilcox (1982) recorded 83 genera of hispines from the New World (Staines, 2002). There have been several regional revisions of New World hispines (e.g. Monrós and Viana (1947); Sanderson (1967); Staines (1996)), however none especially for Ecuador.

### **Eumolpinae HOPE, 1840**

With more than 7000 species in 500 genera the subfamily Eumolpinae is the third in species diversity after Galerucinae s.l. and Cassidinae s.l. (Chaboo, 2007; Jolivet and Verma, 2008). They are worldwide distributed but basically a tropical group



Figure 1.4: **Eumolpinae with shiny, rugose, and hairy elytra (from left to right)**. Specimens 0312\_Eumolpinae\_sp\_021, 0719\_Eumolpinae\_sp\_043, and 0553\_Eumolpinae\_sp\_042.

where they are especially numerous (Jolivet and Verma, 2008). Their typical body forms are oblong, convex, and globose, but some are quite elongated. Antennae are usually filiform and insertions are not closely approximated. Eumolpinae have elytra with well-defined shoulders and are generally smooth, often shiny with metallic colours. In contrast, some are dull coloured, some rugose, and some have elytra and body densely hairy (Jolivet and Verma (2008); Fig. 1.4).

From a basic oligophagy on Asclepiadaceae, Apocynaceae and Convolvulaceae, Eumolpinae became in many genera polyphagous and feed on many wild and cultivated plants (Jolivet, 1988). Food plants are recorded from 116 plant families (Jolivet, 1988). Eumolpine larvae are root feeders (Jolivet and Verma, 2008). Neotropical eumolpine fauna is mostly constituted by the tribe Eumolpini (Flowers, 1999). Blackwelder (1947) lists 44 species for Ecuador. A new genus and several species have been described for Ecuador by Flowers (2004a,b, 2009a,b, 2004c).

#### **Criocerinae LATREILLE, 1804**

With ~1400 species Criocerinae (shining leaf beetles) is a relatively small subfamily (Schmitt, 1988; Vencl et al., 2004). Most species belong to five species-rich genera (*Crioceris*, *Lilioceris*, *Lema*, *Oulema*, and *Neolema*) (Matsumura et al., 2014). Criocerinae live in the temperate, subtropical, and tropical zones of all continents (Schmitt, 1988). They are glabrous with a brilliant metallic sheen (Vencl et al., 2004). They are typically narrow, elongate, depressed to cylindrical, with the pronotum medially or basally constricted ('hourglass-shape') and often differently coloured from the rectangular elytra (Cooter and Barclay, 2006; Reid and Beatson, 2013). Head and pronotum are narrower than the elytra (Fig. 1.5).

A characteristic of all Criocerinae is the ability to produce chirping sounds by means of an elythro-abdominal stridulatory apparatus (Schmitt, 1988). They probably use these sounds to deter predators (Schmitt, 1988; Schmitt and Traue,





Figure 1.5: **Criocerinae**. Specimen 4209\_Criocerinae\_sp\_007.

1990). Detailed information about stridulation of Criocerinae is given in the study by Schmitt and Traue (1990).

Criocerinae are quite well-studied because of their economic interest, feeding on both Mono- (six plant families) and Dicotyledons (12 families) (Jolivet, 1988). The main feeding habit of both adults and larvae is leaf surface grazing (Vencl et al., 2004). For the New World, over 460 species are described (Vencl et al., 2004).

#### **Chrysomelinae LATREILLE, 1802**

Chrysomelinae comprise ca. 3000 species (Daccordi, 1996; Reid and Beatson, 2013). They are generally ventrally flattened and dorsally convex, with ovate body shape (Reid and Beatson (2013); Fig. 1.6). Neotropical chrysomelines comprise beside cassidines some of the largest and most colourful representatives of Chrysomelidae. In Costa Rica they are popularly known as 'confites con patas' (walking candies) (Flowers, 2004c).



Figure 1.6: **Chrysomelinae**. Specimen 0201\_Chrysomelinae\_sp\_002.

For Chrysomelinae, 47 families of dicotyledonous host plants have been recorded (Jolivet, 1988). Most genera are monophagous or polyphagous on related host plants (Jolivet, 1988). In the New World, Solanaceae is the mostly selected family (Jolivet, 1988). In the Neotropical region chrysomelinae are very numerous (Daccordi, 1996). A key for Chrysomelinae genera for Venezuela by Bechyné and Springlová de Bechyné (1965) was adapted for Costa Rica by Flowers (2004c).

### **Cryptocephalinae GYLLENHAL, 1813 and Lamprosomatinae LACORDAIRE, 1848**

Cryptocephalinae and Lamprosomatinae together with Clytrinae and Chlamisinae (often placed within Cryptocephalinae; Bouchard et al. (2011); Reid (1995)) share several morphological characters and are often referred to as 'Camptosomata' (Erber, 1988). They are also called 'case-bearers' because one common feature is a mantle, females cover their eggs with and that is worn as protective case by the larvae (Erber, 1988). As many camptosome species live cryptically or let themselves fall at the least disturbance, relatively little is known about their life-habits, e.g. feeding habits (Erber, 1988).

There are ~3900 species of Cryptocephalinae (Reid and Beatson, 2013). Cryptocephalinae have a cylindrical body that is obtusely rounded in front and behind, and almost circular in cross-section (Erber, 1988). The prothorax is in most cases at its base as broad as the elytra and joined to them without any suture and it tapers slightly in front (Erber, 1988). The head is placed closely against the prothorax, without a neck and hypognathous (Erber, 1988). Antennae are relatively short (Erber, 1988). Although colouring varies there are many shining metallic species (Erber (1988); Fig. 1.7A). Some cryptocephaline species feed on the leaves of woody plants, many on herbs, some feed on petals and there are even pollen-feeders (Erber, 1988).

Lamprosomatinae are a small subfamily with ~120 species (Reid and Beatson, 2013). The body-outline from Lamprosomatinae tapers in front and behind and is oval, like an egg (Erber, 1988). In side-view they are strongly convex, tapering away posteriorly, and they are ventrally flattened (Chamorro and Konstantinov, 2011; Erber, 1988). They are shiny and usually iridescent (Chamorro and Konstantinov (2011); Fig. 1.7B). Lamprosomatine diet seems to be restricted to herbs (Erber, 1988).

## **1.4 Chrysomelidae research in Ecuador**

Ecuador, situated within the peak of species richness at tropical low latitudes, is considered a megadiverse country. On a comparatively small area it harbours pronounced elevational and wet-dry gradients with a large variety of habitats and high beta-diversity (Brehm et al., 2008a). The Ecuadorian leaf beetle fauna can be expected to be megadiverse, however has hitherto scarcely been studied.

Although entomology has a long history in Ecuador as in South America in

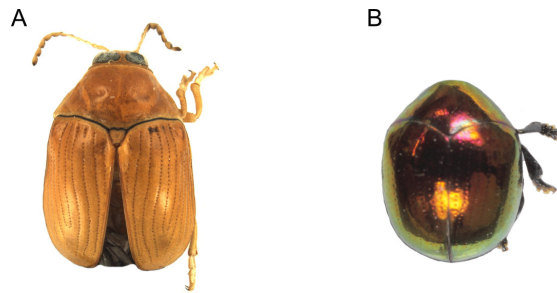


Figure 1.7: **Cryptocephalinae (A)**, **Lamprosomatinae (B)**. Specimens 0924\_Cryptocephalinae\_sp\_001 and 1242\_Lamprosomatinae\_sp\_003.

general (Barragan et al., 2009), with respect to the knowledge of its invertebrate fauna it remains like many other tropical countries a white spot on the map (Beck and Kottke, 2008). Whereas the diversity of certain charismatic groups such as plants, birds, and frogs has been the focus of numerous publications, data on the entomological fauna in Ecuador are scarce (Dangles, 2009). The Ecuadorian research in entomology was dominated by taxonomic studies during the past decades.

In general, Neotropical beetle fauna with exception of certain taxa such as Cerambycidae or Scarabaeoidea is considerably understudied and there are no general books or treatises about that region (Costa, 2000). Accordingly, also the available data records for Ecuadorian chrysomelid fauna is very sparse: Blackwelder's 'Checklist of the coleopterous insects of Mexico, Central America, the West-Indies, and South America' (1947) compiled between 1944 and 1957 (complemented by Bechyné (1952)) is still regarded as standard reference for South American beetle diversity and was only complemented by extensions and revisions for certain taxa or countries (e.g. Bechyné (1953): Eumolpinae, Maes and Staines (1991): Chrysomelidae of Nicaragua, Furth and Savini (1996): Alticinae, Borowiec (1998): Cassidinae). Explicitly for Ecuador, Blackwelder lists ~450 chrysomelid species (another ~100 listed for tropical or South America; Blackwelder (1947)). This number is certainly far below the true species number. Furth et al. (2003) claim that 'all central American countries certainly have a much higher actual diversity than is recorded in the literature'. This surely applies as well to South America in general and Ecuador in particular. A recent review particularly for Ecuador exists only for Cassidinae (Borowiec, 1998) with 200 recorded and ~250 estimated species.

In the Neotropics, a few regions experienced an extensive collecting and research activity. For example, in the 1960s, the entomologists Jan and Bohumila Bechyné who studied Neotropical leaf beetles more extensively than most previous workers collected intensively in Venezuela and described over 90 alticine genera as well as hundreds of species (Furth and Savini, 1996). Mainly due to their work, the collection of the Museo del Instituto de Zoología Agrícola 'Francisco Fernández Yezpez', Venezuela, is one of the most important collections of Neotropical Chrysomelidae, especially Alticinae. It harbours more than 1100 alticine species, giving an idea of the

true species richness of the South American countries. Another example of a better study situation compared to most Neotropical countries is Costa Rica that during the last years became a focus area for Neotropical biodiversity research resulting in a lot of publications, a number of them about Chrysomelidae (e.g. Flowers (1991); Flowers and Janzen (1997); Furth et al. (2003); Staines (2011)). In a study about Alticinae in Costa Rica, the species number recorded in literature was more than doubled resulting in a total of ~350 known species for the whole country (and maybe a total number of 1000 species appearing realistic; Furth et al. (2003)). However, these examples of well-studied countries are exceptions. A similar high diversity as in those countries should be expected for Ecuador, however, Blackwelder lists only ~65 species of alticines (Halticinae) for Ecuador and a recent review is lacking. Most Neotropical countries' leaf beetle diversity remains barely explored.

Although Chrysomelidae are attractive study organisms (see Chapter 1.3), the sheer diversity of the family presents a challenge for studying their diversity in tropical ecosystems. Species-level identification is often impossible. For Neotropical Chrysomelidae, the few existing keys are mostly quite dated: The only key to Neotropical alticine genera by Scherer is from 1962 (Furth et al., 2003; Scherer, 1962); the first revised key for New World genera of Hipines since Weise (1911) was only in 2002 published by Staines. Others are restricted to certain regions (Flowers, 2004c; Staines, 2009), genera (Flowers, 2004a,b; Staines, 2013), or small groups of related genera (Furth, 1992; Konstantinov and Konstantinova, 2011). An exception is the interactive manual 'Cassidinae of the World' (Borowiec and Świętojańska (2014); <http://culex.biol.uni.wroc.pl/cassidae/katalog%20internetowy/index.htm>) that provides a comprehensive key to cassidine genera worldwide. However, reliable identification keys to genera are still lacking for genera of some of the largest and most ubiquitous subfamilies of Neotropical Chrysomelidae (Flowers and Hanson, 2003). None exists particularly for Ecuadorian leaf beetle fauna.

Whereas most studies about Ecuadorian Chrysomelidae are records or descriptions of individual species or genera (e.g. Borowiec (1998, 2000a,b); Flowers (2009a,b); Sekerka and Windsor (2012); Staines and Zamorano (2012); Świętojańska and Borowiec (2000)) or host records (e.g. Flowers and Chaboo (2009)), there are no studies about the diversity of leaf beetle communities in Ecuador. Also a thorough inventory of mountain forests is missing.

The high discrepancy between recorded and true diversity is also reflected by its collections: The Invertebrate Section of the Museum of Zoology at the Pontifical Catholic University of Ecuador in Quito comprises with almost two million specimens Ecuador's largest collection of native taxa. It harbours 24,215 Chrysomelidae, most of them still awaiting identification. 10.83% of all specimens are determined to species, 13.6% to genus, but 75.56% have no identification at all (Clifford Keil, pers. comm.). There are only nine chrysomelid type specimens deposited, all of them belong to the subfamily of Cassidinae (Donoso et al., 2009). The distribution of type localities (for all invertebrates) showed that collection sites are clustered geographically with most of them found towards the northern region of Ecuador. Sites are mainly located in highly accessible areas near highways and towns (Donoso et al.,

2009). Donoso et al. (2009) advise that 'future fieldwork should include localities in the southern region of Ecuador but also target less accessible areas'.

Concerning the study area of the present study, among insects only certain taxa of Lepidoptera (Bodner et al., 2010; Brehm et al., 2003a, 2005, 2003b; Fiedler et al., 2008; Häuser et al., 2008; Hilt and Fiedler, 2006) and Orthoptera (Braun, 2008) have been studied up to now. The large insect orders of Coleoptera, Hymenoptera and Diptera remain completely unstudied so far (Brehm et al., 2008b). Notably Chrysomelidae are mentioned by Brehm et al. (2008b) to be desirable to be studied in the study area as they have high impact on forest ecosystems. A first attempt of studying beetle diversity in the study area was made by Schmidl (2007–2008, pers. comm.). He confined himself on the bark-living fauna and reported ~50 morpho-species of Chrysomelidae.

## 1.5 Aims and structure of the dissertation

### Objectives

This study aims at testing a combined morphological and molecular approach for assessing rapidly the biodiversity of an unknown leaf beetle fauna in a mountain forest in southern Ecuador. The performance of DNA barcodes as substitutes for Linnean taxonomic information is evaluated for identification of species-like units. In detail, the following questions are addressed:

1. How diverse are leaf beetles in the studied Andean mountain forest?  
How can barcode and morphospecies data characterize an unknown leaf beetle community? What assertions can be made about species richness, abundance, incidence, and subfamily composition of the community?
2. How do local and regional species richness change with elevation?  
Can barcode data analyse patterns of species richness, turnover, and community composition along an elevational gradient? Does DNA barcode data reveal diversity patterns in a comparable way as morphospecies do? Which ecological conclusions can be drawn from DNA barcode data?
3. How congruent are the morphospecies method and the DNA based identifications?  
Which discrepancies are there, in which taxa do they occur, and what are their reasons?
4. Which influence do different methods of species delimitation have on species richness estimates?  
How congruent are different DNA based species delimitation methods? How relevant is the choice of the species delimitation method?
5. How do different sampling methods perform?  
Which sampling methods are advisable in terms of sampled specimens and time efficiency?

### Structure of the thesis

The 'Methods' section provides extensive and detailed general information about study area, sampling, and further handling of specimens as well as subsequent laboratory and data analyses. In the individual chapters, specific methodological information relevant for the respective part is given.

Each of the five chapters on leaf beetle biodiversity and ecology and methodological aspects of rapid biodiversity assessment represents a separate study and can be understood by itself. Each follows the standard structure for a scientific publication (introduction, methods, results, discussion, and conclusion) as they are meant to be published as separate publications in scientific journals. Therefore, some content is recurring throughout the thesis.

In the 'General discussion and future prospects' section, conclusions are drawn from the complete study and some future perspectives are outlined.

This thesis was conducted within the framework of the research programme 'ABA-Ecuador: Acceleration of biodiversity assessment – Development of tools and application in a tropical mountain ecosystem'. The project on Chrysomelidae was funded by the German Science Foundation (Deutsche Forschungsgemeinschaft, DFG), grant Wa 530/46-1.

### Teamwork

Chapter 4 is prepared as a manuscript for publication in a scientific journal:

Thormann, Birthe; Ahrens, Dirk; Marín Armijos, Diego; Wagner, Thomas; Wägele, J. Wolfgang; Peters, Marcell K. **Topography effects on elevational alpha-, beta-, and gamma-diversity of Neotropical leaf beetles.**

B. Thormann, M.K. Peters, and J.-W. Wägele developed the study. B. Thormann conducted sampling, preparation of specimens, laboratory work, and data compilation. B. Thormann and Th. Wagner conducted morphospecies sorting. B. Thormann and D. Ahrens conducted molecular species delimitation. B. Thormann and M.K. Peters performed the statistical analyses and developed the first version of the manuscript.

The contents of Chapter 6 are supposed to be published as:

Thormann, B.; Ahrens, D.; Marín Armijos, D.; Peters, M.K.; Wagner, Th.; Wägele, J.-W. **Exploring the leaf beetle fauna (Coleoptera: Chrysomelidae) of an Ecuadorian mountain forest with DNA barcoding.**

B. Thormann, D. Ahrens, M.K. Peters, and J.-W. Wägele developed the study. B. Thormann conducted sampling, preparation of specimens, laboratory work, and data compilation. B. Thormann and Th. Wagner conducted morphospecies sorting. B. Thormann and D. Ahrens conducted molecular species delimitation. B. Thormann and M.K. Peters performed the statistical analyses. All co-authors provided ideas and suggestions for the text.

The study was carried out in close cooperation with the Universidad Técnica Particular de Loja (UTPL), Ecuador.

## 2.1 Study area

The study area is situated within the Reserva Biológica San Francisco (RBSF) and the adjacent Podocarpus National Park (NP) in southern Ecuador. The RBSF is a small private nature reserve ( $\sim 11.2 \text{ km}^2$ ) owned by the foundation Nature and Culture International, NCI. It is located between the province capitals Loja and Zamora and harbours the research station 'Estación Científica San Francisco' (ECSF;  $3^\circ 58' 17.19'' \text{S}$ ,  $79^\circ 4' 44.06'' \text{W}$ ; Fig. 2.1). Podocarpus NP was created in 1982, comprises  $\sim 1463 \text{ km}^2$ , and is part of the Podocarpus – El Condor Biosphere Reserve. Politically, the study area belongs to the provinces of Loja and Zamora-Chinchipe. The study sites are situated in three different areas: (i) ECSF area next to the research station, belonging to RBSF, (ii) Bombuscaro area in the Bombuscaro sector of Podocarpus NP, close to Zamora, and (iii) Cajanuma area in the Cajanuma sector of Podocarpus NP, close to Loja (Fig. 2.1). In Bombuscaro and Cajanuma are the two main entrances to Podocarpus NP.

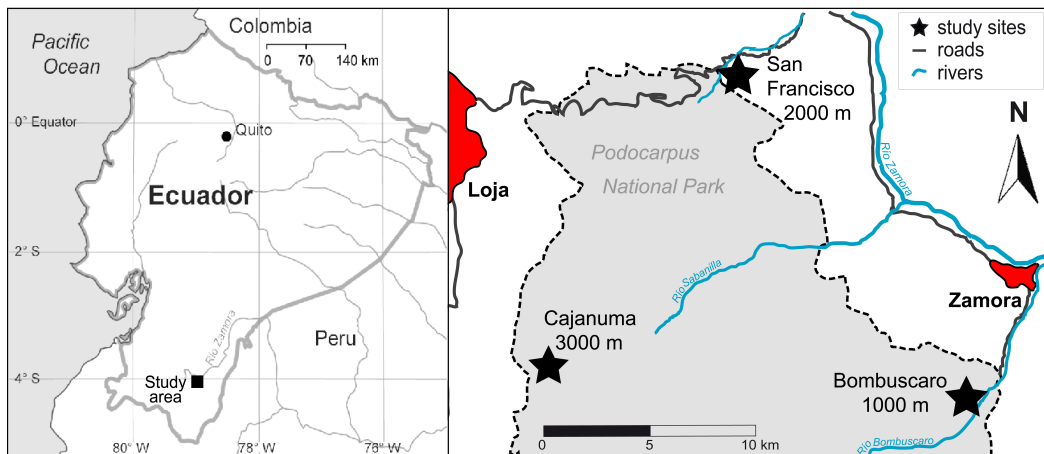


Figure 2.1: Map of Podocarpus National Park, Ecuador, with location of study sites. Bombuscaro (1000 m), ECSF (=San Francisco; 2000 m), Cajanuma (3000 m) (Homeier et al., 2012).

The region is located in the Cordillera Real or Eastern Cordillera of the Andes and is part of the Huancabamba depression. In contrast to central and north Ecuador with impressive volcanoes of up to 6000 m a.s.l. and above, in the Huan-

cabamba depression there are no volcanoes, the crest of the Cordillera Real does not exceed 2800–3400 m a.s.l., and the treeline is comparatively low (Beck et al., 2008b; Richter et al., 2009, 2008). Eastern and Western Cordillera, the two main cordilleras forming the Ecuadorian Andes, are interconnected by transverse mountain bridges forming ten interandean basins. The complex topography creates a landscape with extreme climatic differences (Dangles et al., 2009).

The research area has a tropical humid climate with annual mean precipitation of 2230 mm in Bombuscaro, 1950 mm in ECSF, and up to 4500 mm in Cajanuma (Moser et al., 2007). Precipitation is high throughout the year, however with the main rainy season from April to August and a less humid period from September to December (Bendix et al., 2006, 2008). Annual mean air temperature decreases from 19.4°C in Bombuscaro and 15.7°C at ECSF to 9.4°C in Cajanuma (Moser et al. (2007); Tab. 2.1).

Table 2.1: **Temperature and precipitation at the three study sites (Moser et al., 2007).**

	Bombuscaro	ECSF	Cajanuma
Annual mean air temperature (°C)	19.4	15.7	9.4
Min.	11.5	7.9	3.1
Max.	30.2	29.4	18.8
Relative air humidity (%)	88,7	90,8	93,5
Min.	15.5	15.7	28.6
Max.	100	100	100
Soil moisture (vol %)	29.7	35.4	49.1
Min.	15.3	27.4	39.5
Max.	38.5	44.7	59.5
Annual mean precipitation (mm y <sup>-1</sup> )	2230	1950	4500

The study area is a rugged mountainous area with valleys and differently exposed slopes mostly covered with evergreen (pre-)montane rainforests. An overview of vegetation types of the region can be found in Homeier et al. (2008). Its steep elevational gradients and great topographic heterogeneity create a broad matrix of environmental conditions and a mosaic of different habitats in close proximity to each other leading to a high diversity of animals and plants (Homeier et al., 2010, 2008). This is reinforced by the geographic position of the area: Its location between the humid eastern Andean slope and the dry Inter-Andean region on the one hand, and between the Central and Northern Andes on the other hand allows an intermingling of the respective characteristic species pools (Homeier et al., 2010).

The region is part of a biodiversity hotspot, the Tropical Andes (Myers et al., 2000). It is ranked as a top hotspot of diversity of vertebrates and vascular plants (Brummitt and Lughadha, 2003). There exist only little information about its insect fauna but an exceptionally high level of diversity and endemism is certain (Larsen et al., 2011). The outstandingly high plant diversity has extremely high proportions of endemism (Barthlott et al., 2005; Richter et al., 2009; Weigend, 2002; Young



and Reynel, 1997). In particular the flora of Podocarpus NP is known for its high endemism (Homeier et al., 2008). The tree diversity is higher than in comparable montane forests in northern Ecuador, and also for vascular epiphytes, bryophytes, and orchids extremely high species numbers have been recorded (see references in Brehm et al. (2008b)).

Knowledge of the region's fauna is still very incomplete. Podocarpus NP is one of Ecuador's most important bird areas with more than 550 species recorded (Rahbek et al., 1995). Among the most charismatic mammals of the area are the Spectacled Bear (*Tremarctos ornatus* CUVIER) and the Mountain Tapir (*Tapirus pinchaque* ROULIN). Podocarpus NP is one of the 137 most irreplaceable protected areas of the world (Le Saout et al., 2013).

For the RBSF there exist inventories of birds, bats, and parts of Lepidoptera, Orthoptera, and Arachnida showing high diversity of these taxa (Brehm et al., 2008b). Notably the diversity of geometrid moths (Lepidoptera: Geometridae) has been intensively studied by G. Brehm and colleagues. Besides analysing e.g. the diversity along elevational or successional gradients, their recorded species numbers are the highest ever counted in a single study on such a small spatial scale (e.g. Brehm (2002); Brehm et al. (2003a, 2005, 2003b); Hilt et al. (2006)). In contrast, amphibians, molluscs, and the vast majority of arthropods, including Coleoptera, Hymenoptera, and Diptera, have not been studied at all (Brehm et al., 2008b).

The manifold habitats of the region with their outstanding biodiversity are threatened by deforestation. Ecuador's dense population (55 inhabitants/km<sup>2</sup>) puts strong pressure on natural ecosystems (Dangles et al., 2009). Ecuador suffers the highest deforestation rate (-1.9% per year) in South America (FAO, 2010). The main reason is the conversion of forest into agropastoral land (Mosandl et al., 2008).

## 2.2 Sampling methods and sampling design

Sampling was conducted in November and December 2010 and from 08.05.2011 to 26.04.2012. Mainly three different methods were used to collect leaf beetles from the lower to medium understorey vegetation: (i) sweep netting, (ii) beating of shrubs and smaller trees using a beating-tray, and (iii) hand-collection (picking up beetles individually from the vegetation). Those methods are widely used and promising for sampling of leaf beetles (Thomas Wagner, pers. comm.). All kind of vegetation within reach was sampled (up to ca. 2.5 m). In addition, Malaise- and flight interception-, as well as light-trapping was used. Pitfall traps have been tested but only in individual cases. The sampling methods are described in more detail in Chapter 7.

Sampling was conducted in three different areas within Podocarpus NP and RBSF (see Chapter 2.1): (i) Bombuscaro ('1000 m'; elevation 1020–1075 m a.s.l.; premontane rainforest), (ii) ECSF (in the vicinity of the Estación Científica San Francisco; '2000 m'; elevation 1913–2089 m a.s.l.; lower montane rainforest), and (iii) Cajanuma ('3000 m'; elevation 2805–2891 m a.s.l.; upper montane rainforest or

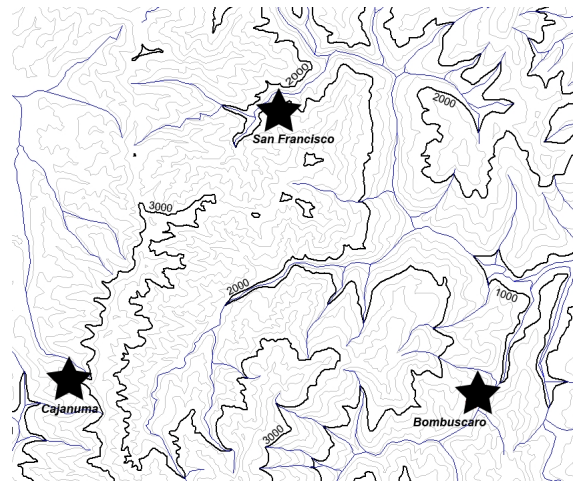


Figure 2.2: **Contour map of the study area.** Contour lines within the study area and location of the three sampling areas Bombuscaro, ECSF (San Francisco), and Cajanuma (Jantz et al., 2014).

cloud forest) (Classification of vegetation: Homeier et al. (2008); Fig. 2.2). Linear distance between each of the three sampling areas is  $\sim 20$  km.

To analyse biodiversity and community patterns and make comparisons along the altitudinal gradient, most of the sampling was conducted in a standardized way on defined study sites: The study sites are the so called Matrix-Plots which have been established in 2007 by C. Leuschner and E. Veldkamp and colleagues (University Göttingen). They are  $20 \times 20$  m squares situated within homogeneous mature forest representative for the elevation in question and without visible recent natural or human disturbance (Homeier et al., 2010). At each elevational area, 12 plots were studied, resulting in a total of 36 plots. At each elevational level, six plots are situated in the valley, near creeks, the so-called 'Lower slope plots' (L-Plots), and six near ridge crests ('Upper slope plots', U-Plots).

The two slope positions (U and L) correspond to two habitat types that differ in several environmental parameters and harbour two distinct forest types. Compared to ridge habitats, valley habitats have a higher productivity, are more nutrient-rich, have a minor organic layer thickness and a lower C:N ratio. The sites in valleys are vegetated with a forest which differs in species composition, has a smaller canopy openness, consists of higher trees, has a higher biomass and harbours a higher diversity of tree species (Homeier et al. (2010); Werner and Homeier (2015); Homeier pers. comm.). Figure 2.3 shows a typical ridge and a typical valley forest. Further information about the location and coordinates of the plots are given in Appendix Tab. B.1.

The sampling on the plots followed a standardized procedure carried out by a team of two persons: Sweep netting alongside two edges of the plot for 30 min by one person; simultaneous beating alongside the other two edges of the plot for 30



Figure 2.3: Typical ridge forest (left) and valley forest (right) in the ECSF area. Photos courtesy of F. Werner.

min by the other person; subsequent hand-collection within the plot for 30 min (by both persons simultaneously for 15 min). The standardized sampling on the plots was complemented with other sampling methods (see above) and additional hand-sampling. A sample is defined as the total of leaf beetles either caught by 30 min sweep netting, beating, or hand-collection on a plot (standardized samples), or by hand-collection during one sampling day (non-standardized hand-collection), or the content of a Malaise-, light-, flight interception- or pitfall trap when emptied.

Being killed and collected in 70% ethanol, beetles were subsequently transferred into 96% ethanol and stored at 4– -20°C to ensure optimal preservation of DNA.

## 2.3 Further handling of the specimens

For each sample, Chrysomelidae were sorted into preliminary morphospecies. Usually one specimen of each morphospecies of each sample was selected and used for the subsequent morphological and molecular analysis. In single cases up to three specimens have been analysed (the remaining individuals of a preliminary morphospecies are termed *duplicate specimens* hereafter and have been included in some analyses; see below, paragraph 'Morphospecies sorting').

Each selected individual was used for both, morphological classification and molecular analyses. For DNA extraction, one to three legs of the beetle were dissected, depending on its size. The specimen was then dry mounted and labeled to allow subsequent morphological investigation. Each sequence remains linked to its voucher specimen and all connected sampling information. Voucher specimens will be deposited in the collections of the Zoological Research Museum Alexander Koenig (ZFMK), Bonn, and the Universidad Técnica Particular de Loja (UTPL), Ecuador (Specimen List in the Appendix Tab. C.1). Photos of a number of the specimens will be available through the ZFMK collection database and all DNA samples are stored in the Biobank of the ZFMK. The sequences from Chapters 4 and 6 are available

from GenBank (accession numbers KJ677272–KJ677945/KR424781–KR425417; see also Appendix Tab. C.1), the rest will also be submitted.

## 2.4 Laboratory protocols

### The DNA barcode marker COI

The cytochrome *c* oxidase I (COI) gene is one of 13 protein-coding genes of the mitochondrial genome. The COI complex consists of several subunits and plays an important role in the respiratory chain (Steinke and Brede, 2006). A 658 base pair (bp) long fragment at the 5' half of the COI gene has established as the standard marker sequence for DNA barcoding of animals. Due to highly conserved sequence positions, this fragment can be amplified with so-called 'universal primers' in a wide variety of animal taxa (Folmer et al., 1994; Hebert et al., 2003a). In this study, the term 'COI' refers to this 658 bp long fragment of the cytochrome *c* oxidase I gene.

Advantages of mitochondrial genes are the lack of introns and the limited exposure to recombination as a result of the maternal mode of inheritance (Hebert et al., 2003a; Steinke and Brede, 2006). In contrast to the rRNA coding genes, the protein-coding genes usually do not contain indels that complicate analyses. The COI marker gene has a greater range of phylogenetic signal than other mitochondrial genes and is supposed to evolve rapidly enough to allow discrimination of not only closely allied species but also phylogeographic groups within a single species (Cox and Hebert, 2001; Hebert et al., 2003a,b; Wares and Cunningham, 2001).

Universal primers are necessary for identifying specimens that are not known *a priori*. Highly conserved sequence positions allow amplification of the COI marker with the universal primers LCO1490 and HCO2198 in a wide variety of animal taxa (Folmer et al., 1994; Hebert et al., 2003a; Zhang and Hewitt, 1997). However, often specific primers have been used for certain taxa and the use of more than one pair of primers can be necessary to derive sequences from all individuals of a taxonomic group (e.g. Hebert et al. (2004); Smith et al. (2005); Ward et al. (2005)). For degenerated DNA mini-barcodes of 100 to 250 bp have been developed (Meusnier et al., 2008).

There are several general problems of mitochondrial markers that must be considered:

*Nuclear mitochondrial pseudogenes (numts)*: Numts are non-functional copies of mitochondrial DNA (mtDNA) sequences that have been translocated into the nuclear genome (Bensasson et al., 2001; Lopez et al., 1994). They can be amplified with conserved universal primers aimed at mitochondrial copies and can complicate or confound analyses in various taxa (Bensasson et al., 2001; Buhay, 2009; Richly and Leister, 2004; Song et al., 2008).

*Wolbachia infections*: DNA barcoding studies usually assume a lower sequence variation within species than between species and a monophyly of mitochondrial DNA within species. Infections with maternally inherited symbionts can have direct influence on reducing the diversity of mtDNA and lead to identical mtDNA sequences

among different species and so disrupt this pattern and confound DNA barcode data (Hurst and Jiggins, 2005). Especially to mention is *Wolbachia*, an intracellular bacterium that is widely spread among insects (Hurst and Jiggins, 2005; Werren and Windsor, 2000; Werren et al., 1995). Detailed information about the impact of *Wolbachia* on DNA barcoding is given by Smith et al. (2012). *Wolbachia* infections have also been reported for Chrysomelidae and there are propositions for *Wolbachia*-mediated pest control and -management (Clark et al., 2001; Keller et al., 2004; Kondo et al., 2011; Roehrdanz et al., 2006; Werren and Windsor, 2000; Werren et al., 1995).

*Mitochondrial heteroplasmy*: The mixture of more than one type of mitochondrial genome within a single individual, and therefore the coamplification of different heteroplasmic copies of mtDNA, can confound species numbers and lead to artificial clades (Hebert et al., 2004; Hulcr et al., 2007; Rubinoff et al., 2006).

Other problems that are linked with single-gene approaches and can complicate DNA barcoding are introgression and hybridization as well as incomplete lineage sorting (Chase et al., 2005; Funk and Omland, 2003; Meyer and Paulay, 2005; Rosenberg and Tao, 2008).

#### DNA-extraction, amplification and sequencing

Total genomic DNA was extracted from one to three legs of each specimen, using the Qiagen DNeasy<sup>®</sup> BloodTissue Kit or Qiagen Biosprint 96BS following the manufacturers' protocol. COI (658 bp) was amplified with the primers LCO1490 and HCO2198, or with LCO and Nancy (for primer information see Tab. 2.2) using the Qiagen<sup>®</sup> Multiplex PCR Kit. Amplification reactions were carried out in a 20  $\mu$ l volume containing 10  $\mu$ l QIAGEN Multiplex PCR Mastermix, 2  $\mu$ l Q-Solution, 1.6  $\mu$ l of each primer (both 10 pmol/ $\mu$ l), and 2.5  $\mu$ l DNA template, and filled up to 20  $\mu$ l with sterile H<sub>2</sub>O. The PCR temperature profile consisted of an initial denaturation at 95°C (15 min), followed by 15 cycles at 94°C (35 s, denaturation), 55°C – 40°C (90 s, annealing temperature decreasing with every cycle about 1°C; Touch down-PCR), 72°C (90 s, extension), 25 cycles at 50°C annealing temperature, and a final extension at 72°C (10 min). Products were checked by electrophoresis on a 1.5% agarose gel containing GelRed<sup>TM</sup> (Biotium Inc.). Successfully amplified DNA fragments were purified using Illustra<sup>TM</sup> ExoStar (GE Healthcare) following the manufacturers' protocol. PCR products were sequenced in both directions by MacroGen Europe (Amsterdam, Netherlands; <http://www.macrogen.com>) using the same primers.

Table 2.2: Primer information.

Name	Sequence	Direction	Reference
LCO1490	5'- GGT CAA CAA ATC ATA AAG ATA TTG G -3'	forward	Folmer et al. (1994)
HCO 2198	5'- TAA ACT TCA GGG TGA CCA AAA AAT CA -3'	reverse	Folmer et al. (1994)
Nancy (C1-N-2191)	5'- CCC GGT AAA ATT AAA ATA TAA ACT TC -3'	reverse	Simon et al. (1994)

## 2.5 Analyses

All sequences were assembled and edited with Geneious version 5.4.4–version 7.1.5 (Biomatters Ltd.; <http://www.geneious.com/>). Detailed information about alignment of sequences and reconstruction of trees as well as programs used are given in the respective chapters. Statistical analyses were performed in R version 2.15.1–version 3.1.1. Detailed information is provided in the respective chapters. Data plottings were created in R, too, or with Microsoft Office Excel 2003–2010.

### Species delimitation

For biodiversity analyses or ecological studies based on invertebrate data, species richness and species turnover are important measures (Oliver and Beattie, 1996). However, the aim of this study is the evaluation of rapid methods for the assessment of an unknown diversity of leaf beetles without existing species information. As a surrogate for species, morphological and molecular working units that are fast to apply are used. They are supposed to be proxies for species and therefore be at species level, but they do not have to correspond exactly to species. It is not the aim to infer true species limits as it is e.g. the aim of integrative taxonomy.

### MORPHOSPECIES SORTING

As the aim of this study is the evaluation of rapid methods for biodiversity assessment, a thorough taxonomic analysis was relinquished and instead a parataxonomic morphospecies approach was used as it is common in tropical arthropod biodiversity studies. The dry mounted voucher specimens were sorted into morphospecies that were revised and verified by Dr. Thomas Wagner who is an experienced taxonomist for Chrysomelidae with afrotropical Galerucinae being his focus of expertise (see e.g. Wagner (2004, 2007a,b); Wagner and Kurtscheid (2005)). Specimens were sorted considering only external characters, without the use of dissected parts and without identification literature. Characters for morphospecies delimitation are shape of head, pronotum and total body, surface structures, and hairs or spines. Body size or colours may be used carefully considering that they may vary e.g. due to recent ecdysis. More information about morphospecies sorting and the concept of parataxonomy is given in Chapter 5. Morphospecies received a subfamily name and a number. It is to note that Hispinae and Cassidinae (meaning Cassidinae s.str.) are treated as separate subfamilies although they both belong to the subgroup Cassidinae (s.l.). The same applies to Alticinae and Galerucinae: For a better understanding the traditional view of two distinct subfamilies is retained although their status as subfamilies of equal rank and also their respective monophyly is in question. For information about the relations between Cassidinae and Hispinae as well as Alticinae and Galerucinae see Chapters 1.3 and 3.

For certain analyses of general diversity and comparison of sampling methods (Chapters 3 and 7) the remaining individuals of the preliminary morphospecies (*duplicate specimens*, see above '2.3 Further handling of the specimens') that have not been dry mounted and sequenced have been used, too (data sets 2a and 3a). They

have been assigned to the same morphospecies as the dry mounted and sequenced voucher specimen. It is to note that their classification to morphospecies is admittedly more superficial, but as the sorting of specimens of a single sample into preliminary morphospecies is quite reliable due to the small number of individuals and species per sample, it is likely that similar specimens of a sample were correctly classified as the same morphospecies. Furthermore, usually only one individual per morphospecies was found in one sample (~80%), in 12% there were two individuals and only in ~8% more than two (only in ten cases ten or more individuals of the same morphospecies were found in one sample). So the number of specimens affected is quite low and the error rate can be considered low (see also Chapter 7.3).

#### MOLECULAR SPECIES DELIMITATION

Four different molecular methods for species delimitation have been used:

*Statistical parsimony analysis* (Templeton, 2001; Templeton et al., 1992) as implemented in TCS v.1.21 (Clement et al., 2000) (95% connection limit) was used to group sequences into separate haplotype networks. The term network is used for all entities delimited by the program, also if they are no true networks or consist of only one sequence.

*Distance-based clustering* was based on the results obtained by SpeciesIdentifier v.1.7.7-dev3 (Meier et al., 2006) from the TaxonDNA package (<http://taxondna.sourceforge.net/>). The program generates clusters of sequences based on pairwise uncorrected distances at user-defined thresholds. All individuals that are connected directly to each other by distances below this threshold are grouped into a cluster (Meier et al., 2006).

*Generalized mixed Yule-coalescent (GMYC) modelling* (Monaghan et al., 2009; Pons et al., 2006) as implemented in the SPLITS package (<https://www.r-forge.r-project.org/projects/splits/>) for the R environment (R Development Core Team, 2009) was used to estimate species boundaries directly from the phylogenetic tree (Monaghan et al., 2009; Pons et al., 2006) produced with COI data. This procedure exploits the differences in the rate of lineage branching at the level of species and populations, recognizable as a sudden increase of apparent diversification rate when ultrametric node height (distance to tips) is plotted against the log number of nodes in a lineage-through-time plot (Nee et al., 1992).

*Poisson tree processes (PTP) modelling* was used as implemented on the PTP web server (<http://species.h-its.org/ptp/>) (Zhang et al., 2013). This method is similar to GMYC modelling but uses directly the number of substitutions (instead of the time) to identify branching rate transition points and therefore avoids the potentially error-prone process of making the tree ultrametric (Zhang et al., 2013).

The results of the different molecular species delimitation methods (networks, distance-, GMYC-, and PTP-clusters) are species-like units and often identical with species discerned by taxonomists. In this study they are summed up in the term molecular operational taxonomic units (MOTUs). The molecular species delimitation methods are described more detailed in Chapter 6.

## HAPLOTYPE DIVERSITY

Additionally, for the analyses in Chapters 4 and 6 haplotype diversity was inferred as a further independent measure for molecular diversity (Papadopoulou et al., 2011).

**Data sets**

Due to capacity restrictions not all collected specimens could be analysed, so a selection had to be made. Generally, only one (in some cases two or three) specimen of each preliminary morphospecies per sample was processed, i.e. assigned to a morphospecies and sequenced. For the different analyses, different data sets of the totality of processed specimens have been used. For some analyses also the duplicate specimens were included, for others only those specimens for that a DNA barcode could be obtained were used (Tab. 2.3):

*Data set 1* (data set of total analysed specimens): Consists of all specimens that have been processed (sorted into morphospecies and sequenced; 1775 specimens). It comprises data set 2 and several additional specimens of specific interest that were processed as well. All specimens have been assigned to a morphospecies and those with a barcode (*data set 1b*; 1475 specimens) to a haplotype network, too. Data set 1 was used for counting the total number of found morphospecies (Chapter 3) and for comparison of the sampling methods (Chapter 7). Data set 1b was used for the comparison of morphological and molecular species delimitation approaches (Chapter 5).

*Data set 2* (complete data set): Comprises specimens of 199 standardized plot samples (consisting of 199 sweep net, 199 beating, and 199 standardized hand-collection subsamples; 186 of the subsamples contained no chrysomelid specimen) and additional 65 non-standardized samples (hand-collection, light-, Malaise-, flight interception-, and pitfall-traps). All these samples have been analysed completely: From each sample, usually one (sometimes up to three) specimen per morphospecies has been processed. The data set 2 comprises all those processed individuals (1583 specimens). For some analyses, also the not processed specimens from these samples (duplicate specimens per morphospecies per sample) have been included (*data set 2a*; 2227 specimens), for others only those specimens with a barcode (*data set 2b*; 1334 specimens). Data sets 2, 2a, and 2b were used for general diversity analyses (Chapter 3), data set 2a also for sampling method analyses (Chapter 7).

*Data set 3* (plot data set): The data set is a subset of the complete data set. It is based only on the 199 standardized plot samples (consisting of 199 sweep net, 199 beating, and 199 standardized hand-collection subsamples). One (in exceptions up to three) specimen of each morphospecies of each subsample was processed (1200 specimens). It was used for comparison of subfamily composition between the different elevations (Chapter 3). For sampling method analyses (Chapter 7) also the not processed duplicate specimens per morphospecies per sample were included (*data set 3a*; 1578 specimens). The biodiversity analyses along the gradient in Chapter 4 were based on a data set with only those specimens for that a barcode could be obtained (*data set 3b*; 995 specimens).

*Data set 4* (preliminary data set): The data set was used for the comparison of



the different molecular species delimitation methods (Chapter 6). It contains 674 specimens that were sampled in November and December 2010 and from May 2011 until 11.08.2011. It is a preliminary data set containing all those specimens that were available at mid of August 2011 for that a barcode could be obtained until June 2012. Several specimens from that sampling time period have been chosen only later to be analysed and for some specimens a barcode could be obtained afterwards. They are not included in this data set. The data set includes four sequences that have been later on excluded from further analyses as they were in retrospect considered to be doubtful or probably contaminated.

Tab. D.1 in the Appendix lists for each specimen the data sets it was used for.

Table 2.3: Overview of the different data sets.

Name		Content	# Specimens
Data set 1	data set of total analysed specimens	all processed specimens (data set 2 + several additional specimens of specific interest)	1775
Data set 1b		all specimens of data set 1 with a barcode	1475
Data set 2	complete data set	specimens from 199 plot samples and 65 additional samples	1583
Data set 2a		all specimens of data set 2 plus duplicate specimens	2227
Data set 2b		all specimens of data set 2 with a barcode	1334
Data set 3	plot data set	specimens from 199 plot samples	1200
Data set 3a		all specimens of data set 3 plus duplicate specimens	1578
Data set 3b		specimens of data set 3 with a barcode	995
Data set 4	preliminary data set	all specimens that were available at mid of August 2011 for that a barcode could be obtained until June 2012	674



# General patterns of leaf beetle diversity

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## 3.1 Introduction

Despite an increasing effort in discovering Earth's biodiversity (Dirzo and Raven, 2003; Novotny and Miller, 2014) it is not yet ultimately known how many species of organisms there are. Estimates range from 3.6 to 100 million species, most probable the species number is an order-of-magnitude of ten million (Chapman, 2009; May, 2010; Mora et al., 2011; Wilson, 2003). Although it is clear that only a small fragment of species has been formally identified and named, there is still uncertainty about the exact numbers (1.4–2.2 million) (Chapman, 2009; Grove and Stork, 2000; Mora et al., 2011; Wilson, 2003).

The largest fraction of all species is made up by the tremendous but largely unexplored fauna of arthropods, predominantly insects, with ~1.1 million named distinct species (Chapman, 2009). The majority of them live in the most species-rich tropical ecosystems, especially the rainforests, where they contribute a large part of diversity in numbers of species as well as individuals (Corlett and Primack, 2011; Novotny et al., 2006). Therefore, in biodiversity research and global species richness estimates tropical arthropods, and especially tropical beetles, play a key role (Grove and Stork, 2000; Hamilton et al., 2010; May, 2010). Beetles are both functionally diverse and the most species-rich animal order, making up about one-quarter of all species on Earth (Hunt et al., 2007; Ødegaard, 2000). Since Erwin's (1982) spectacular estimation based on the number of beetle species associated with an individual tropical rainforest tree species, numerous studies led to widely varying estimates of global insect species numbers (Pimm et al., 1995; Stork, 1988, 1993). Those between four and six million arthropod species seem the most probable (Basset et al., 2012; Hamilton et al., 2010, 2013; Novotny et al., 2002b). Global and regional insect diversity estimates have often been plant-based estimates, i.e. the number of plant species is multiplied by the number of insect species that are effectively specialized to them (Erwin, 1982; Novotny and Miller, 2014). Therefore, especially herbivorous beetles, including Chrysomelidae, and their degree of host-specificity have played a fundamental role in species number estimates (Erwin, 1982; Ødegaard, 2000). Herbivorous insects are an extremely species-rich feeding guild and play essential functional roles in ecosystems (Coley and Barone (1996); Janzen (1970, 1987); Mattson and Addy (1975); Price (2002); see also Chapter 1.3).

Whereas research has focused mainly on lowland rainforests, tropical mountain

forests have received comparatively little attention (Beck et al., 2008a). And yet they are very exceptional habitats with an extraordinary flora and fauna that is extremely diverse, even more diverse than the tropical lowland rainforests (Beck and Kottke, 2008). The study area in the tropical Andes of southern Ecuador belongs to a biodiversity hotspot. Studies on e.g. geometrid moths, birds, vascular epiphytes, bryophytes, or orchids, revealed high species numbers for the respective groups (Brehm et al., 2008b). For beetles as for most other insect taxa there is still a gap and explicitly studies on Chrysomelidae are missing (Brehm et al., 2008b). For more information about biodiversity research in the study area see Chapters 1.4 and 2.1.

Data that can be adduced for comparison are scarce: The state of knowledge of leaf beetle diversity in Ecuador and in the Neotropics in general is poor (more detailed information about research on Neotropical Chrysomelidae is given in Chapter 1.4). Inventories or biodiversity studies on Neotropical leaf beetles are quite rare and often focus on single taxa (e.g. Flowers and Hanson (2003); Furth et al. (2003); Linzmeier and Ribeiro-Costa (2008, 2012, 2013); Staines (2011)). Especially their diversity in montane ecosystems has scarcely been studied (Furth, 2013; Sánchez-Reyes et al., 2014).

This study is a first attempt to assess the leaf beetle diversity of the herbaceous and shrubby understorey vegetation in the mountain forest of the Reserva Biológica San Francisco (RBSF) and Podocarpus National Park in southern Ecuador. Although it does provide neither a complete inventory nor a taxonomic checklist, it can serve as basis for future research on chrysomelid diversity.

## 3.2 Methods

All leaf beetles were sampled between November 2010 and June 2012 in parts of Podocarpus NP and RBSF, Ecuador (detailed information about the study area is given in Chapter 2.1). They were mainly collected by sweep netting, beating, and hand-collection of the lower vegetation. Additionally, light- and Malaise-traps have been used (detailed information about sampling methods and design is given in Chapters 2.2 and 7).

Due to capacity restrictions, not all collected specimens could be analysed but a selection had to be made: Usually the specimens of one sample were sorted into preliminary morphospecies and of each morphospecies per sample only one specimen (in some cases up to three) was selected for sequencing and final morphospecies assignment. However, for some analyses also the not selected specimens (*duplicate specimens*) were included (data set 2a). This procedure is described more detailed in Chapters 2.3 and 2.5. Laboratory analyses are described in Chapter 2.4, for information on morphospecies sorting please refer to Chapters 2.5 and 5.

For different analyses different subsets of specimens have been used; these are described detailed in Chapter 2.5 and summarized in Appendix Tab. D.1. The total number of morphospecies was counted for all processed specimens (data sets 1, 1b). For all obtained barcode sequences (1475 specimens, data set 1b), a statistical

parsimony analysis as implemented in TCS v.1.21 (95% connection limit) was used to group sequences into separate haplotype networks (more detailed information about statistical parsimony analysis is given in Chapters 2.5 and 6). For the further biodiversity analyses the data sets 2, 2a, and 2b were used. For comparisons between the elevational levels the plot data set (data sets 3, 3b) was used. Individuals of this data set with a barcode were grouped into PTP-clusters (Poisson tree processes (PTP) modelling is explained in Chapters 2.5 and 6). Elevational levels were also compared on the basis of haplotypes. The term MOTU (molecular operational taxonomic unit) in this chapter refers to the haplotype networks and PTP-clusters.

Species accumulation curves were calculated with R version 2.15.1 using the package VEGAN 2.0-10 (function *specaccum*). They show the increase in number of found species with growing number of sampled individuals or analysed samples. They can be used to visualize the completeness of the sampling. Species richness estimates were carried out with R and the VEGAN package using the function *specpool*. The function estimates the extrapolated species richness in a species pool. It is based on incidences in sample sites, and gives a single estimate for a collection of sample sites. In a collection of sample plots, many species will remain undetected. The function *specpool* uses some popular ways of estimating the number of the unseen species and adding them to the observed species richness: The variants of extrapolated richness in *specpool* are chao, first and second order jackknife, and bootstrap (Oksanen et al., 2013). These are widely used non-parametric estimators that consider information on the rare species in an assemblage to estimate the minimum number of species in the assemblage (Gotelli and Colwell, 2011). The methods have found to perform well in several comparative studies on species richness estimation (e.g. Colwell and Coddington (1994); Walther and Moore (2005); Walther and Morand (1998)). As different estimators are sensitive to the properties of the assemblage and sampling design, a set of different estimators was used to allow a range of estimates (Samways et al., 2010).

Morphospecies incidence (number of samples in which the morphospecies was found) and abundance (number of individuals per morphospecies collected) were divided into five categories: 1) uniques (morphospecies found in only one sample) respectively singletons (morphospecies of which only one individual occurred in the data set), 2) duplicates (single morphospecies found in two samples) respectively doubletons (single morphospecies of which two individuals were found in all samples), 3) rare morphospecies (found in 3–10 samples, respectively represented by 3–10 individuals), 4) common morphospecies (11–20 samples/individuals), and 5) very common morphospecies (>20 samples/individuals).

Data plottings were created in R 3.3.1 (using the packages VEGAN, RESHAPE2, GDATA, and MUMIN) or with Microsoft Office Excel 2003. For Figure 3.6, a NJ-Tree (Saitou and Nei, 1987) based on a MUSCLE alignment (Edgar, 2004) was constructed for representatives of each network of the plot data set 3b (370 sequences). For a better illustration branch lengths were made ultrametric with PATHd8 software (Britton et al., 2007) using relative ages of nodes and setting the root to an arbitrary age of 1. The Neighbor-Net of Cassidinae plus Hispinae (Fig. 3.10) was

constructed with SplitsTree v.4.12.3 (Huson and Bryant, 2006) using uncorrected p-distances.

### 3.3 Results

#### General Results

Within the complete project, 1174 samples have been taken resulting in a total of 4286 collected Chrysomelidae. Due to capacity restraints not all specimens could be analysed. A total of 1775 specimens (belonging to 515 morphospecies) have been processed (barcoded, mounted, and assorted to morphospecies) (data set 1). They comprise the specimens from the 662 samples that have been analysed completely and additional individuals of specific interest. For 1475 of these individuals (448 morphospecies, 493 MOTUs), barcoding has been successful (data set 1b). Selection of the specimens and the different data sets are explained in Chapters 2.3 and 2.5.

#### Found and estimated species richness

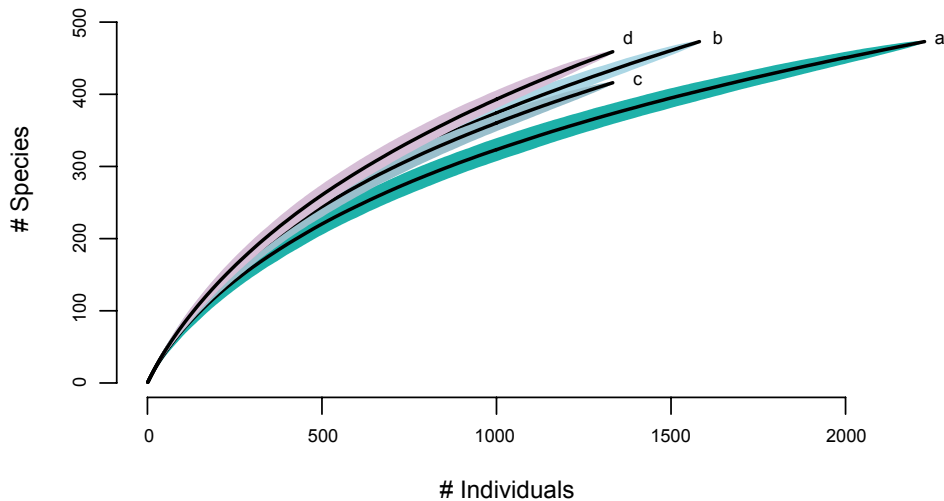


Figure 3.1: **Species accumulation curves (mean  $\pm$ 95% confidence interval) showing increase in number of found species with increasing number of sampled individuals.** Shown are curves for all processed specimens plus duplicate specimens (a; data set 2a), all processed specimens (b; data set 2), and all specimens with barcode (data set 2b; c: morphospecies curve, d: MOTU (haplotype network) curve).

For the following biodiversity analyses a data set was used consisting of all individuals from the 662 samples that have been analysed completely (data set 2a). It contained a total of 2227 specimens of which 1583 belonging to 473 morphospecies were processed (1–3 per preliminary morphospecies per sample) (data set 2). Barcoding has been successful for 1334 specimens (84.3%) belonging to 416 morphospecies and 459 MOTUs (data set 2b). Species accumulation curves did not reach saturation indicating that additional sampling would yield more species (Fig. 3.1). When the duplicate specimens were included (data set 2a; 2227 specimens), the curve levelled out but still did not reach saturation (Fig. 3.1a). It is to note that the inclusion of the duplicate specimens did not increase species number because duplicate specimens were assigned to the same morphospecies as the processed voucher specimen (see Chapters 2.3 and 2.5).

The expected total number of morphospecies estimated with the chao2 estimator was highest when calculated for all 1583 analysed specimens (915.53). When not processed duplicate specimens per morphospecies per sample were included it was slightly lower (905.07). When based on the specimens for which a barcode was obtained it was 705 morphospecies, compared to an estimated MOTU number of 804.72. Species numbers found and estimated by different estimators are given in Table 3.1.

Table 3.1: **Species numbers found and estimated by different estimators for data sets 2, 2a, and 2b.** Data set 2a: all processed individuals plus duplicate specimens, data set 2: processed individuals, data set 2b: individuals with barcode.

	Specimens	Found Species	Estimated Morphospecies Numbers				n Samples
			chao	jack1	jack2	boot	
Data set 2a	2227	473	905.07±80.34	709.5±24.48	880.92	572.18±12.98	477
Data set 2	1583	473	915.53±82.42	710.5±24.58	883.91	572.42±12.96	477
Data set 2b	1334	416	705±55.56	619.55±20.75	751.13	503.46±11.35	454
		MOTUs	Estimated MOTU Numbers				
Data set 2b	1334	459	804.72±61.26	699.47±23.31	855.96	561.44±12.4	454

### Incidence and abundance of morphospecies

The studied leaf beetle community shows an uneven distribution of incidence and abundance with very few common morphospecies while the vast majority is rare. Half of all found morphospecies (237) were collected in only one sample (uniques), and 14% in two samples (duplicates) (Fig. 3.2). Almost one third were rare (found in three to ten samples). The proportion of common and very common species was very low (three, respectively two percent). A similar pattern is visible for the abundance of morphospecies, where ten percent of the morphospecies accounted for 52% of the individuals: The number of singletons (213) is slightly lower than the number of uniques making up 45% of all morphospecies. Proportion of doubletons and common morphospecies remains almost equal, whereas common and very common morphospecies was slightly higher with one morphospecies (*Alticinae* sp. 118)

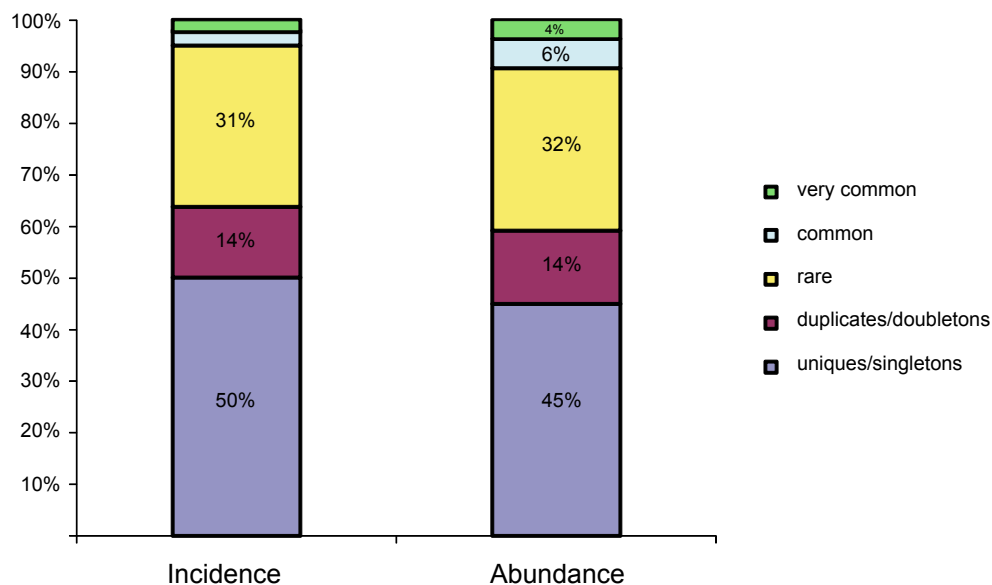


Figure 3.2: **Proportion of uniques/singletons, duplicates/doubletons, rare, common, and very common morphospecies.** Incidence: number of samples in which the morphospecies was found, abundance: number of individuals per morphospecies collected.

showing an extremely distant number of 136 found individuals.

Figure 3.3A is another illustration of the incidence of morphospecies showing that the vast majority was collected in very few samples. A similar pattern is visualized for the abundance of morphospecies (Fig. 3.3B): Most morphospecies were represented by very few individuals.

When the increase in singletons is plotted against the number of individuals, the curve shows a steep incline without any sign of approaching saturation (Fig. 3.4). Therefore, adding further specimens will increase the number of singletons.

Figure 3.5 shows the incidence and abundance of the most frequent, respectively most abundant morphospecies. Of the 19 most abundant morphospecies, 17 were also among the 19 most frequently found ones. Of the most frequently found morphospecies, 14 belonged to Alticinae, four to Eumolpinae, and one to Galerucinae. Of the most abundant morphospecies, 13 belonged to Alticinae, (the same) four to Eumolpinae, and two to Galerucinae. Of the 237 uniques, 213 are singletons. Those uniques that are no singletons are represented by two (in 19 cases), three (in three cases), or four (in two cases) individuals. All morphospecies with five or more specimens were found in at least two samples.

The similar results for incidence and abundance arose from the fact that morphospecies were usually represented by one or few individuals in one sample (one individual: 78%, two individuals: 13%, 3–10 individuals: 8%, >11 individuals: <1%). So, usually a high incidence accounts for the high abundance of morphospecies.

Fig. 3.6 illustrates the incidence of species on the different sampling sites for data set 3b.



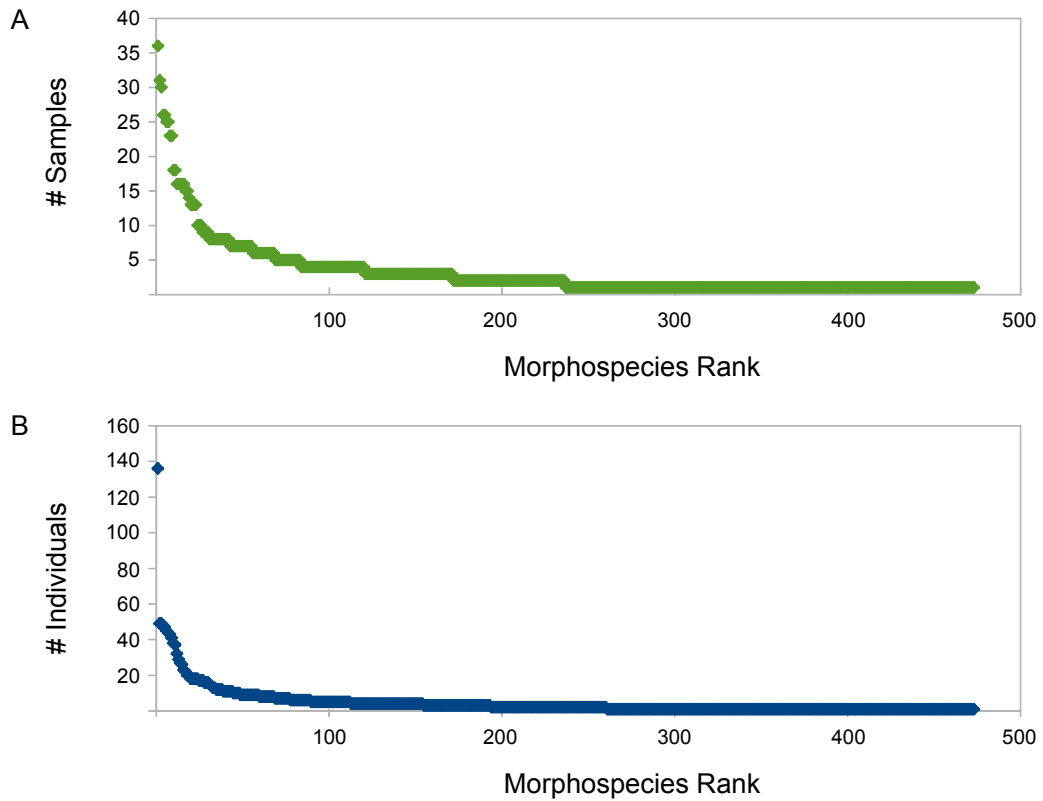


Figure 3.3: **Incidence (A) and abundance (B) of morphospecies.**

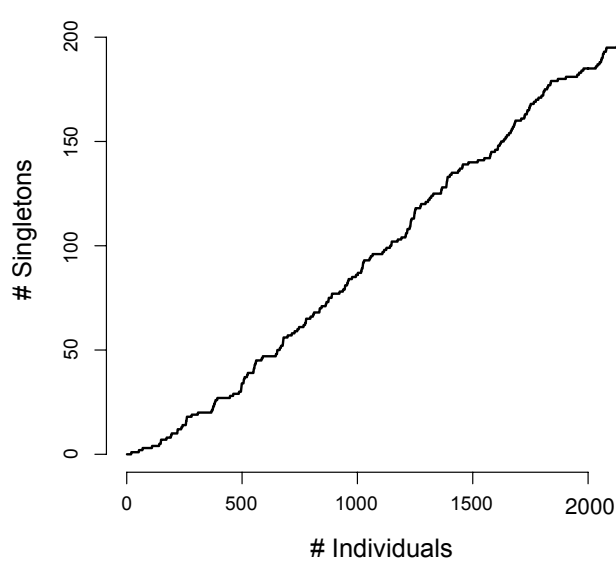


Figure 3.4: **Singleton curve.** The curve shows the increase in singletons with increasing number of individuals.

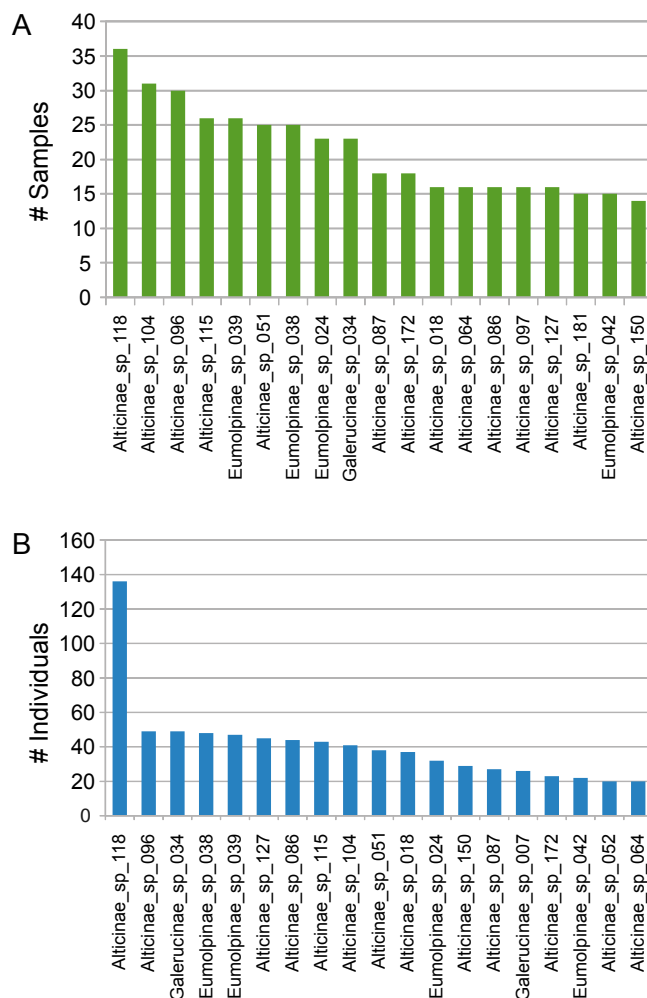


Figure 3.5: Incidence (A) and abundance (B) for the most frequent, respectively abundant morphospecies.

The majority of morphospecies was found at only one elevation (91%) and no morphospecies was found at all three elevations (Fig. 3.7). This stenoecy is even more pronounced for MOTUs (PTP-clusters; 96%) and haplotypes (99%). Only three haplotypes were found in Bombuscaro and also ECSF. If morphospecies, MOTUs, and haplotypes that occur only once in the data set are removed, the proportion of species/haplotypes restricted to one elevation slightly decreases, however still remains the majority (82%, 91%, 98%). Of the species/haplotypes that occur on two elevations, the majority was found in Bombuscaro and ECSF (83% of morphospecies, 93% of MOTUs, and 100% of haplotypes), the others in ECSF and Cajanuma. Only one morphospecies (Alticinae sp. 095), but no MOTUs or haplotypes occur in Bombuscaro and Cajanuma. Probably the two specimens have been erroneously assigned to the same morphospecies. Cajanuma has the highest proportion of species/haplotypes that were found only there: 90% of morphospecies, 98%

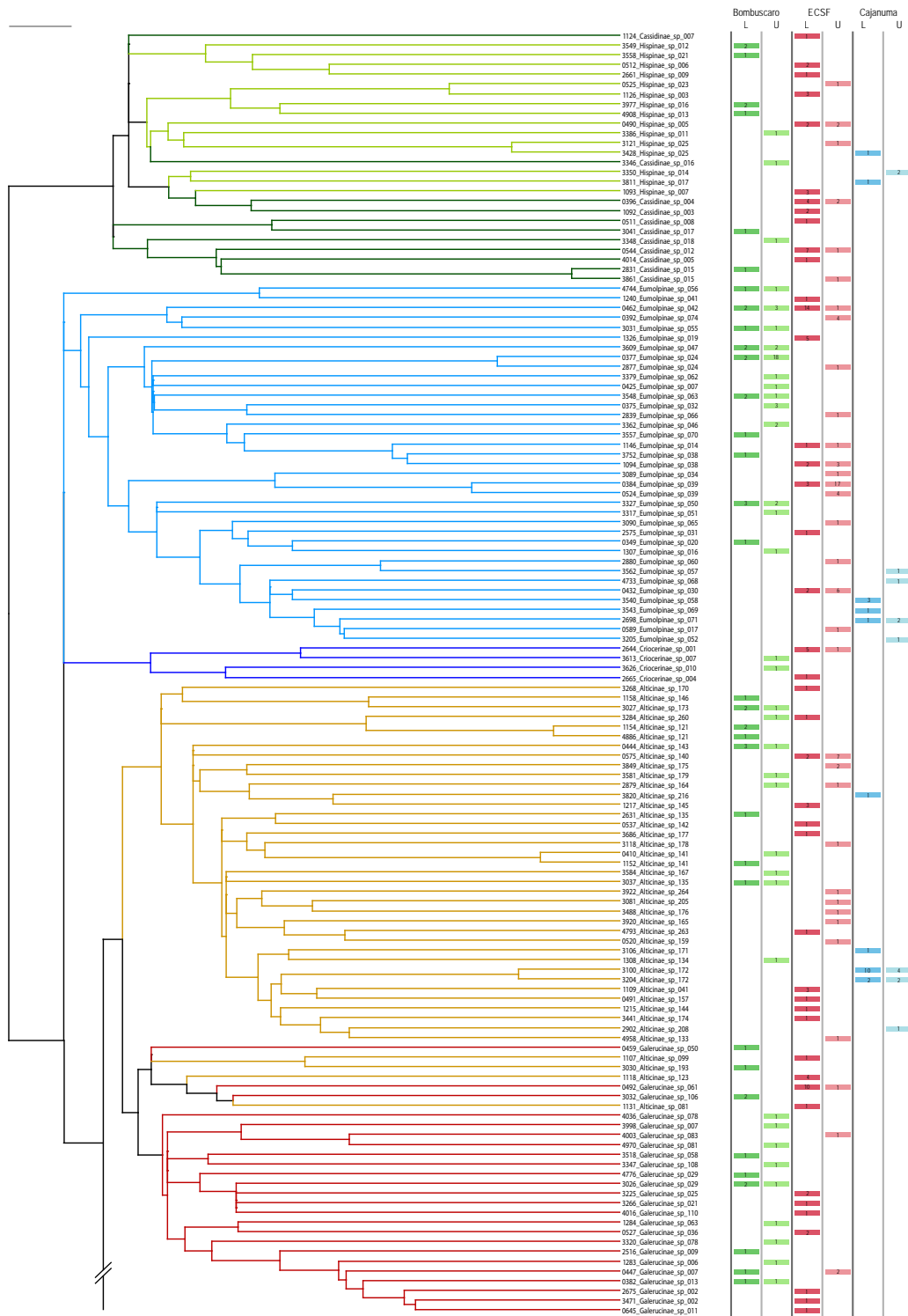
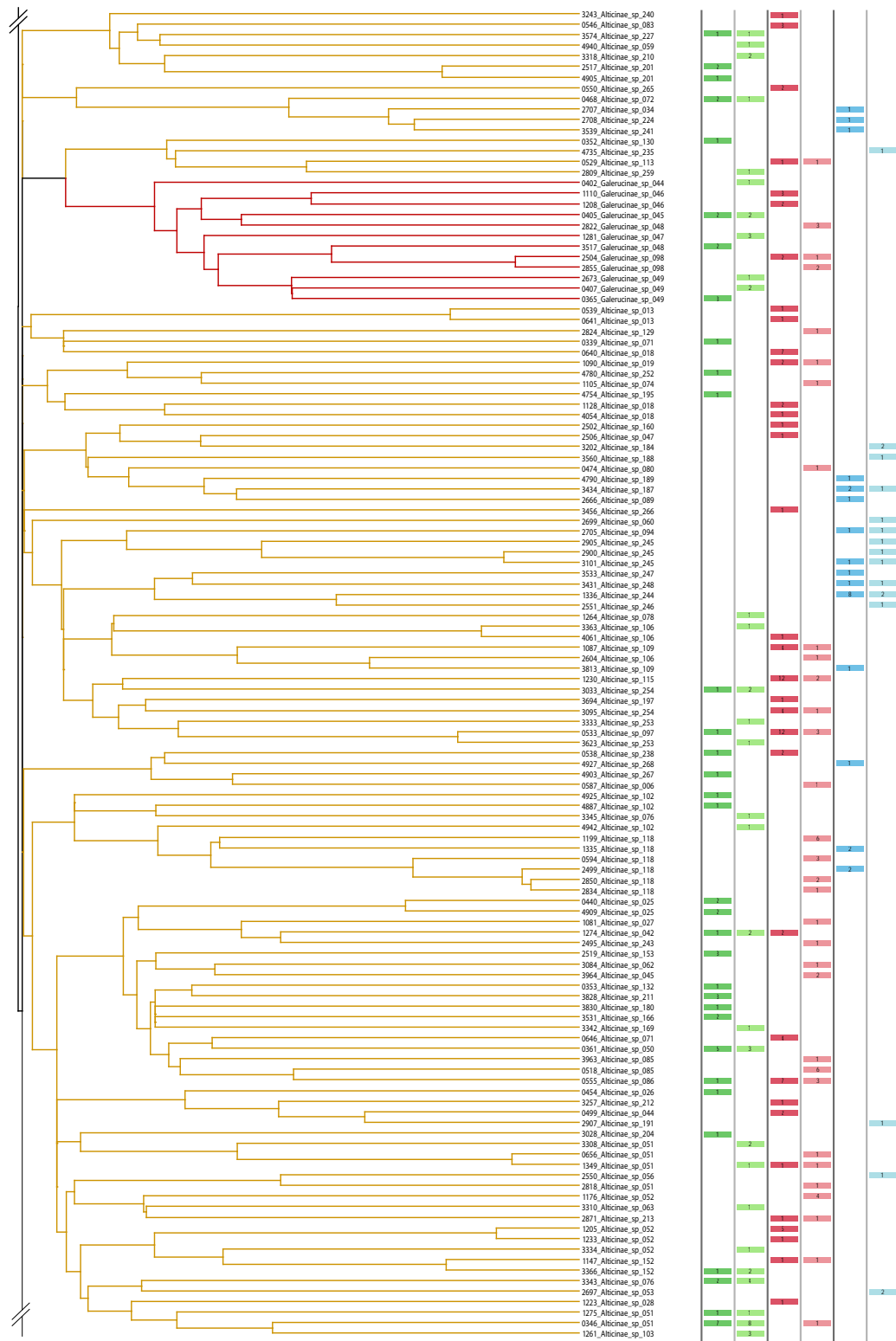
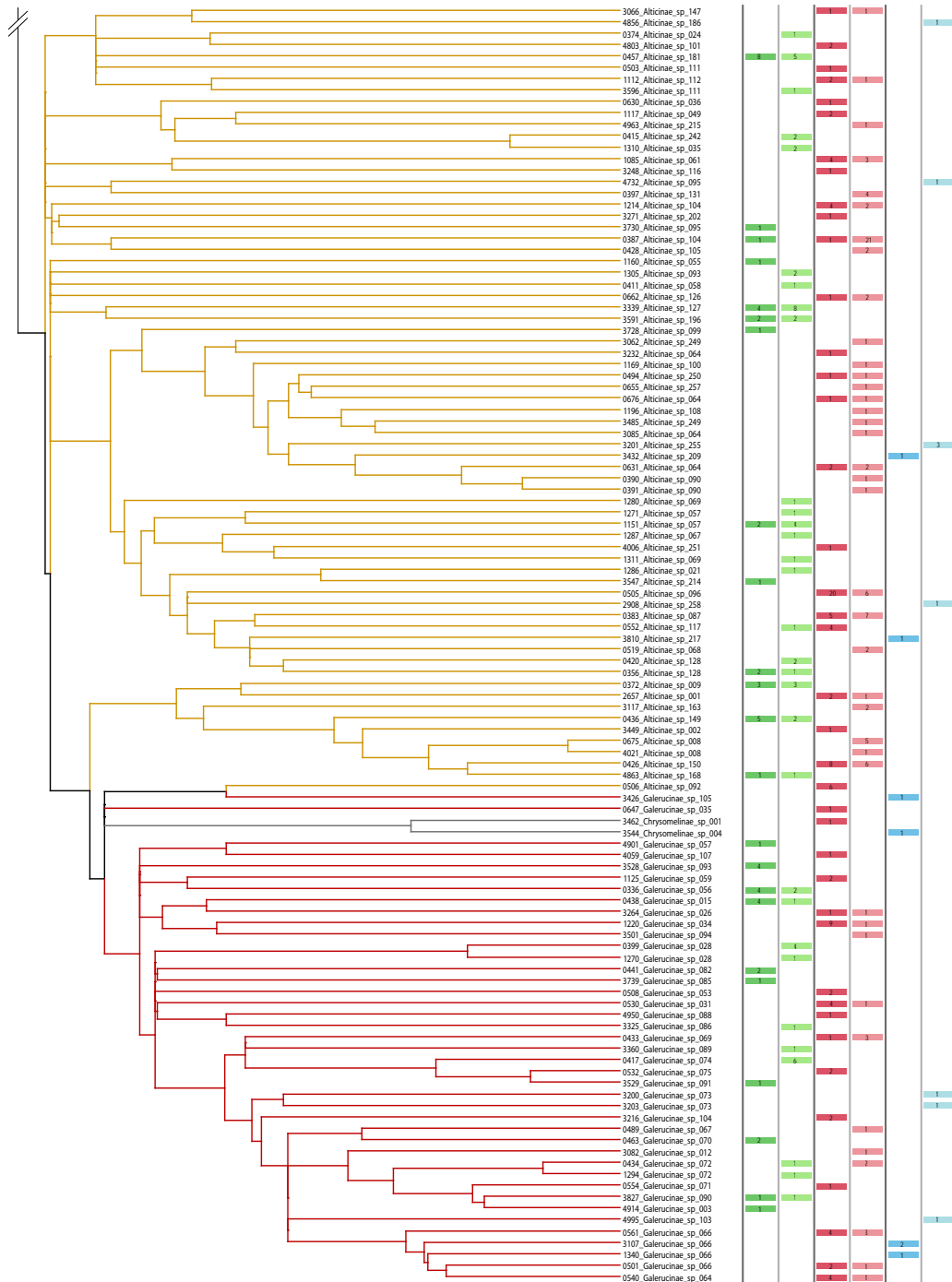


Figure 3.6: NJ-tree of one representative of each MOTU (haplotype network) with incidence of MOTUs. Of each MOTU one specimen was chosen; its voucher-number and its morphospecies name is given as this is rather informative than the MOTU name. Coloured boxes indicate elevational area (Bombuscaro, 1000 m; ECSF, 2000 m; Cajanuma, 3000 m) and habitat (L= 'lower plot', valley; U= 'upper plot', ridge) where the MOTU was found, numbers therein indicate in how many samples (sweep netting, beating, hand-collection) the respective MOTU was found. Based on plot data set 3b.





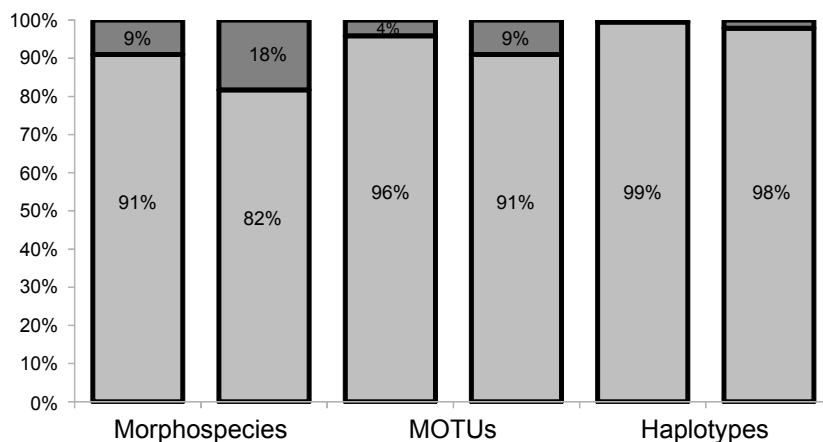


Figure 3.7: **Proportion of morphospecies, MOTUs (PTP-clusters) and haplotypes found at one (light grey), respectively two (dark grey) elevations.** Shown for data set 3b. Left column: complete data set 3b, right column: without morphospecies, MOTUs, and haplotypes that occur only once in the data set.

of MOTUs and all haplotypes found at Cajanuma were found exclusively there. For ECSF (83%, 91%, 99%) and Bombuscaro (82%, 91%, 99%) the percentage was very similar.

### Species richness and abundance of subgroups

The 2227 specimens of the complete data set for biodiversity analyses (data set 2a) belonged to nine subfamilies (Galerucinae s.str., Alticinae, Cassidinae s.str. and Hispinae are regarded as separate subfamilies, see Chapter 1.3): Alticinae, Galerucinae, Eumolpinae, Cassidinae, Hispinae, Criocerinae, Chrysomelinae, Lamprosomatinae, and Cryptocephalinae. The numbers of found individuals and morphospecies are given in Table 3.2.

Table 3.2: **Numbers of found specimens and morphospecies for subfamilies.**

	Morphospecies	Individuals
Alticinae	251	1326
Galerucinae	99	433
Eumolpinae	68	340
Cassidinae	19	65
Hispinae	22	38
Criocerinae	8	19
Chrysomelinae	3	3
Lamprosomatinae	2	2
Cryptocephalinae	1	1
total	473	2227

Alticinae showed the highest species richness and abundance, accounting for more than half of all found morphospecies (53%) as well as individuals (60%; Fig. 3.8). The second species- and individual-rich subgroup was Galerucinae (21% of morphospecies, 19% of individuals). Eumolpinae represented 14% of all found morphospecies and 15% of all found individuals, Cassidinae accounted for four percent, respectively three percent, and Hispinae for five percent, respectively two percent. Criocerinae, Chrysomelinae, Lamprosomatinae, and Cryptocephalinae together accounted for three percent, respectively one percent.

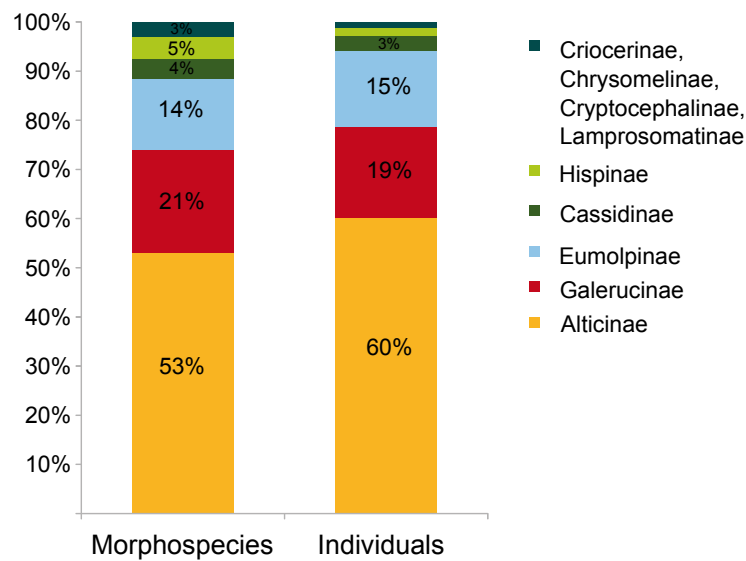


Figure 3.8: **Composition of Chrysomelidae subfamilies.** The proportion of the different subfamilies is shown in terms of morphospecies and number of individuals.

When the proportion of the different subfamilies in the total number of morphospecies is regarded for each altitude separately (for the plot data set 3), the rank order remains the same (Fig. 3.9). However, there are small differences among the elevations: Whereas the proportion of Alticinae plus Galerucinae remains similar (77% at Bombuscaro and ECSF, 74% at Cajanuma), there is a shift towards Alticinae with increasing elevation (51% at Bombuscaro, 59% at ECSF, 68% at Cajanuma). At Bombuscaro and ECSF, there are almost equal portions of Hispinae and Cassidinae; in contrast, at Cajanuma Cassidinae are not found at all.

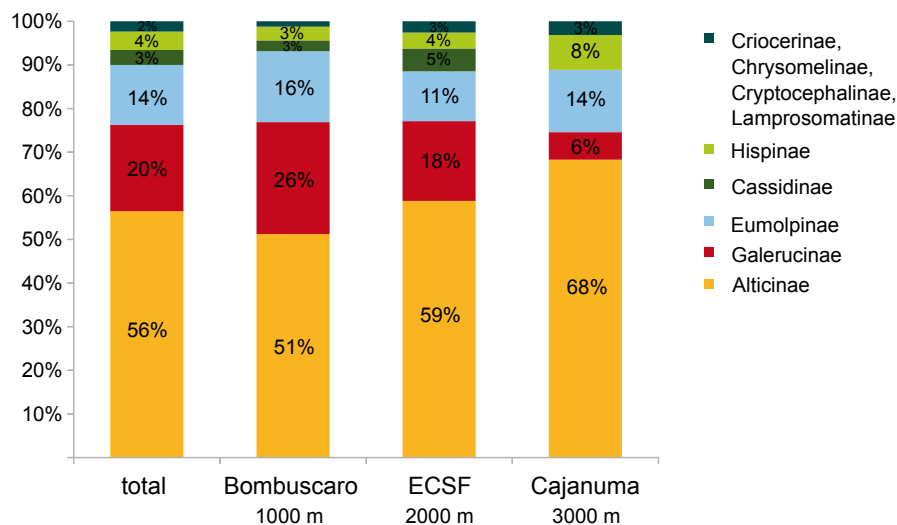


Figure 3.9: Proportion of subfamilies in number of morphospecies for the different elevations.

It is to note that for the most part barcoding grouped the morphospecies correctly into subfamilies (see Fig. 3.6). Cassidinae + Hispinae, Eumolpinae, and Criocerinae formed clusters each. Also Alticinae + Galerucinae formed a cluster but with Chrysomelinae placed within. With single exceptions, within Galerucinae s.l., Galerucinae s.str. and Alticinae formed several well distinguished clusters. Also within the subfamily-clusters, COI-sequences form clearly distinguishable clusters (as an example see Fig. 3.10).

### 3.4 Discussion

#### Species richness

So far, there exist no published records of leaf beetle species for the studied area. The Chrysomelidae found under tree bark in RBSF by J. Schmidl (2007–2008) were sorted into ~50 morphospecies, but not identified taxonomically. The only Chrysomelidae checklist for Ecuador (and whole Central and South America) by Blackwelder (1947) is outdated. Since then, numerous species have been newly described from Ecuador (Borowiec, 1998, 2000a,b; Flowers, 2009a,b; Sekerka and Windsor, 2012; Staines and Zamorano, 2012; Świętojańska and Borowiec, 2000), but only for Cassidinae a more recent checklist has been published (Borowiec, 1998).

Blackwelder lists ~450 chrysomelid species explicitly for Ecuador, a number even below the number of morphospecies found in this rather small-scale study. In this study, 515 morphospecies have been found, and one has to keep in mind that due to capacity restraints not all of the collected specimens have been assigned to morphospecies. The not processed specimens are likely to entail more not yet identified morphospecies. These numbers illustrate the striking discrepancy between the de-



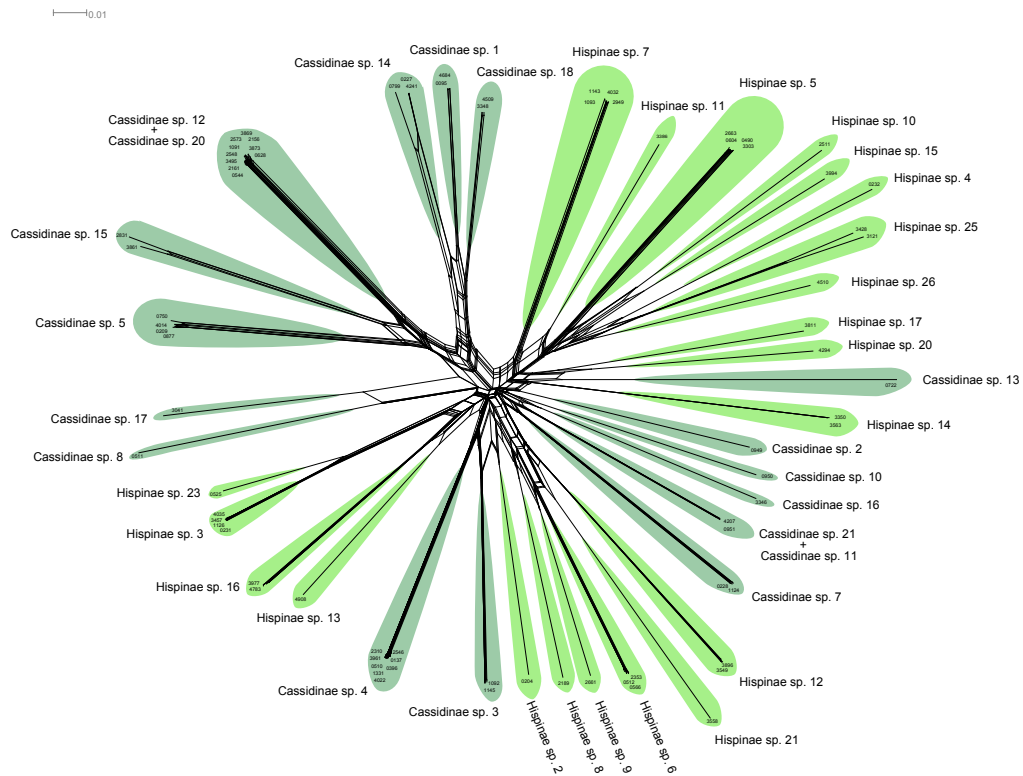


Figure 3.10: **Neighbor-Net of Cassidinae plus Hispinae.** Visualization of clustering within Cassidinae + Hispinae showing that information content of the data is useful to discriminate species but not for phylogenies.

scribed and the true Ecuadorian leaf beetle diversity. That Ecuador still provides much unknown diversity to be discovered is also reflected by its collections: The Invertebrate Section of the Museum of Zoology at the Pontifical Catholic University of Ecuador in Quito comprises with almost two million specimens Ecuador's largest collection of native taxa. It harbours over 24,000 Chrysomelidae, most of them still awaiting identification: Only ~11% of all specimens are determined to species, ~14% to genus, but ~75% have no identification at all (Clifford Keil, pers. comm.). There are only nine chrysomelid type specimens deposited in the collection, all of them belonging to Cassidinae (Donoso et al., 2009). In most other Neotropical countries the situation is similar (see Chapter 1.4).

The present study focused on the analysis of leaf beetle biodiversity and its changes along an altitudinal gradient. It was not attempted to create a complete inventory. The species accumulation curves indicate that additional sampling would further increase the number of found morphospecies. Estimated morphospecies numbers even range up to more than 900 (chao). It is to note that analyses of assemblages with low evenness (few dominating and many rare species) tend to underestimate species richness (Magurran, 2004). The number of found MOTUs for those individuals with a DNA barcode was even higher than the respective number of

morphospecies.

As this is one of the first known studies of site-specific data on leaf beetle richness and diversity for Ecuador there are no published data with which the present results can be compared. Studies of other Neotropical regions are difficult to compare due to differences in geographical scale, sampling effort, and methods. Some of them focus on certain subfamilies or study different habitats (e.g. Furth et al. (2003); Sánchez-Reyes et al. (2014)). If a careful comparison is still attempted, and considering that sampling in this study is far from being complete, species numbers seem comparable to or rather higher than in other Neotropical regions (Charles and Bassett, 2005; Flowers and Hanson, 2003; Furth et al., 2003; Linzmeier and Ribeiro-Costa, 2009, 2011, 2012; Ødegaard, 2006; Sánchez-Reyes et al., 2014).

It is to note that in this study mainly a selected habitat was investigated, the herbaceous and shrubby understorey vegetation. The canopy that is considered to be the most diverse habitat in tropical rainforests and harbouring an especially high leaf beetle diversity (Basset et al., 2001; Charles and Bassett, 2005; Farrell and Erwin, 1988) was completely neglected. Also the inclusion of particular habitats as the tree bark that was studied by Schmidl (2007–2008, pers. comm.) could further increase species number. Studies that include the rainforest canopy or are part of large-scale studies and inventories are likely to yield comparatively higher species numbers (e.g. Farrell and Erwin (1988) who found >650 species in a Peruvian rainforest canopy). Large-scale research programmes (e.g. IBISCA (<http://www.ibisca.net/>): Basset et al. (2012, 2007); Basset and Leponce (2005), ALAS (<http://viceroy.eeb.uconn.edu/ALAS/>): e.g. Furth et al. (2003); Staines (2011)) are capable of more intense sampling due to a much longer available time period and more manpower compared to the present study (Basset et al., 2007; Staines, 2011). They often include a comprehensive set of sampling methods (Basset et al., 2007; Longino and Colwell, 1997). For certain taxa they can aim at complete inventories, e.g. the hispine species at La Selva, Costa Rica, resulting in a quite impressive number of 139 species (Staines, 2011).

In the studied area a high leaf beetle diversity had been expected. The region is part of the Tropical Andes, a biodiversity hotspot, and especially known for its outstanding rich plant diversity (see Chapter 2.1). Particularly the diversity of herbivorous insects is closely tied to plant diversity and can be expected to be especially rich in regions with diverse vegetation. The diversity of the moth family Geometridae has been intensely studied in the study area (Bodner et al., 2010; Brehm and Fiedler, 2003, 2004, 2005; Brehm et al., 2003a, 2013, 2003b; Hilt et al., 2007; Strutzenberger et al., 2011) and found to be much higher than anywhere else in the world documented (Brehm et al., 2005). Although the leaf beetle diversity seems comparatively high in the region, a more complete sampling should be attempted to allow more thorough propositions about the true species numbers for the area.

### **Abundance and incidence of morphospecies**

Abundance and incidence of the studied leaf beetle assemblage showed a pattern with

a few common species and an overabundance of rare species that is characteristic for many taxa of tropical rainforests and was also found for Neotropical leaf beetles (Furth et al., 2003; Linzmeier and Ribeiro-Costa, 2012; Sánchez-Reyes et al., 2014). A low evenness delays the saturation of the species accumulation curve (Magurran, 2004). Whereas 'rare species' are often defined as those species at the lower end of the distribution of species abundance (the cut-off often placed at the first quartile; Magurran (2004)), the term is also used synonymously with singletons (species represented by a single individual in the data set) (Novotny and Basset, 2000). The latter often prevent the species accumulation curves from attaining an asymptote even in much larger-scaled studies than the present one (Novotny and Basset, 2000). Usually, additional sampling turns some singletons into doubletons, but new singletons appear (Lim et al., 2012). In the present study, half of all morphospecies were sampled only once (uniques), most of them were represented by only one individual in the whole data set (singletons; 45% of all morphospecies). This proportion is comparable to many other studies of rainforest arthropods, where the proportion of singletons is on average 32% (Coddington et al., 2009) and often up to half and more (Allison et al., 1993; Erwin, 1997; Janzen and Schoener, 1968).

A high proportion of singletons often results from undersampling even in very large and ambitious tropical arthropod surveys (Coddington et al. (2009) and references therein). Although there are many genuinely rare species in the tropics, most are not as rare as they seem: Sampling flaws can make species appear rare when they were sampled in marginal times or places (insufficient seasonal or spatial replication; Novotny and Basset (2000)) or with inadequate methods (Longino et al., 2002). Rare species in a sample might also be common elsewhere, e.g. in adjacent regions or well-known from collections or literature (Longino et al., 2002). In host plant based surveys of herbivorous insects, many rare species are transient species or 'tourists' that do not use the studied plant as host plant but rather as a site for resting, sun-basking or sexual display (Moran and Southwood, 1982; Novotny and Basset, 2000; Ødegaard, 2004). However, feeding and rearing trials showed that many rare species are indeed associated with the studied hosts (Novotny and Basset, 2000). They might be generalists feeding occasionally on the host examined but with relatively high overall population levels when all potential host plant species are considered or specialists feeding on the host examined but preferring, and being more abundant on other hosts (Novotny and Basset, 2000). Probably few are specialists with genuinely low population levels (Novotny and Basset, 2000).

The circumstance that the assemblage in this study is clearly undersampled (as the species accumulation curves show) is the most likely explanation for the high percentage of singletons, rather than any more biological explanation (Coddington et al., 2009). With additional sampling effort, the percentage of singletons might decrease, but is expected to remain quite high. Biological reasons for singletons in this study cannot be evaluated as insect-plant-relationship is not addressed. Furthermore, species are not described and therefore there is no information about their distribution outside of the study area.

The finding that most morphospecies, MOTUs, and haplotypes were found exclusively at one elevational level, whereas none was found at all three elevations, indicates ecological specialization and the presence of different leaf beetle communities at the different sites. Also the differences in the composition of subfamilies at different levels confirm this. It is not surprising that the three elevations harbour a different leaf beetle fauna: Although the three sites are as close as ~20 km, there are 1000 m elevation difference and the areas exhibit remarkable differences in climate and vegetation. The turnover of tropical insect communities along elevational gradients is generally rapid (Brühl et al., 1999; Ghalambor et al., 2006; Janzen, 1967) and there are often large differences in insect communities in considerably smaller ranges than 1000 m (e.g. Olson (1994); Smith et al. (2014)). For a detailed analysis of the change of communities with increasing altitude see Chapter 4.

### **Methodological considerations**

DNA barcode data generally revealed similar results as morphospecies data. However, there are certain advantages and disadvantages of each method that should be considered: When morphospecies and MOTU richness was compared, for those specimens for that a barcode was obtained (data set 2b), found and estimated MOTU numbers were higher than morphospecies numbers. The morphospecies sorting probably overlooks differences between species ('cryptic diversity') and is likely to underestimate true species richness. However, there are specimens where sequences could not be obtained at all or could not be used as they were contaminated, too short, or of insufficient quality. In contrast, morphospecies can be determined for every specimen (as long it is not severely damaged). Those specimens for that no barcode could be obtained can be assigned to a morphospecies and do not have to be excluded from the data set. If they are included, found and estimated morphospecies richness is higher than MOTU richness.

Aside from those specimens where barcode generation failed, often not all sampled specimens can be barcoded: Temporal or financial restrictions usually require a selection (this might not apply in metabarcoding studies). In the present study, usually only one specimen (sometimes up to three specimens) of each morphospecies per sample was sequenced leading to a number of 644 specimens (duplicate specimens) for which no barcodes were produced. Therefore, in this case, morphospecies can provide abundance data in contrast to barcode data. Although this is actually additional data that the barcode data cannot provide, there are no severe differences between inclusion or exclusion of the duplicate specimens, i.e. incidence and abundance data was very similar. For example, of the 19 most abundant morphospecies, 17 were also among the 19 most incident ones. Furthermore, 90% of the uniques are also singletons. This supports the decision to select one morphospecies per sample for sequencing, at least when, like in this study, usually only few individuals of a morphospecies occur in one sample.

### **Species richness and abundance of subgroups**

It is to note that for a better understanding Hispinae and Cassidinae are treated

as separate subgroups although they both belong to the subfamily Cassidinae (s.l.). The same applies to Alticinae and Galerucinae: The traditional view of two distinct subgroups is retained although their status as subfamilies of equal rank and also their respective monophyly is in question. For information about the classification and relations of Cassidinae and Hispinae as well as Alticinae and Galerucinae see Chapter 1.3.

In the present study Galerucinae s.l. make up the largest fraction of found leaf beetles in terms of morphospecies as well as individuals. This is in accordance with subfamily composition worldwide (Chaboo, 2007; Reid and Beatson, 2013) and other studies on Chrysomelidae in different regions (Kalaichelvan et al., 2005; Linzmeier and Ribeiro-Costa, 2012; Sánchez-Reyes et al., 2014; Wagner, 1999). Especially Alticinae, the largest leaf beetle subfamily (Furth et al., 2003), are often extremely abundant and species-rich (Farrell and Erwin, 1988; Flowers and Hanson, 2003; Freund, 2005; Linzmeier and Ribeiro-Costa, 2012; Wagner, 2003). In this study they accounted for more than half of all individuals and morphospecies at each elevational level. The Neotropical region harbours the most diverse Alticinae fauna in terms of genera and species (Scherer, 1988). Almost half of the known alticine genera occur in the Neotropics (Furth, 2005). More than 200 genera are known from South America, compared to ~40 Nearctic genera and ~65 genera in Africa (Furth, 2005; Scherer, 1988). All of the South American alticine genera are endemic, except *Chaetocnema*, *Epitrix*, *Longitarsus*, and *Terpnochlorus* (Scherer, 1988).

After Alticinae and Galerucinae s.str., the next most abundant and species-rich subfamily in this study was Eumolpinae. This was reported likewise in other studies on Neotropical leaf beetles (Farrell and Erwin, 1988; Linzmeier and Ribeiro-Costa, 2012), however not in the study of Sánchez-Reyes et al. (2014) on leaf beetle assemblages in forest and thorny scrub vegetation, who found Cassidinae more abundant and species-rich. Cassidinae and Hispinae are a dominant element of the Neotropical region, distinguishing it from any other area of the world (Kimoto, 1988). Worldwide, Cassidinae s.l. is the second largest subfamily of Chrysomelidae with 324 genera and ~6000 species (Chaboo, 2007). There is little overlap between Old World and New World cassidine fauna (Chaboo, 2007). Despite the high Cassidinae and Hispinae diversity in the Neotropics, they make up only a small portion in this study, as in studies using canopy fogging in lowland forest or Malaise trapping (Farrell and Erwin, 1988; Flowers and Hanson, 2003). This is probably due to the used methods as different sampling methods are more efficient for certain subfamilies. The composition of subfamilies depends on the sampling methods and a focus on other sampling methods could alter the composition of subfamilies. For example, Cassidinae are reluctant flyers that are mainly collected by hand-collection, whereas light trapping seems to be suited especially for Galerucinae (see Chapter 7).

The comparison of subfamily composition between the different elevations showed an increasing proportion of Alticinae at the expense of Galerucinae. A dominance of Alticinae with increasing elevation was also observed by Flowers and Hanson (2003) along an elevational gradient in Costa Rica. Alticinae and Galerucinae in tropical mountains can survive up to elevations where the vegetation ends (Jolivet and

Hawkeswood, 1995). Notably Alticinae are known to occur at very high altitudes in the Andes as well as on the Venezuelan Tepuys (Jolivet and Hawkeswood, 1995). The fact that most fern-feeding leaf beetles are found among Alticinae could be related to the scarcity of other suitable food plants at high altitudes (Jolivet and Hawkeswood, 1995).

### Phylogenetic considerations

It is well-known that COI has only limited information content at deeper phylogenetic levels (Moritz and Cicero, 2004) and it is not the purpose of this study to infer a phylogeny of leaf beetles. However, it is to note that for the most part DNA barcoding grouped the morphospecies correctly into subfamilies.

Eumolpinae and Criocerinae form monophyletic clusters. Both are taxonomically quite well defined subfamilies considered to be monophyletic (Duckett et al. (2004); Gómez-Zurita et al. (2005); Jolivet and Verma (2008); Matsumura et al. (2014); Reid (1995); Schmitt (1985a,b); but see Gómez-Zurita et al. (2007, 2008) who recovered Eumolpinae as paraphyletic). Also Cassidinae plus Hispinae form a monophyletic cluster that is consistent with current taxonomy. Both taxa are placed by many authors into one subfamily, the Cassidinae (Borowiec (1995); Chaboo (2007); Staines (2002); = Hispinae sensu Reid (1995)). Morphologically and biologically there seems to be no valid reason for retaining Hispinae and Cassidinae as separate subfamilies and there exist intermediate genera (e.g. in the tribes Cephaloleiini, Imatidiini) (Borowiec, 1995; Staines, 2002). At least three morphospecies in this study seem to belong to these taxa: Cassidinae spp. 8 and 17 probably belong to the genus *Imatidium* FABRICIUS (presumably the species *I. buckleyi* SPAETH, respectively *I. thoracicum* FABRICIUS, both known from Ecuador). Morphospecies Hispinae sp. 10 could belong to the genus *Demotispa* BALY. *Imatidium* and *Demotispa* belong to the tribe Imatidiini that has been synonymized with Cephaloleiini that have been traditionally classed with hispines (Borowiec, 1995; Staines, 2002). The genus *Imatidium* has been placed in hispines as well as in cassidines at one time or another (Staines, 2002). For further information about classification and relationships of Cassidinae and Hispinae see Chapter 1.3.

Also Alticinae plus Galerucinae form a monophyletic cluster, however with Chrysomelinae placed as a monophyletic cluster within. With single exceptions, within Galerucinae s.l., the Galerucinae s.str. and Alticinae appear in several well separated clusters. Whereas the monophyly of the group Galerucinae plus Alticinae is generally acknowledged, often subsuming the alticines (flea-beetles) in Galerucinae s.l. (Lingafelter and Konstantinov, 1999; Reid, 1995; Riley et al., 2002), relationships between the groups are controversial (see Chapter 1.3). Chrysomelinae seem to be closely related to Galerucinae s.l. (Duckett et al., 2004; Gómez-Zurita et al., 2008; Reid, 1995). Duckett et al. (2004) recovered Chrysomelinae as the sister group to the Galerucinae s.str. A recent study based on RNA data found a 'chrysomeline' clade with Galerucinae (with alticines) and paraphyletic Chrysomelinae (Gómez-Zurita et al., 2008).

In the NJ-tree in Figure 5.1 (Chapter 5) that includes one specimen of Crypto-

cephalinae and Lamprosomatinae each, both cluster together. This agrees with the placement of both groups in the 'Camptosomata' (Erber, 1988).

For relationships between the subfamilies see the leaf beetle phylogenies of e.g. Duckett et al. (2004); Farrell (1998); Farrell and Sequeira (2004); Reid (1995), and Gómez-Zurita et al. (2008). The relationships between cassidines and hispines, respectively galerucines and alticines are discussed in detail in Borowiec (1995); Staines (2002), and Chaboo (2007), respectively Duckett et al. (2004); Ge et al. (2012); Lingafelter and Konstantinov (1999) and references therein.

### 3.5 Conclusions

This study is the first attempt to investigate the leaf beetle fauna of the herbaceous and shrubby understorey vegetation of Podocarpus NP and RBSF in southern Ecuador. Considering that mainly one type of vegetation was sampled and that sampling is far from being complete, the more than 500 found morphospecies are rather a glimpse on the true diversity of the area. Further sampling as well as inclusion of the canopy fauna is likely to raise species numbers immensely. Given the poor taxonomic recording of leaf beetles in Ecuador, many of the found species are probably not yet recorded for Ecuador or might even be not yet described.

Although incomplete, the analysed selection of beetles provides a good insight into the characteristics of the leaf beetle assemblage: The chrysomelid fauna is species-rich and composed of few common and an overabundance of rare species, as it is typical for tropical arthropod assemblages. However, the number of rare species is likely to be overestimated due to undersampling. Communities differ between the three elevational levels, an issue that is investigated in detail in the following chapter (Chapter 4). The composition of leaf beetle subfamilies seems to be representative of a Neotropical leaf beetle fauna.

DNA barcode data led to higher species richness estimates and similar patterns in the comparison between elevational levels. The barcodes grouped morphospecies correctly into subfamilies.





# Habitat specialization and its influence on elevational diversity patterns inferred by DNA barcode data

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## 4.1 Introduction

One pivotal pattern of biodiversity is the variation of species richness along elevational gradients (Körner, 2000; McCain and Grytnes, 2010). In montane regions across the world the diversity of most plant and animal taxa can be described by one of three different patterns: a decreasing pattern, a low-elevation plateau pattern, or a mid-elevation peak pattern (McCain, 2009; Nogués-Bravo et al., 2008; Rahbek, 2005). Elevational diversity patterns were shown to systematically vary among taxa or geographic regions (McCain and Grytnes, 2010) but are usually assumed to be constant for one taxon on a single mountain. However, tropical mountains exhibit a complex topography shaping spatially heterogeneous habitats within elevational belts (Homeier et al., 2010; Takyu et al., 2002; Werner and Homeier, 2015). While spatial heterogeneity has been identified as a significant determinant of diversity (Stein et al., 2014), the influence of small-scale topography on patterns of diversity remains little understood.

Mountains are ideally described by a simple conical shape (Körner, 2000). However, the terrain formed by geological and hydrological processes over time is more complex and exhibits 'small-scale' geomorphological elements like mountain ridges or valleys (Beck et al., 2008b; Oesker et al., 2008). These habitats may strongly differ in abiotic conditions. For example, soils at ridge crests have higher rates of nutrient losses due to down slope fluxes, lower nutrient contents and lower rates of decomposition which significantly influence structural characteristics of the vegetation, e.g. species richness, forest canopy height and the production of wood and foliage (Takyu et al., 2002; Werner and Homeier, 2015). The variable abiotic and biotic conditions in different habitats may influence species richness of consumer taxa and may lead to habitat-specific differences in the elevational distribution of species. Moreover, community turnover between habitats may be heterogeneous along elevational gradients and lead to systematic changes in diversity patterns when scaling up from the local scale (including one habitat) to the regional multi-habitat scale.

Theory and empirical data support the hypothesis that higher temperatures, higher primary productivity, and more land area at lower elevations may foster habitat specialization and consequently species turnover between habitats. First, species turnover increases with productivity due to a higher importance of stochastic relative to deterministic assembly processes in highly productive environments (Chase and Leibold, 2002; Chase, 2010). Second, the benign warm climate at lower elevations has a positive effect on evolutionary rates (Allen et al., 2006; Rohde, 1992) and increases the importance of biotic interactions driving niche segregation and the evolution of specialization (Dobzhansky, 1950; Mittelbach et al., 2007; Pellissier et al., 2012; Schemske, 2002). Third, everything else being the same, higher amounts of land and the often higher productivity at lower elevations translate into a higher total amount of specific resources (e.g. total leaf biomass of a particular plant species) in lowland elevations than in highland elevations, which increases the probability of specialists' populations to persist. In contrast, populations in higher elevations may have to be generalists to persist over longer intervals of time (Srivastava and Lawton, 1998).

In this study, species diversity and turnover rates of leaf beetle communities are compared between montane ridge crest and valley forest habitats of southern Ecuador and it is inferred how differential turnover rates affect patterns of elevational species richness when scaling up from the local study site to the regional elevational belt scale. Leaf beetles (Coleoptera: Chrysomelidae) are major tropical herbivores and constitute one of the most speciose families in the whole tree of life (Basset and Samuelson (1996); Farrell and Erwin (1988); Wagner (2000); more detailed information on leaf beetles is given in Chapter 1.3). Their considerable specialization on host plants and large contribution to total biodiversity explains their key importance for estimating the total species richness on Earth. However, the extreme diversity, the little developed taxonomy of tropical beetle species and a suspected large proportion of cryptic diversity make the family particularly challenging for ecological studies (Costa, 2000; Flowers and Hanson, 2003; Furth et al., 2003; Gómez-Zurita et al., 2008; Jolivet et al., 1988) advocating the use of molecular approaches for species delimitation (Craft et al., 2010; Hebert et al., 2003b; Pfenninger et al., 2007; Smith et al., 2009, 2005; Tänzler et al., 2012). Molecular methods also allow inference of levels of intraspecific genetic diversity contributing to the understanding of trends in elevational species richness.

DNA barcoding was used to study the elevational diversity of leaf beetles in Podocarpus National Park and RBSF in southern Ecuador. The study region is considered to be one of the most diverse regions of the world (Brehm et al., 2005). Its steep slopes harbour two forest types at different topographic positions which differ in abiotic conditions and species composition of plant communities: ridge habitats at the upper slopes and valley habitats at the lower slopes (Homeier et al., 2010). For more detailed information on study area and sampling sites please refer to Chapters 2.1 and 2.2.

The following four predictions were tested:

1. Patterns of elevational diversity differ between ridge and valley forests.
2. Higher levels of productivity, area, and temperature provide increasing opportunities for habitat specialization and stochastic processes of community assembly (Chase, 2010). Therefore higher levels of species turnover between habitats at lower elevation than at higher elevations can be expected.
3. Elevational patterns of species richness are depending on the spatial scale of analysis (Nogués-Bravo et al., 2008) and may particularly depend on the variation in the rates of species turnover in space (Chase and Leibold, 2002). Due to higher species turnover between habitats at lower elevations stronger increases in species richness at lower than at higher elevation are expected with the spatial scale of the analysis increasing.
4. A decrease in productivity and increasing climatic harshness at high elevations may lead to higher levels of population bottlenecks or extinctions with subsequent recolonization of habitats by lineages from lower elevation (Ehinger et al., 2002; Gilles et al., 2007; Shama et al., 2011). Genetic diversity, here measured as haplotype diversity per species, can be therefore expected to decrease with increasing elevation.

## 4.2 Methods

Beetles were sampled on sampling sites within the Podocarpus NP and the adjacent RBSF, Ecuador. The study area is described in detail in Chapter 2.1. Sampling was conducted between May 2011 and April 2012 following a standardized design at the three different elevational levels Bombuscaro ('1000 m', premontane rainforest), ECSF ('2000 m', lower montane rainforest), and Cajanuma ('3000 m', upper montane rainforest or cloud forest) (classification of vegetation: Homeier et al. (2008)).

At each elevational level, 12 sites were sampled. Sampling sites were 20 × 20 m plots situated within homogeneous mature forest representative for the elevation in question and without visible recent natural or human disturbance (Homeier et al., 2010). At each elevational level, the 12 sites were located in two different types of habitats, six sites in valleys and six on ridges. Habitat types harbour different forest types and differ in several environmental parameters: Compared to sites on ridges, the sites in valleys are vegetated with a forest which differs in species composition, has a smaller canopy openness, consists of higher trees, has a higher biomass, and harbours a higher diversity of tree species. Furthermore, valley habitats are more nutrient-rich, have a minor organic layer thickness, a higher productivity, and a lower C:N ratio (Homeier et al. (2010); Werner and Homeier (2015); Homeier pers. comm.). With increasing elevation, the aboveground biomass of trees decreased as well as height of trees. The tree growth at higher elevation seems to be limited

by decreasing temperatures but also by shortage in nutrients and/or adverse soil chemical and physical conditions (Homeier et al., 2010).

Leaf beetle assemblages were sampled with a combination of three different collection methods: (1) Sweep netting, (2) beating of shrubs and smaller trees using a beating tray, and (3) hand-collection from the vegetation. All vegetation within reach was sampled (up to ca. 2.5 m). Each sampling-site was sampled following a standardized procedure: Sweep netting alongside two edges of the plot for 30 min; beating alongside the other two edges of the plot for 30 min; hand-collection within the plot for 30 min.

Beetles were killed and collected in 70% ethanol but transferred into 96% ethanol the following day. For each subsample (a subsample is either 30 min sweep netting, 30 min beating, or 30 min hand-collection) Chrysomelidae were sorted into preliminary morphospecies and one specimen of each morphospecies of each subsample was used for following morphological classification and molecular analyses.

Processing of the specimens and procedure of DNA barcoding of COI is described in Chapters 2.3 and 2.4. Sequences were submitted to GenBank (accession numbers KR424781–KR425417; see also Appendix Tab. C.1). All plot-samples resulted in a total of 995 leaf beetle specimens for which a barcode could be obtained (data set 3b, plot data set; Chapter 2.5, Appendix Tab. D.1).

Specimens were sorted into morphospecies (for information about morphospecies sorting see Chapters 2.5 and 5) and assigned to MOTUs using the different molecular species identification methods that are described in Chapter 6. As there were only few discrepancies between the methods (the methods were more or less conservative) only the results of Poisson tree processes (PTP) modelling were used for biodiversity analyses.

### **Analyses of biodiversity patterns**

For analyses of biodiversity patterns, a standardized data set was used which was based on four replicate samplings on each study site. All statistical analyses and data plottings were conducted in R 3.0.2 using the add-on packages RESHAPE2, VEGAN and SCIPLOT.

*Species richness patterns on the local scale:* Studies on insect diversity in the tropics always suffer from undersampling and correlation of sampling intensity with species richness causing strong biases in observed data (Brose et al. 2003). Therefore, asymptotic species richness of leaf beetles per study site was estimated using the non-parametric individual-based chao1 estimator provided in the VEGAN package and the estimated species richness was used for all main analyses (Chao et al., 2005; Gotelli and Colwell, 2011; Oksanen et al., 2013). The effect of elevation on species richness per site was analysed with ordinary linear models with estimated species richness per plot as the response (continuous) and elevational level (factorial) as the explanatory variable. In case of significant differences between elevational levels in the general model, post-hoc pairwise t-tests with pooled standard deviations were conducted to test for differences between pairs of elevational levels. In addition to the above mentioned analysis in which it was not differentiated between valley and

ridge crest habitats, trends in species richness with elevation were analysed for the two forest habitats separately using the same procedures as described for the total data set.

*Species richness patterns on the regional scale:* To calculate patterns of species richness with elevation at the regional scale the data of all study plots per elevational level were pooled and the asymptotic cumulative species richness ( $\pm$  SE) was estimated using the non-parametric chao1 estimator (Chao et al., 2005; Gotelli and Colwell, 2011). Regional species richness was estimated for plots of ridge and valley forests separately and for all plots per elevational level combined. To compare patterns of species richness for a standardized (rarefied) number of sampled individuals per elevational level individual-based species accumulation curves were calculated for each elevational level. Species accumulation curves were calculated using the method 'random' which adds up individuals in a random order with 1000 iterations and calculates the mean  $\pm$ 95% confidence interval.

Trends in species turnover across elevations and habitats were visualized using non-metric multidimensional scaling (NMDS) and statistically analysed using permutational multivariate analysis of variance (MANOVA) provided in the VEGAN package. For measuring the dissimilarity in species composition between the communities of different plots the chao dissimilarity index was used (Chao et al., 2005). The chao dissimilarity index is recommended for samples that differ in the intensity and completeness of sampling and which is particularly suited for data which are known or suspected to contain high numbers of rare or undetected species, and therefore seems appropriate for highly speciose leaf beetle communities in which typically a large number of singletons appear and many species remain undetected. The function *adonis* in the R package VEGAN (Oksanen et al., 2013) was used to partition the variation in overall beta-diversity among the effects of elevation, habitat and the elevation-habitat interaction and the significances were tested with permutation tests with 10,000 permutations. In case the beta-diversity between habitats was heterogeneous along the elevational gradient the interaction term was expected to be significant. With a partial Mantel test it was tested if the patterns of beta-diversity could be explained by differences in distance and elevation between the plots of different altitudinal levels.

To test for differences in relative genetic diversity an index of haplotype diversity was calculated for each elevational level. Haplotype diversity was defined as the probability that two individuals of one MOTU show different haplotypes. Probabilities were calculated based on a data set of all individuals collected per elevational level. Only species were considered for which more than two individuals per elevation were collected. The effect of elevation on haplotype diversity was analysed with ordinary linear models.

### 4.3 Results

#### Local and regional patterns in species richness

A total of 271 morphospecies and 453 haplotypes were detected which were differentiated by Poisson tree processes (PTP) modelling into 294 MOTUs. The estimated species richness (i.e. MOTU richness; for better readability hereafter simply 'species richness' is used) per site (local level) significantly varied among the three elevations ( $F_{2,33} = 11.79$ ,  $p = 0.0001$ ) and was highest at 2000 m a.s.l. and significantly lower at 1000 and 3000 m a.s.l. (post-hoc pairwise t-test with pooled SD: 1000–2000 m:  $p < 0.05$ , 1000–3000 m:  $p < 0.01$ , 2000–3000 m:  $p < 0.0001$ ; Fig. 4.1A). When patterns were analysed for the two forest habitats separately, a mid-elevation peak pattern was evident in forests on ridges but not in valleys where species richness peaked at the lowest elevation but did not significantly differ from species richness at 2000 m a.s.l. (paired t-test:  $p = 0.8$ ). However, for both habitats the lowest species richness was found at 3000 m a.s.l.

When species richness patterns were analysed at a regional level (i.e. by estimating the cumulative MOTU richness of all plots per elevational level), species richness did not differ between 1000 and 2000 m a.s.l. (Fig. 4.1B). When corrected for differences in the number of sampled individuals, species richness was even slightly higher at 1000 m a.s.l. than at 2000 m a.s.l. (Fig. 4.1C). When cumulative species richness was analysed for each habitat separately, the mid-elevation peak of diversity which was detected in ridge forests at the local scale was strongly reduced at the regional level. Regional level diversity in valley forests did not differ between 1000 and 2000 m a.s.l. Regional species richness was in all cases lowest at 3000 m a.s.l.

#### $\beta$ -diversity between habitats and elevational levels

NMDS and permutational MANOVA analyses revealed a clear differentiation in the composition of leaf beetle communities among the three elevations (Fig. 4.2A, Tab. 4.1). The differences between 2000 and 3000 m a.s.l. were more pronounced than between 1000 and 2000 m a.s.l. Communities at 1000 and 3000 m a.s.l. differed most strongly in the composition of MOTUs. A significant interaction between habitat and elevation was found (Tab. 4.1): While communities of ridge and valley habitats strongly differed at 1000 m a.s.l., differences were less strong but still significant at 2000 m a.s.l. and non-significant at 3000 m a.s.l., suggesting a reduced habitat differentiation of species communities in higher elevations. Even though no significant difference was found in the leaf beetle composition of ridge and valley forests at 3000 m a.s.l., the turnover of MOTU between plots was generally very high and similar among the plots of one habitat as among the plots of two different habitats (Fig. 4.2B).

The higher beta-diversity between habitats in lower elevations could not be explained by differences in distance and elevation between the sites at different elevational levels (Mantel test,  $p > 0.05$ ).

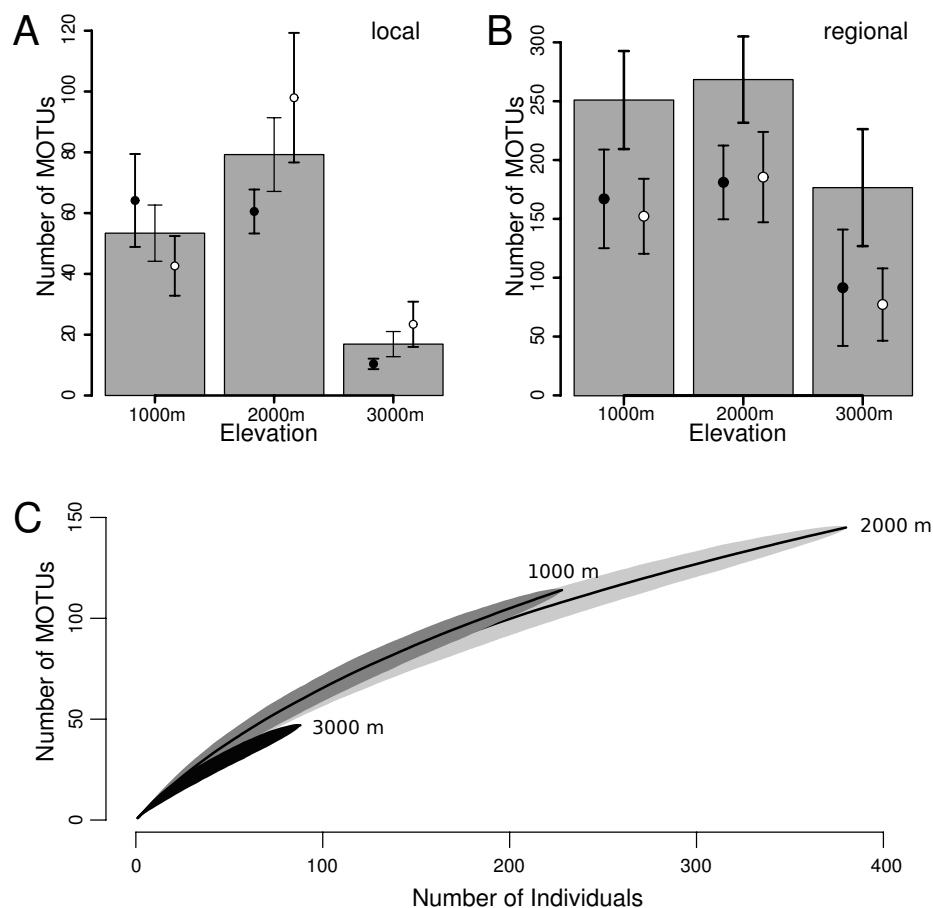


Figure 4.1: Mean ( $\pm$  SE) number of estimated MOTUs per study plot (A), total number of estimated MOTUs per elevational level (B) and the cumulative number of MOTUs with increases in the number of sampled individuals (C). A: Barplots and the error bars show the mean number of estimated species richness  $\pm$  standard error found on plots of both habitats; the black and white dots with error bars show the values for plots in valley and ridge forests, respectively. The hump-shaped pattern is most pronounced for the plots on ridges (pairwise t-test: 1000–2000 m:  $p < 0.02$ , 2000–3000 m:  $p < 0.01$ , 1000–3000 m:  $p = 0.4$ ). B: Barplots and the corresponding error bars show the total number of estimated species  $\pm$  standard error for each elevational level; the black and white dots with error bars show the values for plots in valley and ridge forests, respectively. C: Species accumulation curves (mean  $\pm$  95% confidence interval) show the increase in species richness with increasing number of sampled individuals.

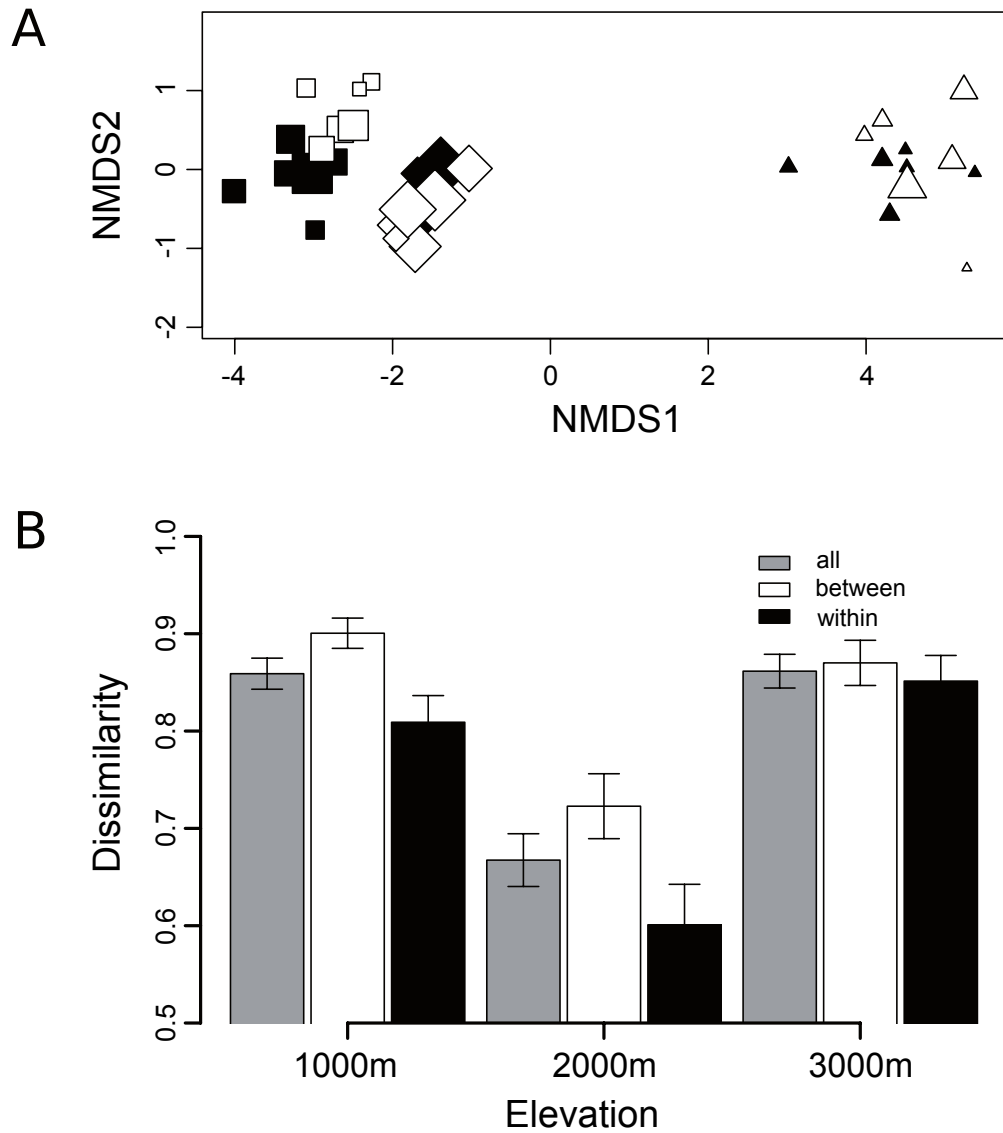


Figure 4.2: NMDS ordination of MOTU data (A) and mean ( $\pm$  SE) community dissimilarity among plots within elevational levels (B). A: Different symbols indicate different elevational levels (square = 1000 m, diamond = 2000 m, triangle = 3000 m) and habitats (black = forests in valleys, white = forests on ridges). Size of symbols is proportional to the number of estimated MOTUs per plot. B: Bars show mean estimates of community dissimilarity  $\pm$  SE among the plots on one elevational level: all = dissimilarity among all plots; between = dissimilarity among plots of different habitats; within = dissimilarity among plots of the same habitat.



Table 4.1: **Results of permutational MANOVA analysis testing on the effects of elevation and habitat on the species composition of leaf beetle communities.** At 1000 m and 2000 m communities in ridge and valley habitats significantly differed in their composition, which was, however, not the case at 3000 m (1000 m:  $F = 2.27$ ,  $p = 0.006$ ; 2000 m:  $F = 3.19$ ,  $p = 0.031$ ; 3000 m:  $F = 1.25$ ,  $p = 0.211$ ).

	<b>F</b>	<b>df</b>	<b>r<sup>2</sup></b>	<b>p</b>
Elevation	7.43	2	0.29	0.001
Habitat	2.05	1	0.04	0.006
Elevation $\times$ Habitat	2.07	2	0.08	0.002
Residuals		30	0.59	

### Genetic diversity

Haplotype diversity (defined as the probability that two individuals of one MOTU show different haplotypes) did not significantly differ among elevational levels (ANOVA,  $F_{2,33} = 0.57$ ,  $p = 0.57$ ; Fig. 4.3).

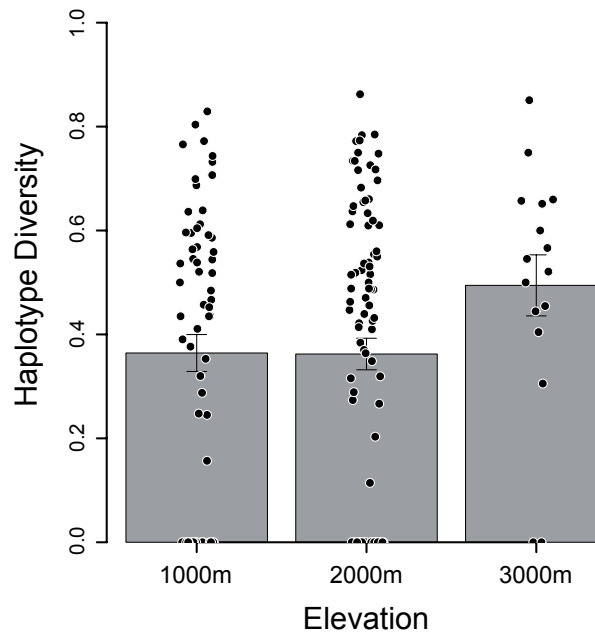


Figure 4.3: **Estimates of relative genetic diversity in different elevation levels, i.e. the haplotype diversity for MOTUs for which more than one individual was sampled per elevational level.**

## 4.4 Discussion

The complex topography of mountains produces habitats with strongly differing environmental conditions across and within elevational levels. The patterns of elevational diversity in the studied leaf beetles differed between ridge and valley forests. Moreover, a higher turnover of communities between and within forest habitats at lower elevations shifted mid-elevational diversity patterns towards lower plateau patterns when scaling up from the local study site level ( $\alpha$ -diversity) to the regional elevational belt level ( $\gamma$ -diversity).

### **Differences in patterns of elevational $\alpha$ -diversity between ridge crest and valley habitats**

While being largely ignored in elevational gradient studies, the results show the importance of small scale topography (here: topography within elevational belts) for patterns of elevational diversity. Sites on ridges strongly vary in several environmental parameters from sites in valleys: They have less nutrient-rich soils, a thicker humus layer, a lower productivity, and a higher C:N ratio of foliage. The forests on ridges have a different plant species composition from those in valleys, a lower and more open canopy layer, lower biomass and lower diversity of tree species (Homeier et al., 2010; Werner and Homeier, 2015). Differences in soil nutrient availability influence leaf quality and concentrations of phenolics (Werner and Homeier, 2015) which likely affect herbivore diversity. While there is insufficient data to directly relate leaf beetle diversity to the variation in characteristics of the vegetation, this study underscores the value of considering the small scale topography in studies of elevational diversity. Differences in the kind of sampled habitats among studies may explain some of the variation in patterns of elevational diversity found for single taxa in meta-analyses (McCain and Grytnes, 2010). Moreover, as some environmental variables vary more strongly between the habitats of one elevational level than others (primary productivity and plant species richness strongly differ between ridge and valley habitats while average temperature or O<sub>2</sub> concentration remain similar), incorporating the spatial habitat heterogeneity within elevational belts in the design of macroecological studies may allow more clear tests of the hypotheses commonly used to explain diversity gradients.

### **Habitat-differentiation of communities and the influence on patterns of elevational $\gamma$ -diversity**

Mid-elevational peaks of diversity as found for the pooled data in the present study are frequently reported in the ecological literature (Nogués-Bravo et al., 2008; Rahbek, 2005), in particular also for beetles on tropical mountains (Escobar et al., 2005; Furth, 2009; Sánchez-Reyes et al., 2014). In most montane regions, these patterns are in conflict with major climate-based hypotheses to explain large scale diversity gradients, e.g. the temperature hypothesis, the productivity hypothesis, or the area hypothesis (Brown et al., 2004; Currie et al., 2004; Mittelbach et al., 2007; Rosenzweig, 1995). Like in most other montane areas, in the study region tempera-

ture, primary productivity, and area decrease with elevation (Homeier et al., 2010; Körner, 2000; Moser et al., 2007; Wolf et al., 2011) suggesting monotonous positive correlations of elevation and species richness. Also plant species richness, a variable often positively correlated to herbivorous insect diversity is declining with elevation in the study area (Homeier et al., 2010, 2008) and cannot explain the mid-elevation peak pattern of species richness found at local scales.

By using leaf beetle diversity from two different forest habitats at multiple sites the study shows that mid-elevational peak patterns may disappear when increasing the spatial scale of diversity from local alpha-diversity to regional gamma-diversity. Study sites in the middle of the elevational gradient exhibited highest levels of species richness at local scales but showed lowest rates of species turnover between sites. It is widely recognized that spatial scale is a crucial factor influencing species richness patterns along environmental gradients (Gaston, 2000; McCoy, 1990; Nogués-Bravo et al., 2008; Rahbek, 2005). Local assemblage structure is inseparable from the regional context (Gaston, 2000), and it has been shown that, similar to the present study, the same data can show different diversity patterns on a local than on a regional level (Black and Prince, 1983; Chase and Leibold, 2002).

As beta-diversity often increases with increasing productivity, rates of species turnover were expected to decrease with elevation (Bai et al., 2007; Bonn et al., 2004; Chase and Leibold, 2002; Evans et al., 2008; Harrison et al., 2006). However, this prediction could only be confirmed for the two lower elevational levels while species turnover at 3000 m a.s.l. was nearly as high as at 1000 m a.s.l. Interestingly, at 1000 m a.s.l., the high turnover is related to a clear differentiation of leaf beetle communities between forests on ridge crests and in valleys. This habitat differentiation is most pronounced at 1000 m a.s.l., less strong but still significant at 2000 m a.s.l. (where turnover is generally lower), whereas absent at 3000 m a.s.l. The larger and more productive lower elevations seem to offer more opportunities for specialization (More Specialization Hypothesis; Srivastava and Lawton (1998)) and possibly allow habitat differentiation of herbivore communities into valley and ridge forest habitats.

However, in general turnover rates were very high between elevations with only a small overlap in leaf beetle community composition between 1000 and 2000 m a.s.l. and no species overlap at all with 3000 m a.s.l.: Although the linear distance between each of the three areas is only ~20 km, there are striking differences in climate and vegetation (pers. observ.). The turnover of tropical insect communities along elevational gradients is generally rapid (Brühl et al., 1999; Ghalambor et al., 2006; Janzen, 1967) and there are often large differences in insect communities in considerably smaller ranges than 1000 m (e.g. Olson (1994); Smith et al. (2014)).

It should be emphasized that spatial issues are not considered to be of sole importance for explaining mid-elevational peak patterns. In some mountain regions, land area and/or productivity exhibit hump-shaped distributions along the elevational gradient and reflect patterns in species diversity (Brown, 2001; McCain and Grytnes, 2010; Sanders, 2002). Whereas the mid-domain effect has been rejected as a general explanation of mid-elevational peak pattern it may, nevertheless, ex-

plain patterns of diversity in some taxa, particularly in those with large elevational ranges (Mid-domain effect; Brehm et al. (2007); Colwell and Lees (2000); Colwell et al. (2004)). Nevertheless, the high proportion of mid-elevational peak patterns in the ecological literature and the fact that they are often incongruent to major biogeographic hypotheses for explaining diversity gradients calls for further general explanations and issues of spatial scale could be an important factor to consider in future analyses. In this respect, the mismatch between a high proportion of hump-shaped diversity patterns reported from elevational gradient studies and a near lack of those patterns along latitudinal gradients may be due to systematic differences in the spatial scale of analyses. While most studies along elevational gradients are conducted at local scales and measure alpha-diversity, a high percentage of latitudinal gradient studies measure regional richness in large quadrats (often >100 km<sup>2</sup>), i.e. in areas which incorporate multiple habitats. If species turnover among habitats varies along environmental gradients (or the diversity of habitats) this may cause systematic differences in diversity patterns of studies conducted at local versus regional scales.

### Haplotype-diversity

In contrast to the expectations, the haplotype-diversity of species did not differ significantly between elevational levels. Species populations at higher elevations were expected to be less genetically diverse because limited resources and a harsher climate with extreme climatic events may lead to smaller and temporally less stable populations (Ehinger et al., 2002; Frankham, 1996; Gilles et al., 2007; Shama et al., 2011; Srivastava and Lawton, 1998). Population bottlenecks or local extinction with subsequent immigration from other mountain areas were expected to cause a lower diversity of haplotypes within populations (Glenn et al., 1999; Hoelzel et al., 2002; Nei et al., 1975; Weber et al., 2004). In contrast, the warm and favourable climatic conditions at low elevations were assumed to lead to larger and more stable populations holding higher levels of genetic diversity over time. In addition, higher metabolic rates and related nucleotide substitutions in warmer climates may foster higher levels of genetic diversity (Allen et al., 2006; Rohde, 1992). Indeed, elevational gradients in genetic diversity have been reported e.g. for species of shrews (Ehinger et al., 2002) and salamanders (Giordano et al., 2007).

An explanation for the lack of any systematic differences in haplotype diversity among elevational levels could be the high connectivity between the Andean mountains regions. The large and connected high elevation habitats may facilitate the sustenance of large and stable populations with high levels of genetic diversity.

It is also possible that high altitude communities contain generalist rather than specialist species that are less susceptible to adverse environmental influences and have a lower risk of extinction as they can adapt to the harsh conditions at higher elevations (Packer et al., 2005). For example, it has been shown that alpine Chrysomelidae species show a more broadly oligophagous or polyphagous feeding behaviour (Lopatin, 1996). An indication for this could be the slightly higher haplotype-diversity of species at 3000 m a.s.l. that may indicate the prevalence of generalist

species. It has been shown that specialist species have a reduced genetic variation (due to a smaller effective population size) (Kelley et al., 2000; Packer et al., 2005; Zayed et al., 2005). This agrees with the notion that usually low productivity communities are dominated by generalists as some plant resources are too scarce to support viable specialist populations (Srivastava and Lawton, 1998). The finding that habitat specialization in this study is most pronounced at 1000 m a.s.l. and absent at 3000 m a.s.l. is in line with the idea of a higher prevalence of generalist species at higher elevations, too.

#### Methodological considerations

All results have been revealed in a similar way by morphospecies- and MOTU-based analyses. The total number of found species is ~8% higher for MOTUs compared to morphospecies. This may be explained by species indiscernible by the morphospecies approach (cryptic diversity). Possibly, integration of DNA barcoding into biodiversity studies can prevent an underestimation of diversity (Hebert et al., 2004; Witt et al., 2006). Due to a high number of uniques in the data set, cryptic diversity has a rather small influence on species number as morphospecies found only once cannot be split by the molecular approach. This is enhanced by the study design, analysing only one specimen per morphospecies per sample. However, communities with many species but a low abundance of each individual species and many rare species are typical for tropical insects. It has been shown that in such data sets and on a geographically restricted level, different methods of morphological and molecular species delimitation can lead to very similar results (Chapter 6).

DNA barcoding is often integrated in biodiversity studies not only to consider cryptic diversity but also to provide additional information at the infra-specific level (García-Lopez et al., 2013; Monaghan et al., 2009; Papadopoulou et al., 2011). It is especially useful for a rapid analysis of unknown tropical insect faunas where taxonomic identification is still missing and where it usually reveals diversity patterns in a similar way as morphological approaches (Smith et al., 2005; Tänzler et al., 2012).

## 4.5 Conclusions

First, the study highlights the complexity of insect communities in tropical montane regions that are so far severely understudied. It confirms the findings by Werner and Homeier (2015) who showed the relevance of topographic positions because contrasting biotic and abiotic conditions found along short topographical gradients are an important source of beta-diversity in tropical mountains and should therefore be considered in the sampling design. Furthermore, the study affirms the importance of spatial scale for the analysis of diversity patterns along elevational gradients.

Second, the study shows the suitability of DNA barcoding to examine even complex ecological questions. It is a practical example demonstrating the useful implementation of routine DNA barcoding for analysis of biodiversity patterns and their ecological implications. It is a further example showing that barcoding goes beyond

the mere function of species discovery and identification. Patterns of community composition and turnover can be analysed and interpreted even without taxonomic information.

Finally, the study suggests that despite a strong loss of diversity of local communities at the species level, genetic diversity within species may remain relatively stable along the elevational gradient. It would be interesting to use the DNA barcode data for further phylogenetic measures of the leaf beetle community structure along the elevational gradient, such as phylogenetic diversity (PD) or nearest taxon index (NTI) (Brehm et al., 2013; Smith et al., 2014).

# Comparison of morphological and molecular species delimitation approaches

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## 5.1 Introduction

In studies on biodiversity of tropical arthropods that are hyperdiverse and extremely abundant (although most individual species taken by themselves are not abundant) often many thousand individuals may accrue (see summary in Coddington et al. (2009)). The sheer mass of individuals and the high species richness usually prevents thorough taxonomic analyses as this is very time consuming. Especially in the light of the fast rates of habitat conversion and destruction threatening particularly ecosystems with the highest biodiversity (e.g. tropical rainforests; Corlett and Primack (2011); Laurance and Peres (2006); Primack (2014)), less time intensive methods of assessing this diversity are urgently required.

Difficult access to taxonomic expertise also impedes taxonomic identification: In large taxa, a single taxonomist is specialized only on certain subgroups and usually is not able to identify all species of a set of samples. For example, in Neotropical Chrysomelidae, the taxonomic expertise for the whole family is covered by the sum of experts for certain subfamilies (e.g. Borowiec (1998); Chaboo (2007); Chaboo and Borowiec (2003); Flowers (2004a,b,c); Flowers and Chaboo (2009); Furth and Savini (1996); Furth (2007); Staines (2002); Windsor et al. (1995)). Additionally, identification literature and keys for many taxa are incomplete, old, or not existing at all. In Neotropical Chrysomelidae, identification keys are scarce and exist only for certain subgroups or regions and are often not at species-level (Borowiec and Świętojańska, 2014; Furth, 1992; Staines, 2002, 2013, 2009). There is also a lack of easily accessible and reliably identified reference collections (Furth et al., 2003).

Due to such adverse circumstances, for most studies on tropical arthropod diversity, working units as surrogate for species serve well while taxonomic identification and description of new species has to be postponed. Traditionally, morphospecies are used as working units, but these have been recently complemented by molecular working units.

Morphospecies in the broad sense are species discerned with morphology-based taxonomic techniques but not necessarily named (Basset et al., 2008, 2004; Oliver and Beattie, 1996). The distinction of morphospecies may be performed on different levels of preciseness, for example, dissection of genitalia might be included. However,

usually the term morphospecies refers to species-like groups of specimens sorted more superficially on the basis of external morphology and without the use of identification keys (Krell, 2004; Oliver and Beattie, 1996; Pik et al., 1999). In this study the term 'morphospecies' is used in this sense.

The morphospecies approach is sometimes used synonymously with parataxonomy. The term parataxonomy was originally coined by Janzen (Janzen, 1991, 2004; Janzen et al., 1993). It described a concept of training local people to support inventorying and monitoring tropical biodiversity and so improve the flow of primary information on tropical biodiversity (Basset et al., 2004). Beside preliminary sorting into morphospecies, the expertise of the trained parataxonomists comprises collecting and preparing specimens and databasing the associated information (Basset et al., 2004). Parataxonomy is widely used in terrestrial arthropod research (Basset et al., 2008; Longino and Colwell, 1997; Novotny et al., 2002a). However, the term parataxonomist may also be applied to local collectors, students, professional zoologists and botanists focusing on ecological studies, or taxonomists operating outside of their range of expertise (Basset et al., 2004).

The morphospecies approach is widely used and is a standard method in studies on tropical arthropod biodiversity (Basset et al., 2004; Springate and Basset, 2004; Wagner, 2000). It is not only applied to study tropical rainforest arthropods but also other taxa that are extremely abundant, speciose, and/or morphologically hard to identify (e.g. terrestrial nematods: Bernard and Schmitt (2005); Lawton et al. (1998), or benthic macroinvertebrates: Costa and Melo (2008); Duncan and Brusven (1985)). However, some authors criticize a low accuracy and the problem of lacking comparability and replicability (Krell, 2004).

Indeed, the superficial morphospecies sorting relying only on external characters (as used in this study) has its shortcomings. Morphospecies sorting may be confounded by cryptic diversity, sexual dimorphism, polymorphism, or juvenile forms. This may result into splittings of morphospecies into two or more species and lumping of morphospecies into a species. Furthermore, for certain organisms, even morphospecies sorting might be difficult. These may include premature stages, very small organisms (meio- and micro-fauna, zooplankton) and species only distinguishable by subtle or geographically variable morphological characters (Blaxter et al., 2004; Bucklin et al., 2007; Decaëns et al., 2013; Plaisance et al., 2009; Vences et al., 2005).

In such cases, species-like working units based on molecular characters are a useful tool. The DNA barcode marker cytochrome *c* oxidase I (COI) has been established as a species-specific identification marker (Hebert et al., 2003a). According to differences in the sequences, specimens can be classified into molecular operational taxonomic units (MOTUs; Floyd et al. (2002)). There exist a variety of methods to derive species limits from DNA sequence data, several methods are described in Chapter 6.

The barcode/MOTU approach is comparable between studies and sites (Floyd et al., 2002) whereas morphospecies at this point can be flawed (Krell, 2004). However, both methods should be followed by a thorough taxonomic analysis with de-



scription and naming to allow connection of the species-specific facts with existing literature.

Neotropical Chrysomelidae are a very diverse and species-rich taxon. Due to the lack of data and/or difficult access to taxonomic information, an approach with rapid identification of (morphological and molecular) working units is standing to reason. As leaf beetles are extremely diverse, very abundant, feature many small species and species where cryptic diversity, colour-polymorphism or sexual dimorphism occurs, it is advisable to complement the widely used morphospecies approach with a DNA barcode approach.

## 5.2 Methods

Chrysomelidae were sampled between November 2010 and June 2012 in parts of Podocarpus National Park and RBSF, Ecuador (detailed information about the study area is given in Chapter 2.1). Beetles were mainly collected by sweep netting, beating, and hand-collection of the lower vegetation. Additionally, light-, Malaise-, flight interception-, and pitfall traps have been used (detailed information about sampling methods and design is given in Chapters 2.2 and 7). Laboratory analyses and handling of the specimens are described in Chapters 2.3 and 2.4.

The dry mounted specimens were sorted into morphospecies. Classification into morphospecies was revised and verified by Dr. Thomas Wagner who is an experienced taxonomist for Chrysomelidae with afrotropical Galerucinae being his area of expertise (see e.g. Wagner (2004, 2007a,b); Wagner and Kurtscheid (2005)). Only ectoskeletal characters were considered, without the use of dissected parts and without identification literature. Characters for morphospecies delimitation are shape of head, pronotum, and total body, surface structures, and hairs or spines. Body size or colours may be used carefully considering that they may vary e.g. due to recent ecdysis (more information about the morphospecies concept is given in the introduction of this chapter). Morphospecies received a subfamily name and a number. Hispinae and Cassidinae are treated as separate subfamilies, likewise Alticinae and Galerucinae.

For all specimens for which a barcode sequence could be obtained (1475 specimens, data set 1b), a statistical parsimony analysis as implemented in TCS v.1.21 (95% connection limit) was used to group sequences into separate haplotype networks (more detailed information about statistical parsimony analysis is given in Chapters 2.5 and 6). The haplotype networks are termed MOTUs hereafter. All 1475 specimens that were assigned to a morphospecies and a MOTU were included in the analysis of congruence between both methods.

A Neighbor-Joining-Tree (NJ-Tree; Saitou and Nei (1987)) based on a MUSCLE alignment (Edgar, 2004) of all 1475 sequences was constructed in Geneious version 7.1.5 (Biomatters Ltd.; <http://www.geneious.com/>). Figures 5.2–5.5, 5.9, 5.10, 5.13, 5.14, and 5.15 showing splittings and lumpings in certain groups are excerpts from the NJ-Tree of all 1475 specimens. Figure 5.1 shows a NJ-Tree of

representatives of all MOTUs (493 sequences).

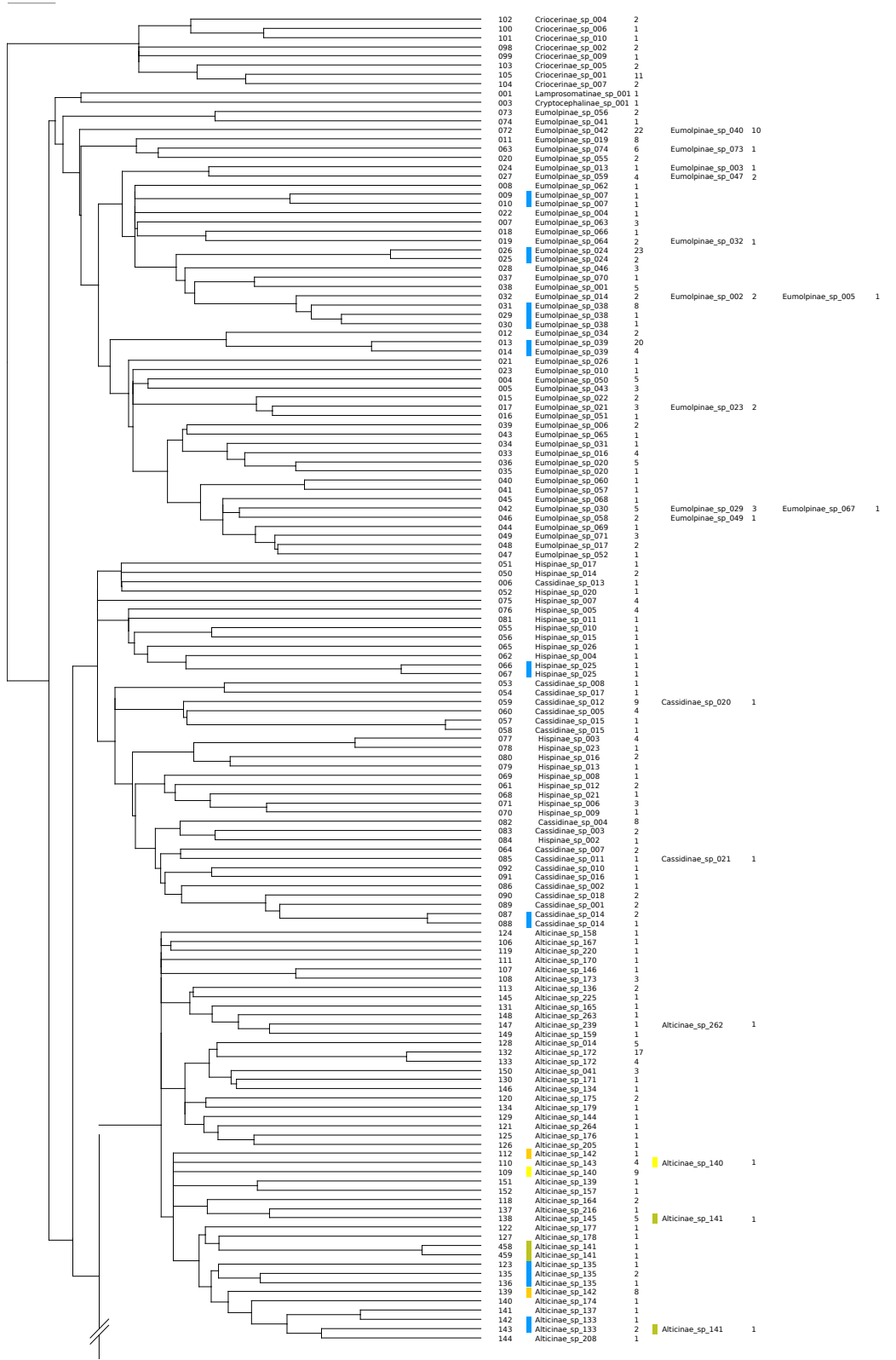
### 5.3 Results

The barcode approach with species delimitation using haplotype networks estimated ten percent higher species numbers compared to the morphospecies approach (448 morphospecies, 493 MOTUs). This discrepancy arose due to splittings of morphospecies into two or more MOTUs or lumpings of two or more morphospecies (respectively parts of them) into one MOTU. The occurrence of more splittings than lumpings resulted in a higher number of MOTUs than morphospecies. The similar numbers of morphospecies and MOTUs arose partly due to the fact that splittings and lumpings compensated one another. However, the number of perfect matches was comparatively low: 63% of all morphospecies corresponded exactly with one MOTU, i.e. contained specimens of only one MOTU and at the same time all of them. Five percent of all morphospecies were both split and some specimens were placed into other morphospecies at the same time. An overview of all splittings and lumpings is given in Figure 5.1.

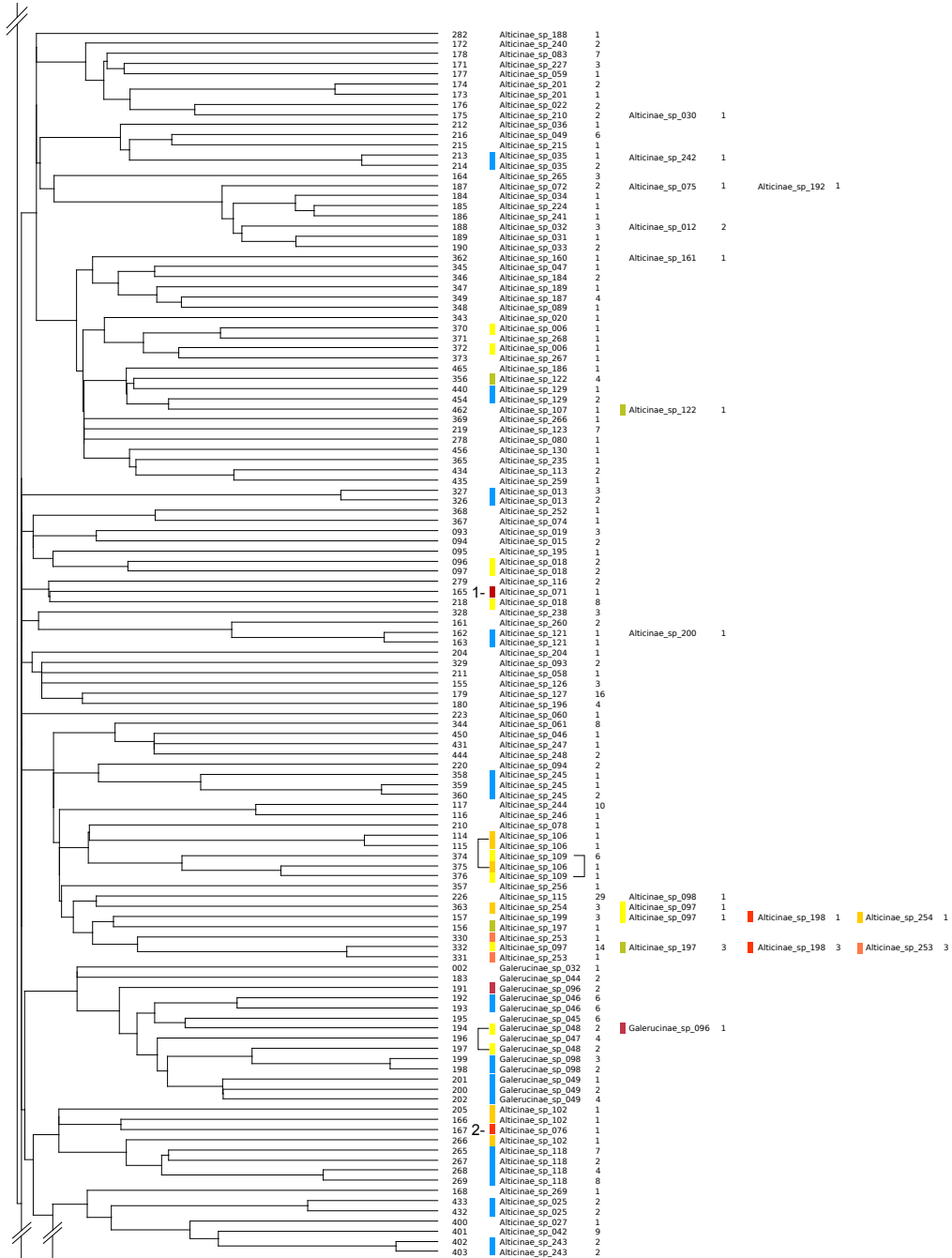
In most cases, individuals of two (76% of all lumpings) or three (16%) morphospecies were united into one MOTU, in four percent four, respectively five morphospecies were fused. Similarly, most splittings divided a morphospecies into two (69%) or three (21%) MOTUs. In eight percent of all cases, a morphospecies was split into four MOTUs and only once a morphospecies was split into five (1%) or six MOTUs (1%).

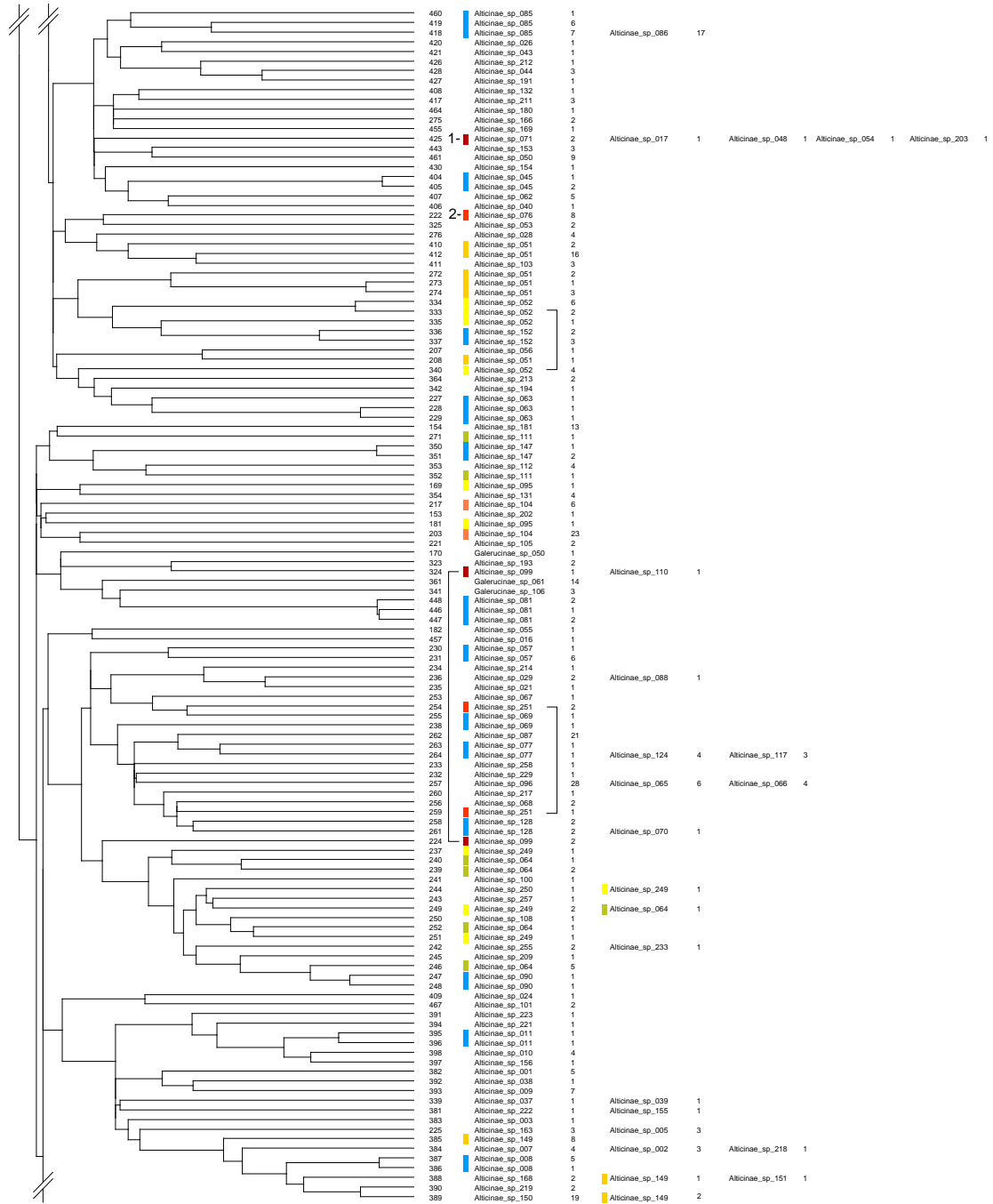
There were slight differences between the five most abundant subfamilies. Whereas in Alticinae and Galerucinae MOTUs estimated species number 14% higher compared to morphospecies, in Hispinae it was only five percent and in Cassidinae splittings and lumpings compensated resulting in an equal number of MOTUs and morphospecies. In Eumolpinae there were more lumpings than splittings resulting in a lower estimation of species number by MOTUs by nine percent. Morphospecies that were split and lumped at the same time occurred only in Alticinae (19 cases) and Galerucinae (three cases). The proportion of perfect morphospecies (matching exactly with one MOTU) was highest in Hispinae (95% of all morphospecies). It was considerably lower in Cassidinae (67%), Galerucinae (65%), Alticinae (59%), and Eumolpinae (54%).

Despite a comparatively low 'accuracy' relative to morphospecies, Eumolpinae seemed to be one of the less challenging subfamilies beside Cassidinae and Hispinae. Only two or three complete morphospecies were lumped into one MOTU or morphospecies were split into two or three MOTUs. No morphospecies was split and lumped at the same time. The most challenging subfamilies were Galerucinae and especially Alticinae. The difficulties most probably arose due to the large number of specimens and species. Furthermore, the Alticinae in the data set contained many very small species (<5 mm) and many species looking similar (e.g. Alticinae spp. 097, 197, 198, 199, 253, and 254) exacerbating morphospecies sorting.



# Chapter 5. Comparison of morphological and molecular species delimitation approaches





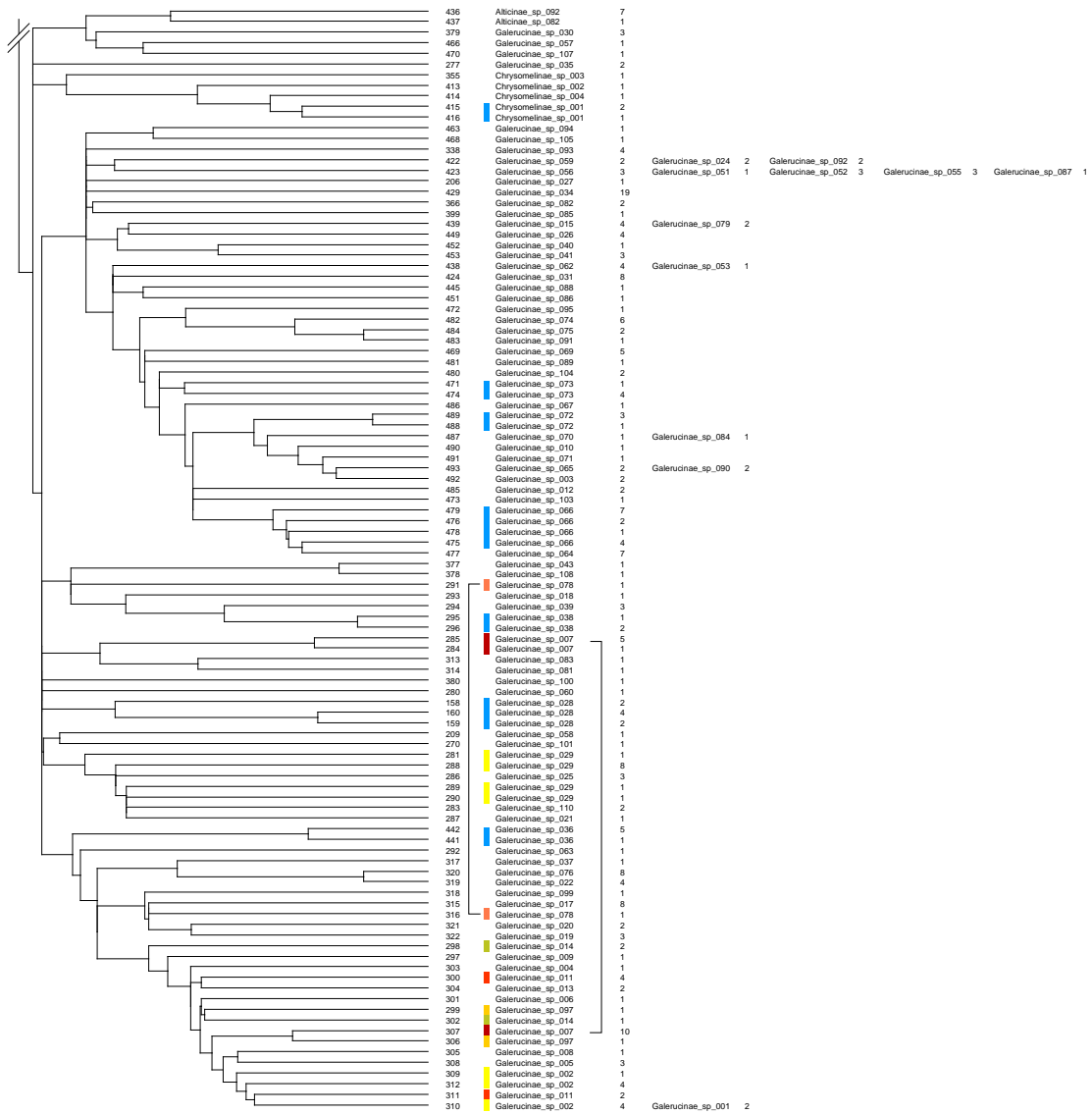


Figure 5.1: Overview of splittings and lumpings. NJ-Tree with ultrametric branch lengths of all MOTUs. Each MOTU is represented by one sequence. Branch labels indicate name/number of the MOTU. Morphospecies names behind the MOTU name represent the one or more morphospecies the specimens of the MOTU belong to and the numbers how many specimens of the respective morphospecies there are. Coloured bars indicate splitting of a morphospecies into several MOTUs; morphospecies split into MOTUs next to each other are indicated by blue bars.

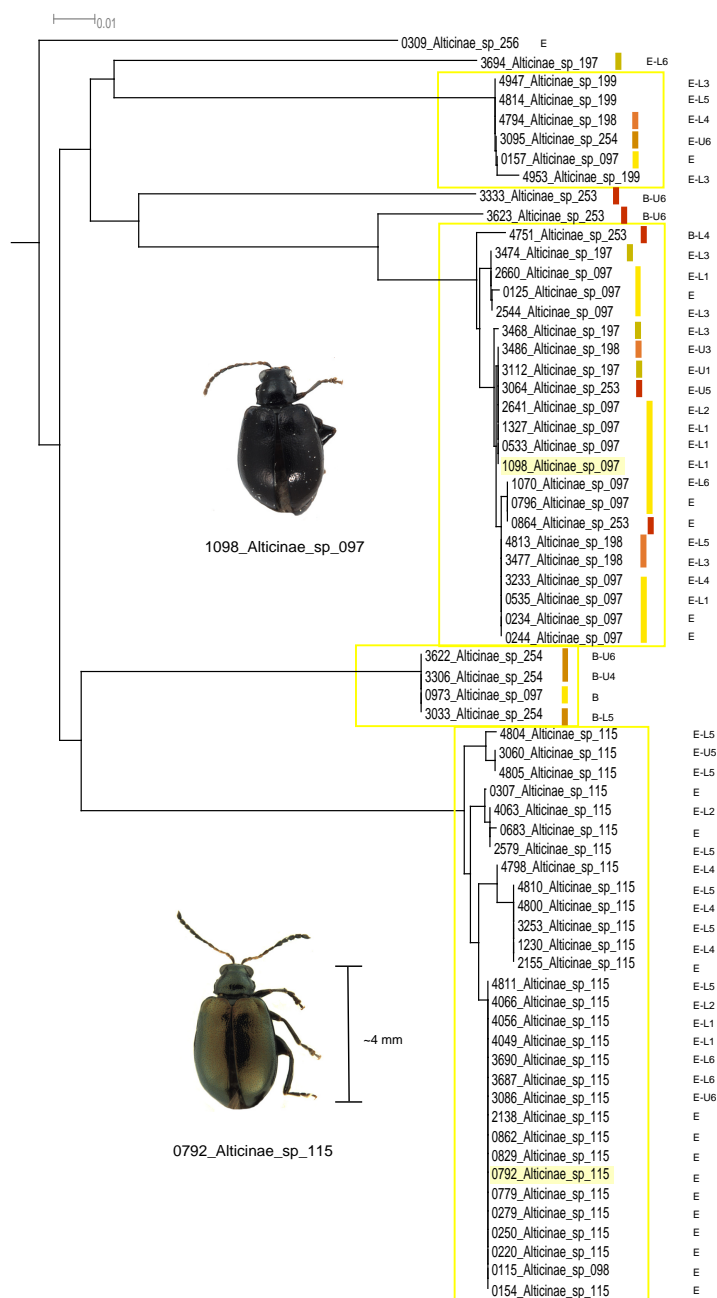


Figure 5.2: **Splittings and lumpings within a group of several small alticine morphospecies.** Coloured bars behind morphospecies names indicate splitting of the morphospecies into more than one MOTU; morphospecies that are lumped into one MOTU are bordered by a coloured box. Abbreviations behind the individuals indicate the sampling area (B = Bombuscaro, E = ECSF), respectively the plot where the specimen was collected. Information on collection plots is given in Tab. B.1 (Appendix).

The case of Alticinae spp. 097, 197, 198, 199, 253, and 254 is an example where morphospecies were both split and lumped as well (Fig. 5.2). Each of them (except Alticinae sp. 199) was split into two or three MOTUs and also lumped (two, respectively four morphospecies lumped into one MOTU). The morphospecies were completely resorted into six MOTUs. The six morphospecies consisted of very small (<5 mm), black specimens all looking very much alike, making morphospecies sorting very confounding. Another example for morphospecies that were split and lumped (Alticinae sp. 149) is given below ('Precarious taxa').

Two hundred and six morphospecies occurred only once in the data set (uniques). One hundred sixty-five of them were perfect morphospecies, 41 were lumped. Certainly, uniques cannot be split into two or more MOTUs. Therefore, for experimental reasons, uniques have been removed from the data set and again splittings and lumpings were counted. Estimation of species number by MOTUs increased to 33% higher than morphospecies. The number of splittings remained equal, but the number of lumpings decreased in this data set. Not only the agreement between MOTU and morphospecies number decreased, also the proportion of perfect morphospecies decreased (52%).

Perfect morphospecies were mostly uniques (59%). However, 39% occurred with two to ten individuals and two percent with even more than ten specimens. Whereas in the second most frequent perfect morphospecies (Galerucinae sp. 034, 19 individuals) all individuals looked very similar, the individuals of the most frequent one (Alticinae sp. 087, 21 individuals) varied in colour and size.

### Splittings of morphospecies

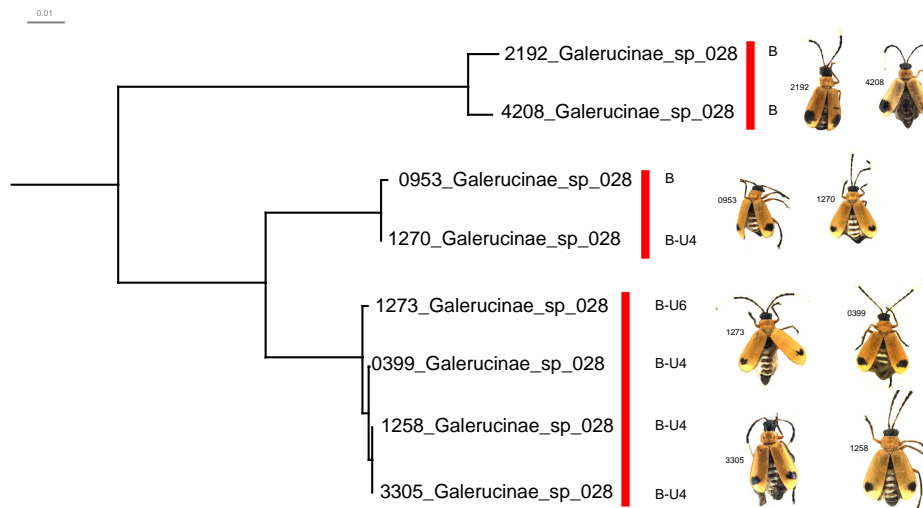


Figure 5.3: **Splitting of the morphospecies Galerucinae sp. 028 into three MOTUs.** Abbreviations behind the individuals indicate the sampling area (B = Bombuscaro), respectively the plot where the specimen was collected.



An example for splitting of morphospecies by the molecular approach is *Galerucinae* sp. 028 that was split into three MOTUs (Fig. 5.3). *Galerucinae* sp. 029 was even split into four MOTUs (Fig. 5.4). All specimens of *Galerucinae* sp. 029 show a conspicuous bicoloured pattern (black and yellow) that is shared by *Galerucinae* spp. 021, 025, and 110 (it is as well found in *Alticinae* sp. 163, see Fig. 5.13). The four morphospecies formed a monophyletic cluster in the molecular analyses.

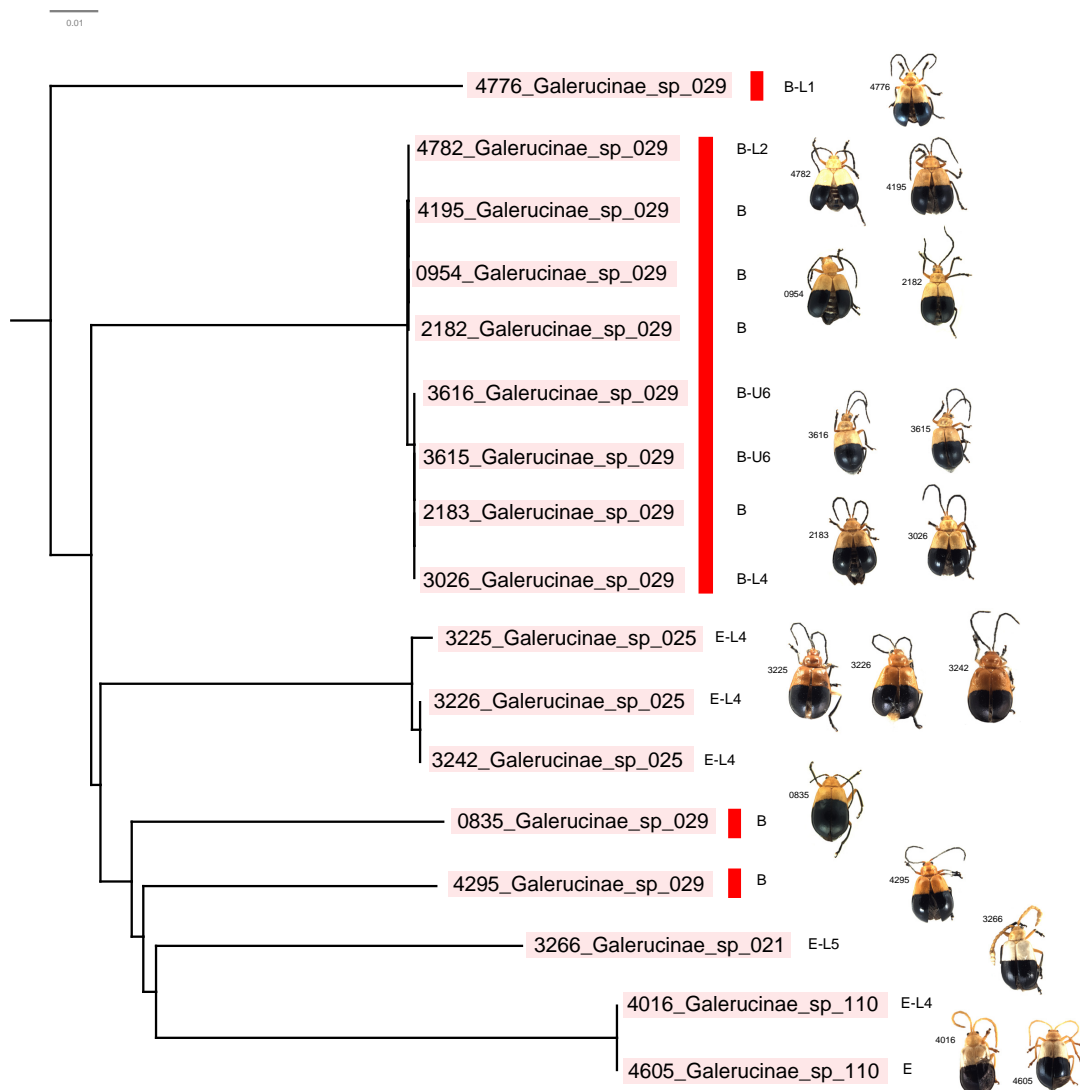


Figure 5.4: **Splitting of the morphospecies *Galerucinae* sp. 029 into four MOTUs.** Abbreviations behind the individuals indicate the sampling area (B = Bombuscaro, E = ECSF), respectively the plot where the specimen was collected.

Alticinae spp. 051, 052, and 152 formed a monophyletic cluster with another orange (Alticinae sp. 056), a fawn (Alticinae sp. 213), and three black (Alticinae spp. 063, 103, 194) morphospecies (Fig. 5.5). They were split into six, four, respectively two MOTUs, however other morphospecies were not lumped into these groups.

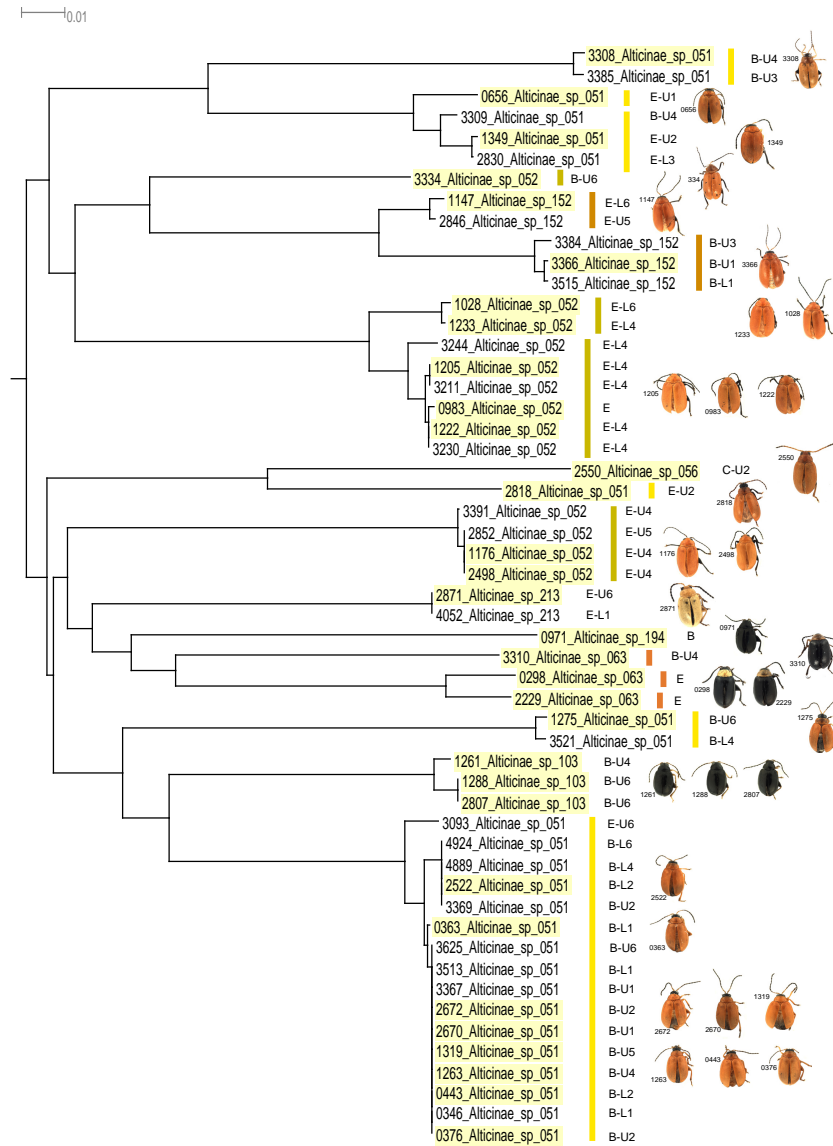


Figure 5.5: Splittings within a group of several orange alticine morphospecies. Coloured bars behind morphospecies names indicate splitting of the morphospecies into more than one MOTU. Abbreviations behind the individuals indicate the sampling area (B = Bombuscaro, E = ECSF, C = Cajanuma), respectively the plot where the specimen was collected.

### Lumpings of morphospecies

Alticinae spp. 017, 048, 054, 071, and 203 that all have different colours belong to the same MOTU (Fig. 5.6). Likewise, Eumolpinae spp. 002, 005, and 014 are lumped by the molecular approach (Fig. 5.7).



Figure 5.6: **Five alticine species with different colours that belong to the same MOTU.** From left to right: Alticinae spp. 017, 048, 054, 071, and 203.



Figure 5.7: **Three eumolpine species with different colours that belong to the same MOTU.** From left to right: Eumolpinae spp. 014, 002, 002, and 005.

Galerucinae spp. 051, 052, 055, 056, and 087 show four different colourations but belong to the same MOTU. Also Galerucinae spp. 015 and 079 with two different patterns belong to one MOTU (Fig. 5.8A, B). Galerucinae spp. 065 and 090 that have different colour patterns belong to the same MOTU but within the MOTU the specimens with the respective pattern clustered (Fig. 5.9). Galerucinae sp. 065 (two individuals) occurs at 2000 m, and Galerucinae sp. 090 (two individuals) at 1000 m. Also the specimens of Galerucinae spp. 092, 024, and 059 belong to the same MOTU but form two distinct clusters (Galerucinae sp. 092 distinct from Galerucinae spp. 024 and 059) (Fig. 5.10). On the other hand, specimens with different colours or patterns may even share the same haplotype e.g. Galerucinae spp. 070 and 084 (Fig. 5.11).



Figure 5.8: **Galerucine morphospecies with different colourations that belong to the same MOTU.** A: Galerucinae spp. 052, 055, 056, and 087 (from left to right). B: Galerucinae sp. 015 (left), Galerucinae sp. 079 (right).

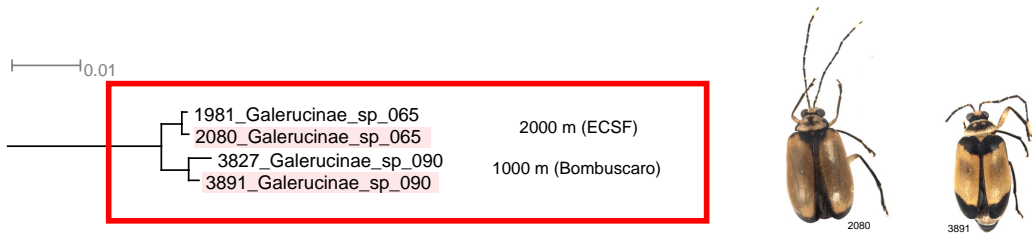


Figure 5.9: **Two galerucine morphospecies with two different patterns that were found at different elevational levels cluster in the same MOTU.**

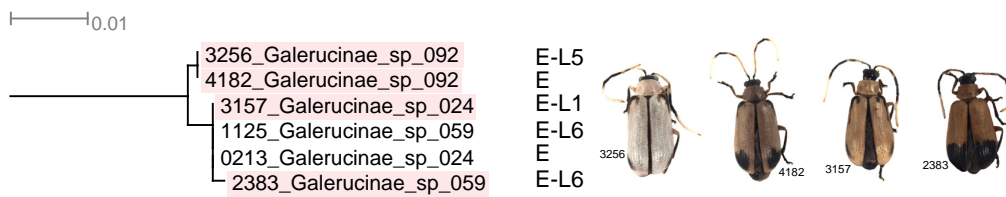


Figure 5.10: **Specimens of three galerucine morphospecies form two distinct clusters within one MOTU.**



Figure 5.11: **Two galerucine specimens show different colouration despite of sharing the same haplotype.** Specimen 0463\_Galerucinae\_sp\_070 (left), specimen 3038\_Galerucinae\_sp\_084 (right).

### Precairious taxa

Several cases of discrepancy between morphospecies and MOTUs were found in a certain group of large Alticinae (~5–11 mm) characterized by a pronotum with a broad margin (Fig. 5.13). Many of them show striking patterns with a partly broad variance between individuals. The group contains morphospecies that were split and lumped at the same time (Alticinae sp. 149) and several cases of colour dimorphism (Alticinae spp. 002, 007, and 218; Alticinae spp. 005 and 163; Alticinae spp. 222 and 155; Alticinae spp. 037 and 039). Noteworthy are the patterns of Alticinae spp. 163, 155, and 222: They are very similar to those of certain galerucine species (Galerucinae sp. 029, respectively Galerucinae spp. 005 and 011; see Figs. 5.4 and 5.14). Alticinae sp. 155 and Galerucinae sp. 005 share an eye-spot pattern. It is to note that an eye-like pattern is also found in some cassidines: Cassidinae spp. 009, 010, and 011 have an eye-like appearance in their general habitus (Fig. 5.12).



Figure 5.12: **Three cassidine species with an eye-like appearance in their habitus.** From left to right: Cassidinae spp. 009, 010, and 011.

A challenging group concerning morphospecies sorting was a group of Galerucinae most probably belonging to the genus *Diabrotica*. They have a similar body-shape and a variety of striking colours and patterns confounding morphospecies sorting (Fig. 5.14). Many splittings occurred in a group of slender Galerucinae that are all dark blue or green and have an orange to yellow pronotum and head (only Galerucinae sp. 032 has a black pronotum and head) (Fig. 5.15). Several of them have also orange legs and antennae.

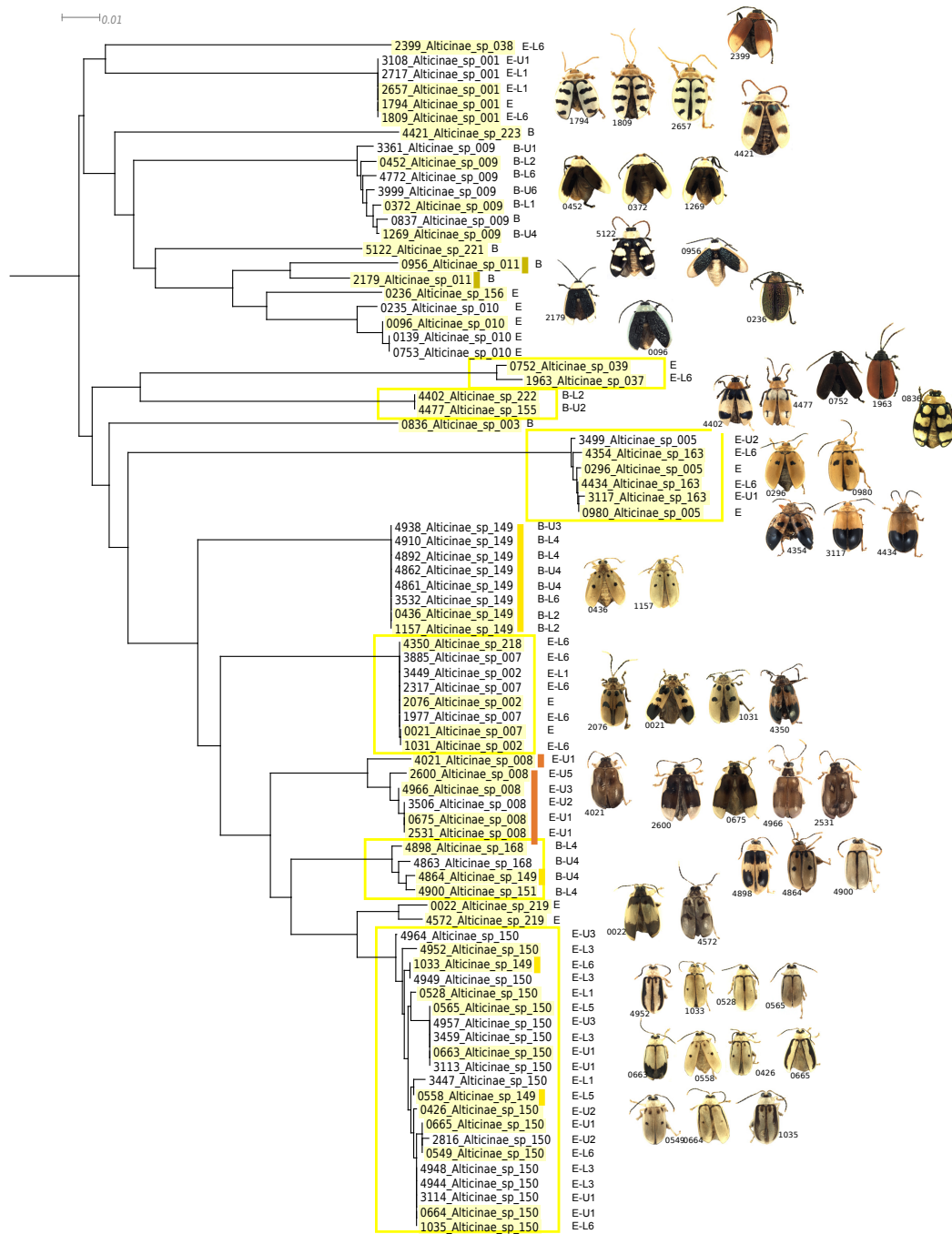


Figure 5.13: Splittings and lumpings within a group of several large alticine morphospecies. Coloured bars behind morphospecies names indicate splitting of the morphospecies into more than one MOTU; morphospecies that are lumped into one MOTU are bordered by a coloured box. Abbreviations behind the individuals indicate the sampling area (B = Bombuscaro, E = ECSF, C = Cajanuma), respectively the plot where the specimen was collected.



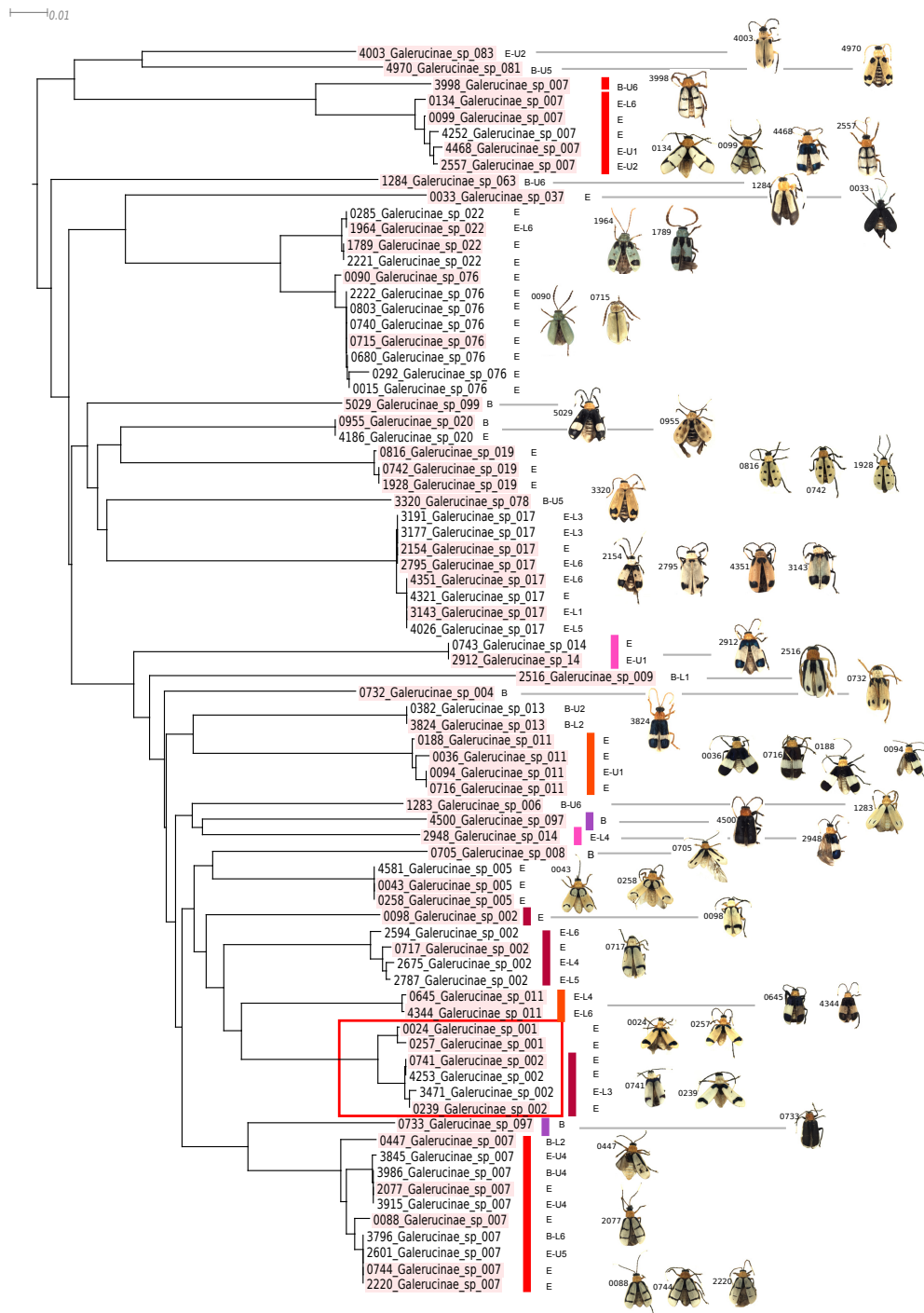


Figure 5.14: **Splittings and lumpings within a group of galerucine morphospecies, presumptively diabroticites.** Coloured bars behind morphospecies names indicate splitting of the morphospecies into more than one MOTU; morphospecies that are lumped into one MOTU are bordered by a coloured box. Abbreviations behind the individuals indicate the sampling area (B = Bombuscaro, E = ECSF, C = Cajanuma), respectively the plot where the specimen was collected.

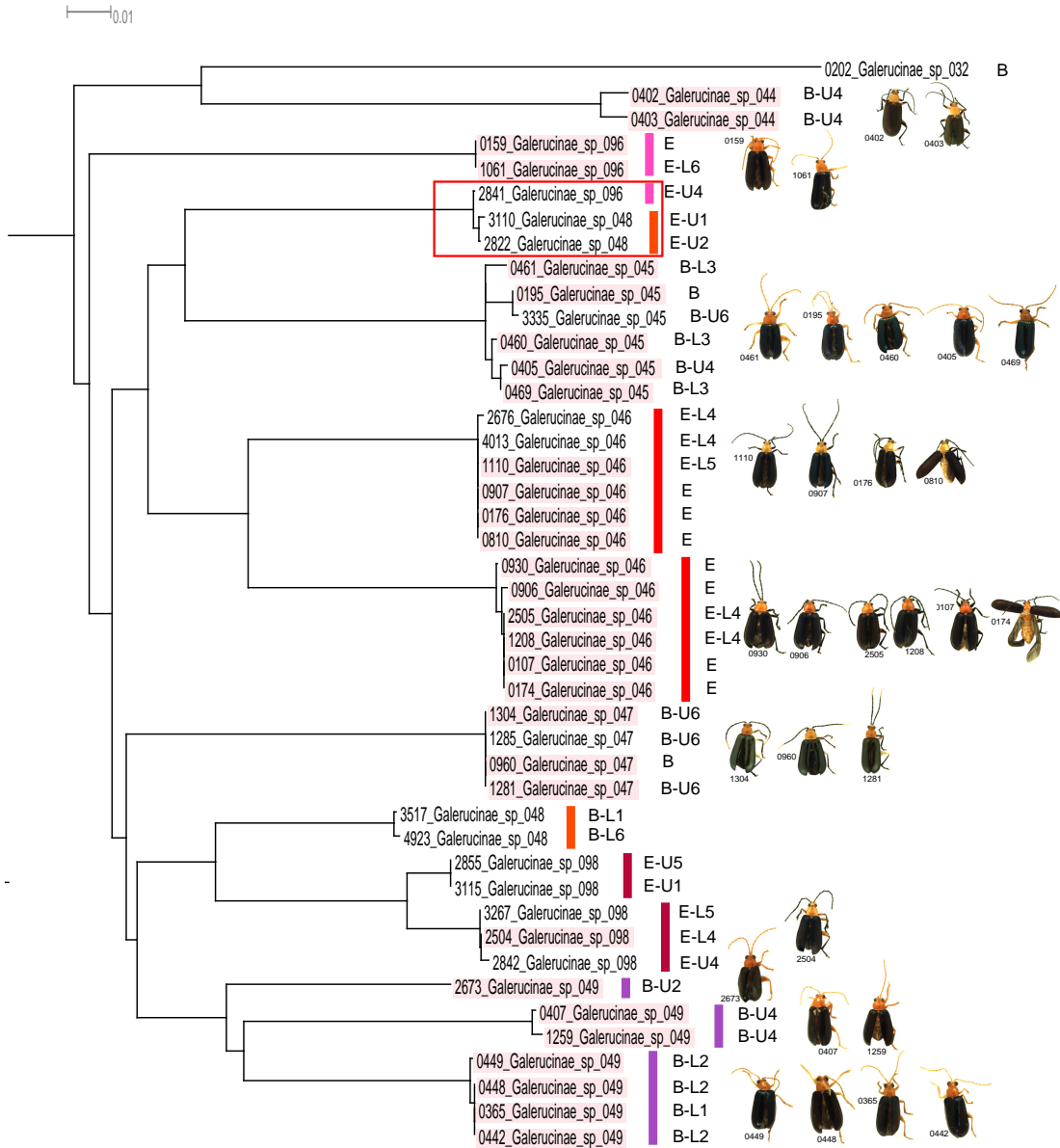


Figure 5.15: Splittings and lumpings within a group of several galerucine morphospecies that are dark blue or green and have an orange to yellow pronotum and head. Coloured bars behind morphospecies names indicate splitting of the morphospecies into more than one MOTU; morphospecies that are lumped into one MOTU are bordered by a coloured box. Abbreviations behind the individuals indicate the sampling area (B = Bombuscaro, E = ECSF, C = Cajanuma), respectively the plot where the specimen was collected.



## 5.4 Discussion

### Splittings of morphospecies

The splitting of morphospecies into two or more MOTUs may arise due to several reasons: cryptic diversity, too liberal morphospecies sorting / a too restrictive molecular approach, mistakes in morphospecies sorting, and mistakes during sequencing or sequence analyses.

*Cryptic diversity:* Cryptic species are two or more distinct species that have been classified as a single species because they are at least superficially morphologically indistinguishable (Bickford et al., 2007; Pfenninger and Schwenk, 2007). In some taxa, cryptic species can be discriminated by differences in mating pheromones or behaviour, e.g. mating calls (Haruyama et al., 2008), however, the increasing availability of DNA sequence data has become a valuable tool for detecting cryptic diversity fuelling research on this phenomenon (Bickford et al., 2007; Hebert et al., 2004). Nowadays, a high proportion of newly described species is being discovered in cryptic complexes (Ceballos and Ehrlich, 2009). Cryptic diversity is widely distributed in most types of organisms and habitats (Bickford et al., 2007) and is also found in Chrysomelidae (Laroche et al., 1996; McKenna and Farrell, 2005; Takano et al., 2011). The unexpected high genetic diversity within species has implications for estimates of biodiversity and potentially for future conservation decisions (Bickford et al., 2007; Witt et al., 2006). In herbivorous insects, cryptic diversity may concern host plants (Blair et al., 2005).

Galerucinae sp. 028 that was split into three MOTUs (Fig. 5.3) might be an example for cryptic diversity, likewise Galerucinae sp. 029 that was split into four MOTUs forming a monophyletic cluster (Fig. 5.4). Cryptic diversity is also found within the orange Alticinae (spp. 051, 052, and 152) that are split into six, four, respectively two MOTUs (Fig. 5.5). Their lack of conspicuous external features seems to hamper the distinction of species.

In most cases of splittings, the resulting MOTUs were next to each other in the NJ-Tree or at least very close, sometimes containing also other morphospecies. In Eumolpinae, Cassidinae, Hispinae, and Chrysomelinae all split morphospecies were split into MOTUs next to each. They are probably sister species. So, usually it seemed to be closely related species that have not been separated by the morphospecies sorting (e.g. Eumolpinae sp. 007, Galerucinae sp. 038, Hispinae sp. 025).

Eumolpinae sp. 024 was split into two MOTUs, one of which (23 individuals) occurred only at Bombuscaro, the other one (two individuals) at ECSF. It is an example for very similar species or cryptic species that occur at different altitudinal levels. They are possibly two recently ecologically diverged species that do not yet have differentiated morphologically.

The term cryptic diversity / cryptic species as used in this study acknowledges that there might be morphological differences but they were not recognized during the applied superficial morphospecies separation. A further taxonomic analysis and dissection of genitalia would possibly resolve those cases.

*Too liberal morphospecies sorting / a too restrictive molecular approach:* Characters of different but related species might be taken as intraspecific variation. Sequence data can help to discover the real significance of slight morphological variation. On the other hand, the molecular species delimitation methods could be too restrictive and overestimate sequence differences, e.g. the genetic distance in COI could be within the range of intraspecific variance. With the applied approach it cannot be told whether the morphospecies or the MOTUs represent true species. The use of another marker could give more information.

*Mistakes in morphospecies sorting:* In this case, specimens have been erroneously grouped into the same morphospecies because differences have not been noticed.

*Mistakes during sequencing or sequence analyses:* Contaminations of DNA-samples cannot be excluded and due to the high number of analysed specimens, the risk of erroneous assignment or denotation of sequences, specimens, and names is given, too. However, all cases with hints at such errors (e.g. conspicuously different individuals sharing the same haplotype) have been rigorously excluded prior to analyses.

### Lumpings of morphospecies

There are several reasons for lumpings of morphospecies into one MOTU: sexual dimorphism, intraspecific morphological variability / polymorphism, too strict morphospecies sorting / insufficient resolution of the molecular approach, mistakes in morphospecies sorting, mistakes during sequencing or sequence analyses. Sexual dimorphism and intraspecific morphological variability / polymorphism may cause erroneous sorting of conspecific individuals into different morphospecies. The molecular method will unite those morphospecies into one MOTU.

*Sexual dimorphism:* Morphological differences between males and females occur in many animal species (Emlen and Nijhout, 2000; Lande, 1980; Poissant et al., 2010; Shine, 1989). Especially a difference in body size (sexual size dimorphism) is a widespread phenomenon (Fairbairn, 2005; Stillwell et al., 2010).

Beetles contain a large number and a great diversity of sexually dimorphic species (Kawano, 2006) ranging from inconspicuous dimorphism to highly developed male traits such as the horns of rhinoceros beetles (Scarabaeidae: Dynastini), the enlarged mandibles of Lucanidae (e.g. *Cyclommatus elaphus* GESTRO) or some Cerambycidae (e.g. *Macrodonia cervicornis* L.), or elongated legs of e.g. long-arm beetles (Scarabaeidae: Euchirinae) (Emlen and Nijhout, 2000; Kawano, 2006).

In Chrysomelidae, there are several forms of sexual dimorphism: Mandibular sexual dimorphism is frequent in leaf beetles (Reid and Beatson, 2013), as well as tarsal specialization (Hammack and French, 2007; Voigt et al., 2008) or modified antennae (Mohamedsaid, 2004). Modified male antennal segments that occur especially in Galerucinae are larger in size and differ from the usual shape of unmodified antennae (Maulik, 1932; Mohamedsaid, 2004). Galerucinae sp. 022 could be an ex-



Figure 5.16: **Different shape of antennae in two individuals of Galerucinae sp. 022.**

ample for sexual dimorphic antennae (Fig. 5.16). Generally, in Galerucinae strongly expressed sexual dimorphism is common with abundant secondary sexual modifications of head, thorax, abdomen and appendages (Mohamedsaid and Furth, 2011; Reid and Beatson, 2013). In contrast, in some subfamilies (e.g. Criocerinae) there is little external sexual dimorphism (Reid and Beatson, 2013).

Certain male modifications in Cassidinae s.l. are associated with behaviour: *Acromis sparsa* BOHEMAN males use elongated corners of the elytra to flip over rivals (Windsor, 1987) and in several hispoid genera a head elongation is used for dislodging rival males (Beaman (1980), cited in Chaboo (2007); Reid and Beatson (2013)).

The sexual dimorphism in Chrysomelidae is often subtle, not affecting the characters considered for morphospecies separation (e.g. mandibles: Reid and Beatson (2013); tarsi: Hammack and French (2007)). Therefore it is less likely to affect morphospecies sorting in this study.

*Polymorphism:* Phenotypic plasticity (the ability of a single genotype to produce more than one alternative form of morphology, physiological state, and/or behaviour in response to environmental conditions; West-Eberhard (1989)) and emerging from this polymorphism (existence of morphologically distinct alternatives in a population; West-Eberhard (1989)) are quite common in insects and also Chrysomelidae (Verma and Kalaichelvan, 2004; Whitman and Agrawal, 2009).

Often, polymorphism concerns colouring and patterns. As in many other insect groups, intraspecific variation in elytral colour pattern is considerable in Chrysomelidae (Verdyck et al., 1996) and although most species of leaf beetles are not polymorphic for body colour and elytral markings, some of them show different morphs (Petitpierre, 1988).

Whereas the frequency of the two colour forms in *Chrysolina aurichalcea* MANNERHEIM correlates with altitude in mountainous areas (Fujiyama, 1979; Fujiyama

and Arimoto, 1988), the colour pattern polymorphism in *Chelymorpha cribraria* FABRICIUS (Cassidinae) with eight different phenotypes is explained by mimicry (Vasconcellos-Neto, 1988). The case of Galerucinae spp. 065 and 090 that have different colour patterns and form two distinct clusters occurring at different elevational levels (Fig. 5.9) does indicate ecological speciation.

Another case where specimens belonging to the same MOTU were assigned into different morphospecies due to colour-polymorphism concerns Alticinae spp. 017, 048, 054, 071, and 203. Due to the different colours of the specimens, the morphological similarity that indicates belonging to the same morphospecies has been overlooked (Fig. 5.6). In a large data set with hundreds of species it is likely to overlook similarities. Barcoding may reveal those similarities that then can be confirmed by a direct comparison of specimens or morphospecies to preclude errors or contaminations. Another example where specimens due to colour-polymorphism are sorted into erroneous morphospecies that are lumped by the molecular approach are Eumolpinae spp. 002, 005, and 014 (Fig. 5.7). Other examples of colour-polymorphism confounding morphospecies sorting are Galerucinae spp. 051, 052, 055, 056, and 087 that show three different colourations but belong to the same MOTU and Galerucinae spp. 015 and 079 with two different patterns (Fig. 5.8A, B).

From published literature other known prominent cases of colour polymorphisms in Chrysomelidae are e.g. *Chrysomela lapponica* L. (Gross et al., 2004; Zvereva et al., 2002), *Plateumaris sericea* L. (Kurachi et al., 2002), *Phyllotreta cruciferae* GOEZE (Verdyck et al., 1996), and *Chrysophtharta agricola* CHAPUIS (Nahrung and Allen, 2005) (Nie et al., 2012). Also the Colorado potato beetle *Leptinotarsa decemlineata* SAY shows polymorphism (Verma and Kalaichelvan, 2004).

It is noteworthy that beside colour polymorphism, also the phenomenon of *colour change* has been observed in Chrysomelidae: Maybe unique among insects, adult cassidines of certain species are able to reversibly change their structural, metallic colours within minutes (Barrows, 1979; Manson, 1929). This behaviour was observed during mating or was caused by disturbance in *Charidotella* (= *Metriona*) *bicolor* FABRICIUS (Barrows, 1979). There is also colour changing in phases during the life cycle, e.g. in *Physonotha helianthi* RANDALL (Cassidinae) (Kirk, 1971). Although most frequently occurring in Cassidinae, there is also colour change in individual species of Chrysomelinae and Alticinae (Buzzi, 1988).

Polymorphism is not only restricted to colouring and patterns. For example there is also wing polymorphism in Chrysomelidae (Furth, 1980) or polymorphism regarding the ability to use an atypical host plant (de Jong and Nielsen, 1999).

*Too narrow morphospecies sorting / a too liberal molecular approach:* Specimens have been assigned to different morphospecies based on differences that only represent intraspecific variation. It can also be the case that COI does not provide sufficient resolution.

In a few cases, when two or more morphospecies were lumped into one MOTU, the specimens of each morphospecies formed separated clusters within the MOTU. Here, a more restrictive molecular analysis (e.g. using a very low distance thresh-

old) would have recovered the morphospecies. However, usually lumped morphospecies did not form distinct clusters when placed in one MOTU. Erroneous sorting into different morphospecies occurred due to colour-polymorphism, overlooking of morphological similarities, or misinterpretation of intraspecific variability. When morphospecies showed a distinct colour-dimorphism or different patterns, this was usually not reflected in the genetic distances. Specimens with different colours or patterns may even share the same haplotype (e.g. Galerucinae spp. 070 and 084; Fig. 5.11). On the other hand, in some cases, morphospecies with different colouring or pattern were lumped into one MOTU but within the MOTU the specimens with the respective pattern clustered, e.g. Galerucinae spp. 065 and 090 (see Fig. 5.9) or Galerucinae spp. 092, 024, and 059 (Galerucinae sp. 092 distinct from Galerucinae spp. 024 and 059; Fig. 5.10).

*Mistakes in morphospecies sorting:* Due to the large amount of species and specimens there is the probability of an erroneous assignment of a specimen into a new morphospecies instead of assigning it to an already existing one. In this case the molecular approach of course lumps those (erroneous) morphospecies into one MOTU.

#### **Precairous taxa**

In the group of large Alticinae with the pronotum with a broad margin (Fig. 5.13), the discrepancies are mainly caused by the striking patterns with a partly broad variance between individuals and several cases of colour dimorphism.

A challenging group concerning morphospecies sorting was a group of Galerucinae most probably belonging to the genus *Diabrotica*. Diabroticites are New World Chrysomelidae that include several significant agricultural pests (Hammack and French, 2007). The diabroticites in the data set had a similar body-shape and a variety of striking colours and patterns confounding morphospecies sorting (Fig. 5.14). Certain patterns are found in different morphospecies and MOTUs, at the same time individuals of one morphospecies or MOTU can show more than one of those patterns.

The manifold splittings within the blue or green galerucines with orange to yellow pronotum and head (Fig. 5.15) can be explained by the similarity of body shape and colouring that complicated morphospecies sorting. Within this group morphospecies sorting maybe was too liberal with features indicating belonging to different species being interpreted as intraspecific variation.

Some patterns of alticine species (spp. 163, 155 and 222) are very similar to those of certain galerucines (Galerucinae sp. 029 respectively Galerucinae spp. 005 and 011; see Figs. 5.4 and 5.14). There have been cases observed where galerucine and crioerine species share the same colouring. This could be explained by mimicry where species imitate toxic or unpalatable species (Balsbaugh, 1988). Eye-spot patterns that are found in Alticinae sp. 155 and Galerucinae sp. 005 are widely distributed among insects and other animals as well (Balsbaugh, 1988). There are several ex-

amples of Chrysomelidae with eye-like markings which could be involved in eye-spot mimicry (Balsbaugh, 1988). Several Neotropical cassidine species have an eye-like appearance in their general habitus as it is found in Cassidinae spp. 9, 10, and 11 (Fig. 5.12; Balsbaugh (1988); Sekerka and Windsor (2012)).

## 5.5 Conclusions

Although biodiversity can be assessed at different levels of classification, the significance of the species as a biological unit is widely recognized and for ecological studies based on invertebrate data species richness and species turnover are important measures (Gaston, 2000; Oliver and Beattie, 1996). Therefore, it is reasonable that the morphospecies as well as the MOTUs are at species-level. In this regard, the morphospecies- as well as the barcode approach are facing the same challenge: to decide where to draw a line between species using character differences. Using sequence data, it must be decided if e.g. differences in genetic distances (Meier et al., 2006) or in branching rates (Pons et al., 2006) are used to delimit species and which values allow assigning of specimens to different species. Different molecular species delimitation methods (Distance-based clustering: Barrett and Hebert (2005); Blaxter et al. (2005); Statistical parsimony analysis: Templeton (2001); Templeton et al. (1992); Generalized mixed Yule-coalescent (GMYC) modelling: Monaghan et al. (2009); Pons et al. (2006); Poisson tree processes (PTP) modelling: Zhang et al. (2013)) are described in Chapter 6 where also their performance and influence on species richness estimates is evaluated. In analogy, the taxonomist or parataxonomist uses morphological characters. As in molecular species delimitation, it must be decided how small or large the differences in those characters must be to assign specimens to the same or to different species.

Exacerbating the morphospecies approach is that external morphological differences are not always categorical (e.g. number of spines) but gradual (e.g. width of pronotum). Furthermore, an experienced taxonomist or parataxonomists has an eye or intuition e.g. for the significance of certain shapes of body or body parts. The problem is that his decisions are not always open to scrutiny and replicable by third persons in the same way (Krell, 2004). In addition, different persons might interpret differences differently: They may be less or more strict, i.e., based on visible differences they may separate or fuse two morphospecies (so-called 'splitters' or 'lumpers'). In contrast, all molecular species delimitation methods, disregarding their particular advantages and disadvantages, have the merit that they are third-party-verifiable and comparable among sites, and they can be applied irrespective of a person's taxonomic knowledge.

Another advantage of DNA barcoding is that it may reveal cryptic diversity that otherwise remains undiscovered. It also performs well in cases where morphospecies sorting is confounded by pronounced intraspecific variation in colour, shape, or size. Additionally, when a huge amount of specimens and species is handled, the probability increases that a specimen is assigned erroneously to a new morphospecies

and it is overlooked that this morphospecies already exists. Especially in species that are small and/or lack conspicuous external characters barcoding is a useful tool to complement morphospecies sorting. Its inclusion, especially in large, specimen- and species-rich data sets is advisable, if possible, as it can facilitate morphospecies sorting and can result into a more accurate species delimitation. The collection and storing of barcode data in taxonomic databases together with information as comprehensive as possible (e.g. photos, sound records) will facilitate future identification. The general accessibility of information is supposed to be an advantage of DNA barcoding (Hebert and Gregory, 2005; Savolainen et al., 2005).

DNA barcode data can be used to infer biodiversity patterns in a similar way as morphospecies data (see Chapter 4) and thus provides a variety of information even without the species being exactly identified and described. Mass-sequencing methods completely rely on sequence data alone and provide a considerable saving of time (Ji et al., 2013; Yu et al., 2012). However, in most cases a combined voucher-based morphospecies and barcode workflow is desirable and necessary. Morphological and molecular approaches should be seen as a feedback loop with both analyses profiting from each other (Page et al., 2005). When an unknown fauna with many probably undescribed species is studied, barcode data might give valuable information about its diversity, but at least in the longer term it is necessary that the species are taxonomically identified and described. This is important as for conservation decisions it is necessary to not only compile species-lists and inventories but also to understand the species' ecology. Although the species' ecology can be studied without valid names, the taxonomic identification, species description, and naming allow the connection with existing knowledge and therefore comments on endemism or threat levels to specific species, for instance (Samways et al., 2010; Schlick-Steiner et al., 2010).

In addition, the voucher-based workflow provides the possibility to check for contaminations or errors in sequencing or naming. It should also be kept in mind that there is the possibility that COI might not be able to discriminate between certain species. It is only one of several possible markers, with advantages and shortcomings (see Chapter 2.4). Sometimes it may be advisable to include other markers than COI into analyses.

It is noteworthy that an advantage of the morphospecies sorting in comparison with barcoding is its possible application for each specimen of the data set (provided that the specimen is not damaged). In contrast, there are cases where for some individuals no barcode can be obtained. It may also be, that when specimens are sequenced individually via Sanger sequencing, usually due to financial restrictions a selection must be chosen and not all specimens can be barcoded. Therefore, in a certain way, the morphospecies approach may also provide additional information not given by the barcode approach, an aspect that also supports the combination of morphospecies sorting and barcoding.

A large amount of singletons/uniques in the data set is disadvantageous for morphospecies as well as molecular approaches: Morphospecies assignment is easier with long series of specimens (Charles and Bassett, 2005) and most techniques for

molecular species delimitation consider rarity insufficiently (Lim et al., 2012).



# Comparison of rapid species delimitation methods and their influence on species richness estimates

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## 6.1 Introduction

Species richness is an important measure in biodiversity research and conservation biology. At a global level, as well as at a local level where diversity of communities is assessed and compared, often the first intuitional question is 'How many species are there?'. Besides, knowledge of species numbers is important for providing a reference point to estimate biodiversity loss (Mora et al., 2011). There is still debate about the total number of species living on Earth. While estimates range from 3.6 to 100 million species, most probable is a number between 5 and 15 million (Chapman, 2009; May, 2000, 2010; Mora et al., 2011; Wilson, 2003). Only a small percentage of them has been formally identified and named (Chapman, 2009; Grove and Stork, 2000; Mora et al., 2011; Wilson, 2003).

It is widely accepted that the most species-rich ecosystems are in the tropics, especially the tropical rainforests (Connell, 1978; Corlett and Primack, 2011; Dirzo and Raven, 2003; Novotny et al., 2006; Wilson, 1988; Wright, 2002), and furthermore, that arthropods with  $\sim 1.1$  million named distinct species make up the largest fraction of all species on Earth (Chapman, 2009). Therefore, efforts to estimate the total number of species are often based on tropical arthropods, especially tropical beetles (Grove and Stork, 2000; Hamilton et al., 2010; May, 2010). Beetles are extremely rich both in functionality and species numbers, making up about one-quarter of all species on Earth (Hunt et al., 2007; Ødegaard, 2000). Since Erwin's (1982) spectacular estimation based on the number of beetle species associated with an individual tropical rainforest tree species, numerous studies led to widely varying estimates of global insect species numbers (Pimm et al., 1995; Stork, 1988, 1993). Those between four and six million arthropod species seem the most probable ones (Basset et al., 2012; Hamilton et al., 2010, 2013; Novotny et al., 2002b). Global and regional insect diversity estimates have been often plant-based, i.e. the number of plant species is multiplied by the number of insect species effectively specialized to them (Novotny and Miller, 2014). Especially herbivorous beetles, including

Chrysomelidae, and their degree of host-specificity have played a fundamental role in species number estimates (Erwin, 1982; Ødegaard, 2000).

At a local level where diversity of assemblages is assessed and compared, the number of species is an intuitive and natural index of community structure (Gotelli and Colwell, 2011). However, despite the familiarity with species richness, analysis of this variable is complex (Gotelli and Colwell, 2011). Species are important biological units and the 'currency' of conservation biology (Agapow et al., 2004). There are several species concepts and debates on how a species should be defined (for overviews see e.g. Mayden (1997); Wägele (2005)). Different species concepts may lead to different species numbers and have potential impact on decisions of conservation management (Agapow et al., 2004). However, it is not aim of this study to delve into this subject and to determine 'real' species. The focus is rather on the practical application of working units that are recognizable and resemble species but do not necessarily correspond exactly to species. Traditionally, morphospecies have been used as such units in studies where detailed taxonomic identification is prevented, e.g. in studies of tropical arthropod diversity (Basset et al., 2008; Longino and Colwell, 1997; Novotny et al., 2002a). Recently, such studies have also profited from DNA barcoding (Janzen et al., 2005, 2009). When an unknown diversity is studied, the interpretation of the sequences is crucial, i.e. how the sequences can be linked to species. There exist several methods for species delimitation, e.g. the use of genetic distances between sequences and a defined threshold to form molecular operational taxonomic units (MOTUs) or clusters, a method which is highly discussed due to the arbitrary choice of a threshold (Blaxter et al., 2005; Hebert et al., 2003a; Meier et al., 2006); statistical parsimony analysis grouping sequences into haplotype networks that are supposed to correspond to species (Hart and Sunday, 2007; Templeton, 2001); the Generalized mixed Yule-coalescent (GMYC)- and the PTP (Poisson tree processes) modelling, a deduction of species boundaries inferred from the data itself by identifying a shift in branching rates between coalescent and speciation (Monaghan et al., 2009; Pons et al., 2006; Zhang et al., 2013). The choice of delimitation method can have a smaller or larger effect on estimates of local and regional species richness.

In this study, a set of different sequence-based species delimitation methods is used to investigate the unexplored leaf beetle fauna in the study area. Resulting predictions of species diversity are compared to estimates from morphospecies sorting. It is evaluated how these different treatments might affect estimates of species richness.

So far, there is no information at all about leaf beetle diversity in the study area or in other regions of Ecuador. In general, leaf beetle diversity is severely understudied in most Neotropical countries (Costa, 2000). More information about leaf beetle research in Ecuador is given in Chapters 1.4 and 3. In contrast to the scarce taxonomic information available for this group, a high actual species richness of Chrysomelidae may be expected: The study area is part of a mega-diverse biodiversity hotspot (Brummitt and Lughadha, 2003; Myers et al., 2000), where climates and habitat types change rapidly along elevational gradients resulting

in a high turnover of communities (Brehm and Fiedler, 2003; Brehm et al., 2003a).

This study should be understood as a first glance on the Chrysomelidae fauna in the studied area and an estimation of species richness and differences between elevations. The focus is on the comparison of the different species delimitation methods. General diversity patterns and especially their change along the elevation gradient is analysed in detail in Chapters 3 and 4.

## 6.2 Methods

### Study area and specimen sampling

Analysed leaf beetles represent a set of beetles sampled in November and December 2010 and between May and August 2011 (data set 4, see Chapter 2.5). Beetles were collected in the Reserva Biológica San Francisco (RBSF) and parts of Podocarpus National Park. Chapter 2.1 gives detailed information about the study area. Sampling was conducted at all three elevational zones (Bombuscaro: 1000 m a.s.l., ECSF: 2000 m a.s.l., Cajanuma: 3000 m a.s.l.); however, Cajanuma was sampled only marginally due to logistical reasons.

Chrysomelidae were collected by standardized sampling with sweep netting, beating, and hand-collection from the vegetation. To complete the overview of species diversity for DNA based species delimitation, standardized sampling was complemented with additional hand-collection and Malaise- as well as light-trapping collections (detailed information about sampling methods and design is given in Chapters 2.2 and 7).

One specimen (in some cases up to three specimens) of each preliminary morphospecies per sample was used for morphological sorting and molecular analysis. More detailed information about laboratory analyses and handling of the specimens is given in Chapters 2.3 and 2.4.

Specimens were sorted into morphospecies on the basis of external morphology but without genital dissection or the use of identification literature. Morphospecies were subsequently provided with the subfamily name and numbered (detailed information about morphospecies sorting is provided in Chapters 2.5 and 5).

Sequences were assembled and edited with Geneious version 5.4.4 (Biomatters Ltd.; <http://www.geneious.com/>) being subsequently aligned using the implemented MUSCLE algorithm (Edgar 2004). The default settings were retained except for the maximum number of iterations (maxiters) that were set to 500. A Maximum Likelihood (ML) Tree was generated in RAxML version 7.3.2 (Stamatakis, 2006) using a GTR+I+ $\Gamma$  model and 5000 bootstrap replicates. Three species of weevils (Coleoptera: Curculionidae; sequences were obtained from GenBank and BOLD) were chosen as outgroup taxa to root the tree (*Anthonomus eugenii* CANO, *Dichromacalles dromedarius* BOHEMAN, and *Acalles camelus* FABRICIUS; Appendix Tab. C.1). They were not included in the further analyses. Branch lengths were made ultrametric with PATHd8 (Britton et al., 2007) using relative ages of nodes and setting the root to an arbitrary age of 1.

### Sequence-based species delimitation

Four different sequence-based methods of species delimitation were used and results were compared with results obtained from morphospecies sorting:

*Statistical parsimony analysis* (Templeton, 2001; Templeton et al., 1992) as implemented in TCS v.1.21 (Clement et al., 2000) (95% connection limit) was used to group sequences into separate haplotype networks. These networks consist of closely related haplotypes connected by mutational paths free from homoplasy with a probability of 95% (Posada and Crandall, 2001; Templeton, 2001). TCS-networks have been shown in various studies to correspond reliably to species across a broad range of taxa (e.g. Ahrens et al. (2007); Astrin et al. (2012); Cardoso and Vogler (2005); Hart and Sunday (2007); Pons et al. (2006); Templeton (2001)). All entities that were given out by TCS were called haplotype networks, even though they may contain only one haplotype or haplotypes that are connected linearly and not necessarily by loops.

*Distance-based clustering* is, despite wide criticism (Cognato, 2006; Meier et al., 2006), widely used as it is fast and easy to apply (Barrett and Hebert, 2005; Blaxter et al., 2005). SpeciesIdentifier v.1.7.7-dev3 (Meier et al., 2006) from the TaxonDNA package (<http://taxondna.sourceforge.net/>) was used to generate clusters of sequences based on pairwise uncorrected distances at user-defined thresholds (function 'Cluster'). All individuals that are connected directly to each other by distances below this threshold are grouped into a cluster (Meier et al., 2006). Clusters may also contain individuals that are connected to each other indirectly, i.e. some distances may exceed the threshold (e.g. A–B: 2.9%, A–C: 2.9%; B–C: 4.8%) (Meier et al., 2006). Different threshold values of 3%, 5%, and 7.5% were tested. As optimal thresholds could not be unambiguously estimated with the underlying data set (Fig. 6.1), only the results of the 3%-threshold are presented. The 3%-threshold has been initially suggested in early barcoding studies by Hebert et al. (2003a) and is often used as standard in insect barcoding (e.g. Hendrich et al. (2010); Smith et al. (2005); Strutzenberger et al. (2011); Tänzler et al. (2012)). It was successfully used to discriminate beetle species of well-known faunas (Astrin et al., 2012; Raupach et al., 2010) and analyses of Papadopoulou et al. (2013) using a combination of mtDNA loci confirmed this value.

*Generalized mixed Yule-coalescent (GMYC) modelling* (Monaghan et al., 2009; Pons et al., 2006) as implemented in the SPLITS package (<https://www.r-forge.r-project.org/projects/splits/>) for the R environment (R Development Core Team, 2009) was used to estimate species boundaries directly from the phylogenetic tree (Monaghan et al., 2009; Pons et al., 2006) produced with COI data. This procedure exploits the differences in the rate of lineage branching at the level of species and populations, recognizable as a sudden increase of apparent diversification rate when ultrametric node height (distance to tips) is plotted against the log number of nodes in a lineage-through-time plot (Nee et al., 1992). Its likelihood is compared then with that of the null hypothesis assuming no shift in branching rate (no separate species), and subsequently the threshold value (time) is estimated which is the cut-

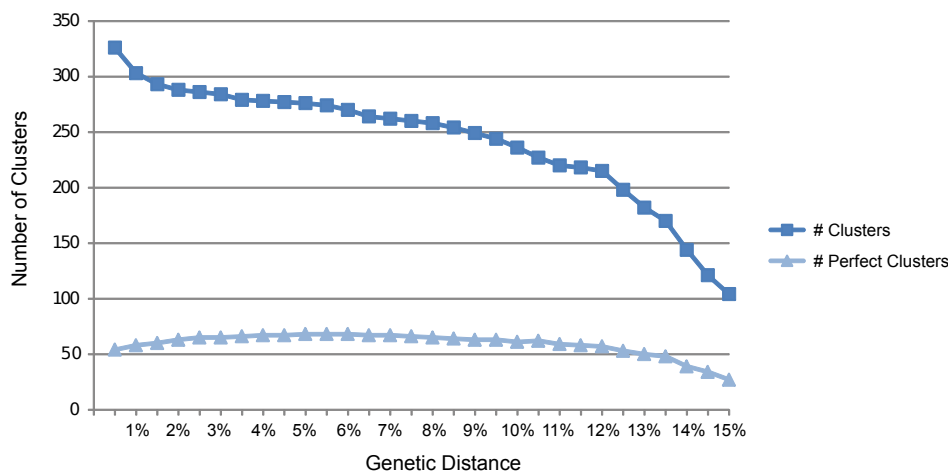


Figure 6.1: **Calibration of distance clusters with morphospecies to determine the best threshold.** Squares = number of delimited clusters, triangles = number of clusters congruent with morphospecies.

off point between speciation and coalescence (Fujisawa and Barraclough, 2013). As the single-threshold value does not differ significantly from the multiple-threshold value, the single threshold value was used for the input tree (Monaghan et al., 2009) which has been already applied successfully to selected groups of organisms (Ahrens et al., 2007; Astrin et al., 2012; Fontaneto et al., 2011; Monaghan et al., 2009; Papadopoulou et al., 2013; Pons et al., 2006).

*Poisson tree processes (PTP) modelling* was used as implemented on the PTP web server (<http://species.h-its.org/ptp/>) (Zhang et al., 2013). This method is similar to GMYC modelling but uses directly the number of substitutions instead of the time to identify branching rate transition points and therefore avoids the potentially error-prone process of making the tree ultrametric (Zhang et al., 2013).

Results of the different molecular species delimitation methods (haplotype networks, distance-, GMYC-, and PTP-clusters) are summed up in the term molecular operational taxonomic units (MOTUs). Additionally, haplotype diversity was inferred as a further independent measure for molecular diversity (Papadopoulou et al., 2011).

### Species richness estimates

For species richness estimates only sweep netting, beating, and hand-collection samples were included, as light-trapping was conducted at Bombuscaro infrequently and Malaise-trapping not at all. The samples from Cajanuma were excluded because the area was significantly understudied. Because the sampled individuals result from the first field trips where the workflow just had to be developed, it was not possible to sample more frequently at Cajanuma due to logistic restrictions. The adverse weather conditions at each of the few sampling trips to Cajanuma contributed to the extremely low number of sampled specimens there.

Species accumulation curves were used to visualize the increase in total species diversity in relation to the number of analysed individuals and to check the completeness of the faunal survey. The method 'random' adds up the samples in a random order with 1000 iterations and calculates the mean  $\pm 95\%$  confidence interval (Fig. 6.3), whereas the method 'collector' adds up samples in the order they appear in the data (Fig. 6.5). The expected total number of species was estimated using *chao2* (Chao, 1987), and first- and second-order Jackknife estimator using the incidence-based estimation provided by the *specpool* function of the R package *VEGAN* 2.0-5 (Oksanen et al., 2012). These are widely used non-parametric estimators that use information on the rare species in an assemblage to estimate the minimum number of species in the assemblage (Gotelli and Colwell, 2011) and have found to perform well in several comparative studies on species richness estimation (Colwell and Coddington, 1994; Walther and Moore, 2005; Walther and Morand, 1998). As different estimators are sensitive to the properties of the assemblage and sampling design, a set of different estimators was used (Samways et al., 2010). Please refer also to Chapter 3.2.

As sampling effort was different between Bombuscaro and ECSF, species richness is hard to compare. To still get an assessment, the number of analysed individuals was standardized to allow comparison of mean species richness: A Jackknifing analysis was performed by randomly drawing 10,000 times 153 individuals (the number of individuals collected at Bombuscaro) from the individuals from ECSF and calculating mean and 95% confidence interval of these samples. For this procedure, the *sample* function of the R base was used within a simple loop.

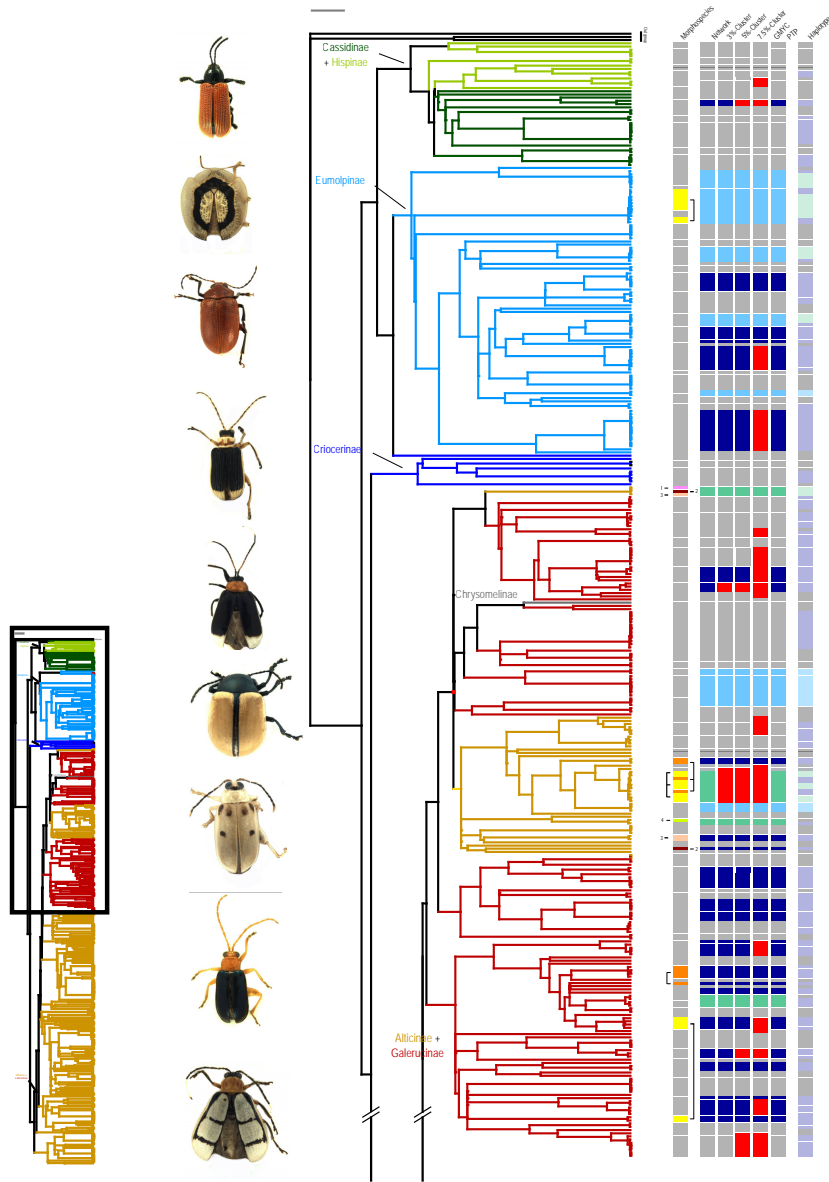
## 6.3 Results

### General results

The 674 Chrysomelidae specimens belonged to seven different subfamilies: Galerucinae s.str. (represented by 163 specimens), Alticinae (371 specimens), Eumolpinae (90 specimens), Cassidinae s.str. (25 specimens), Hispinae (14 specimens), Criocerinae (ten specimens) and Chrysomelinae (one specimen). Specimens showed 426 different haplotypes. Galerucinae + Alticinae (= Galerucinae s.l.), Eumolpinae, as well as Cassidinae + Hispinae (= Cassidinae s.l.) formed monophyletic clusters in the COI Maximum Likelihood tree (Fig. 6.2), only Criocerinae appeared paraphyletic and the chrysomeline specimen was placed within the Galerucinae. This can be ascribed to the inaptitude of COI to resolve phylogenetic groups reliably. Galerucinae s.str. and Alticinae formed several well distinguished clusters within Galerucinae s.l.

### Species delimitation

Morphospecies sorting resulted in a total number of 266 morphospecies. TCS-Network analyses led to a total number of 289 networks and distance-based cluster analyses to a number of 284 clusters. GMYC- and PTP modelling resulted in a



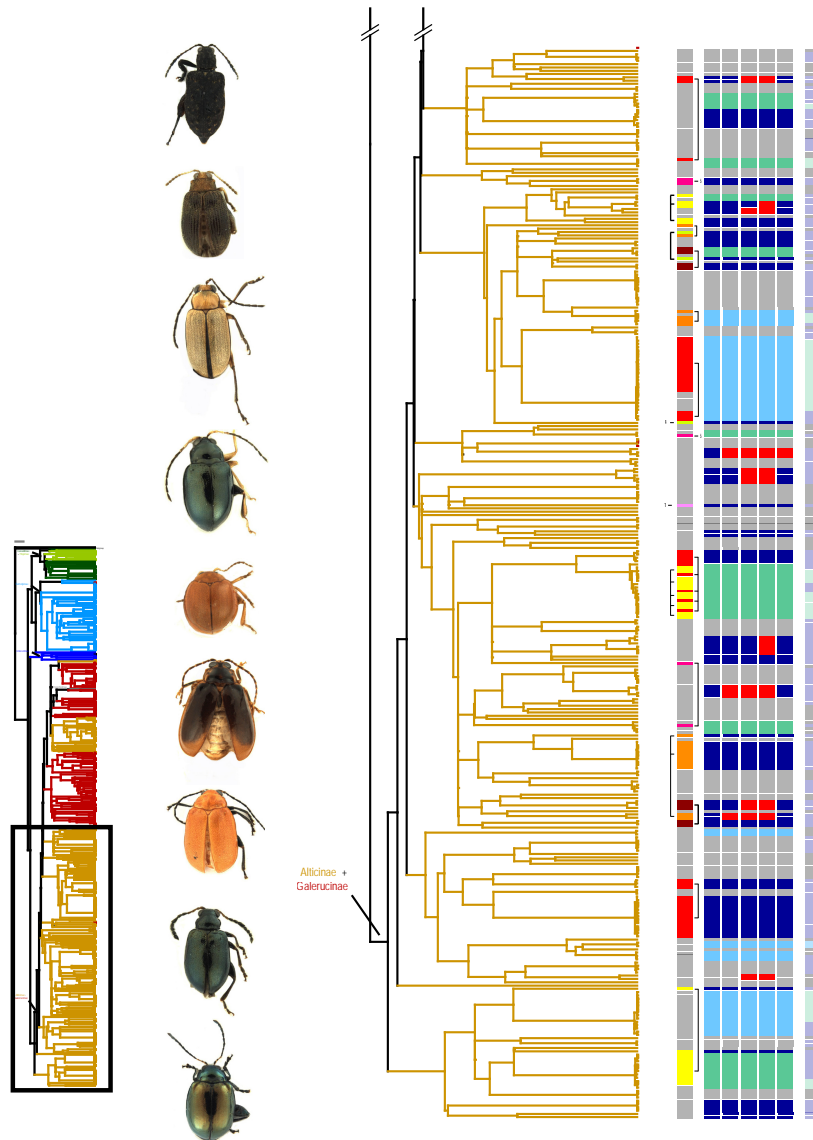


Figure 6.2: ML-Tree providing an overview about morphospecies and MOTUs and differences between the methods. Column 1: Split morphospecies are connected by brackets or numbers and share the same colour. Columns 2-6 + 7: MOTUs (Networks, 3%-, 5%-, 7.5%-, GMYC-/PTP-clusters) and haplotypes splitting a morphospecies are indicated by dark blue fields, those lumping morphospecies by light blue fields, those splitting and lumping morphospecies at the same time by green fields. Red fields indicate differences between the different molecular species delimitation methods.



total of 288 identical GMYC- and PTP-clusters (for results of species delimitation for each specimen see Appendix Tab. E.1).

Despite the high congruence in species numbers, it must be noted that there were several cases of conflicts between morphospecies and MOTUs (Fig. 6.2). These contradictions arise from splitting (in sequence-based analyses a morphospecies is split into two or more MOTUs, respectively parts of them) or lumping events (in sequence-based analyses two or more morphospecies, respectively parts of them, are lumped into one MOTU) (Tab. 6.1). Therefore, despite a high agreement between the number of MOTUs and the number of morphospecies (partially due to the fact that splitting and lumpings compensate one another) perfect congruence was rather low: In total there were 178 perfect matches between morphospecies and networks, 180 between morphospecies and distance-clusters, and 179 between morphospecies and GMYC-/PTP-clusters (see Tab. 6.1).

Table 6.1: Overview of splittings and lumpings.

	Morpho-species	Networks	Distance-clusters	GMYC-/PTP-clusters	Haplotypes
Species number	266	289	284	288	426
Singleton specimens	140	161	156	160	324
Doubleton specimens	94 (47 pairs)	104 (52 pairs)	98 (49 pairs)	102 (51 pairs)	94 (47 pairs)
Congruence with morphospecies number	–	108.65%	106.77%	108.27%	160.15%
# perfect matches morphospecies / MOTUs	–	178	180	179	154
# perfect matches that are not singletons	–	62	65	63	28
# perfect matches relative to # morphospecies	–	66.92%	67.67%	67.29%	57.9%
# Split morphospecies	–	42	39	41	88
# Lumped morphospecies	–	60	61	60	42
# Morphospecies that were both split and lumped	–	14	14	14	18

Splittings and lumpings were almost identical for networks, distance-, and GMYC-/PTP-clusters. For all approaches, the number of morphospecies being split into several MOTUs was higher than the number of cases where several morphospecies were lumped into one MOTU. The congruence between the different species delimitation methods (groups that have been identically delimited by the respective methods) was very high (see Tab. 6.2). There were only five cases of discrepancies where one or another method was more or less restrictive than the others, and there was no case where three methods disagreed, i.e. grouped specimens in three different ways (Fig. 6.2).

The morphological sorting revealed a large amount of singletons in the data set: 140 morphospecies (52.6%) were represented by only one specimen (representing 20.8% of all analysed individuals), 47 (17.7%) by only two specimens (doubletons, 14% of all analysed individuals). Of the 140 singleton morphospecies, 115 (distance-clusters), respectively 116 (networks and GMYC-/PTP-clusters) were also 'molecu-

Table 6.2: **Congruence between the different species delimitation methods.** Shown are the numbers of perfectly matching morphospecies/MOTUs, i.e. groups that have been identically delimited by the respective methods.

	Morpho-species	Networks	Distance-clusters	GMYC-/PTP-clusters
Morpho-species	266	178	180	179
Networks	–	289	279	287
Distance-clusters	–	–	284	280
GMYC-/PTP-clusters	–	–	–	288

lar singletons’, i.e. they were the unique representatives of a MOTU, while 126 were the unique representatives of a haplotype. The remaining 25, respectively 24 singletons were lumped with other specimens into one MOTU. One hundred and sixty-one networks (55.7%), 156 distance-clusters (54.9%), and 160 GMYC-/PTP-clusters (55.6%) were represented by only one specimen; 324 haplotypes (76.1%) occurred only once (see Tab. 6.1).

### Species richness

Sweep netting, beating, and hand-collection samples of Bombuscaro and ECSF resulted in 525 individuals belonging to 219 morphospecies. The species accumulation curve did not reach saturation, suggesting that additional sampling would significantly increase the number of morphospecies (Fig. 6.3). Molecular species delimitation resulted in 241 networks and GMYC-/PTP-clusters as well as 239 distance-clusters represented by 334 haplotypes. The curves of the methods were in their slope similar to the morphospecies curve, none of them showed saturation.

The expected total number of morphospecies estimated with the chao2 estimator was  $413.6 \pm 49.8$  (first-order Jackknife:  $338.2 \pm 21.2$ ; second-order Jackknife: 420.3) while the expected number of networks, GMYC- and PTP-clusters was  $481.1 \pm 56.9$  (first-order Jackknife:  $382 \pm 24$ ; second-order Jackknife: 480.9) and of distance-clusters  $469 \pm 54.9$  (first-order Jackknife:  $377 \pm 23.7$ ; second-order Jackknife: 473). Total number of haplotypes was estimated  $1134.1 \pm 164.1$  (first-order Jackknife:  $585.2 \pm 35.1$ ; second-order Jackknife: 795.5). Leaf beetle communities in the sampled areas of the Podocarpus National Park were estimated to be considerably richer by the molecular approaches than by the morphological one.

As sampling effort was different at the two elevations, the number of analysed individuals was standardized to compare species richness at the two elevational levels. At ECSF, 372 individuals were sampled belonging to 146 morphospecies, 151 networks and GMYC-/PTP-clusters, 150 distance-clusters, and 215 haplotypes. The 153 individuals from Bombuscaro were assigned to 90 morphospecies, 96 networks

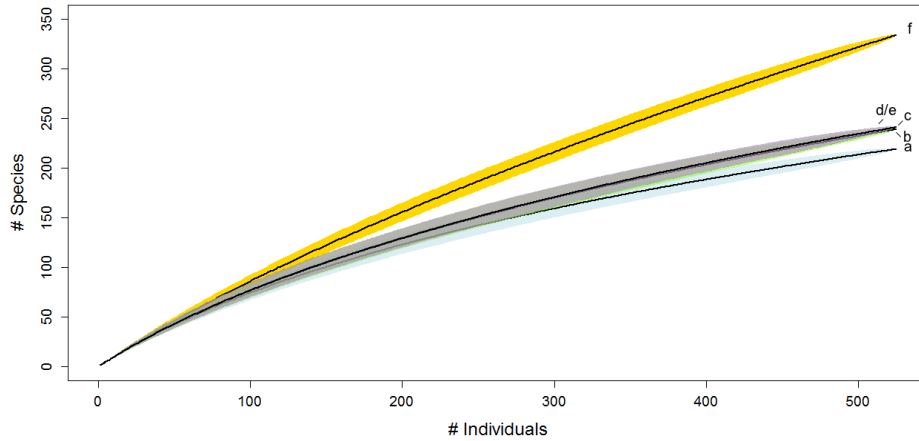


Figure 6.3: **Species accumulation curves.** Increase in the number of morphospecies (a), distance-clusters (b), networks (c), GMYC- and PTP-clusters (d/e), and haplotypes (f) with increasing number of analysed individuals. Coloured polygons indicate 95% confidence intervals.

and GMYC-/PTP-clusters, 95 distance-clusters, and 120 haplotypes. Standardizing the results of Bombuscaro and ECSF to the same number of analysed individuals (153; Jackknifing) revealed no significant difference in mean morphospecies richness between the two areas (Tab. 6.3). The same was valid for networks, distance- and GMYC-/PTP-clusters as well as for haplotype numbers.

Table 6.3: **Comparison of species- and haplotype richness between Bombuscaro and ECSF.** Results standardized with Jackknifing to the same number of analysed individuals (153 analysed individuals from Bombuscaro).

	Species richness				Haplotype richness
	Morphospecies	Networks	Distance-clusters	GMYC-/PTP-clusters	Haplotypes
<b>Bombuscaro</b>	90	96	95	96	120
<b>ECSF</b>	87.9	89.9	89.7	89.9	111.6

The majority of all found morphospecies occurred exclusively at a single elevational level (only 8% occurred at two elevational levels and no morphospecies was found at all three elevational levels; Fig. 6.4). This pattern was even more pronounced when using genetic clusters: Almost all MOTUs occurred at only one elevational level, only 3% at two levels. All haplotypes were restricted to one elevational level. When singletons and doubletons (morphospecies, MOTUs or haplotypes represented by one or two specimens) were removed from the data set results were similar: The percentage of species found at one single elevational level was still the vast majority (80% of all morphospecies and 91% of all distance-clusters and 92%

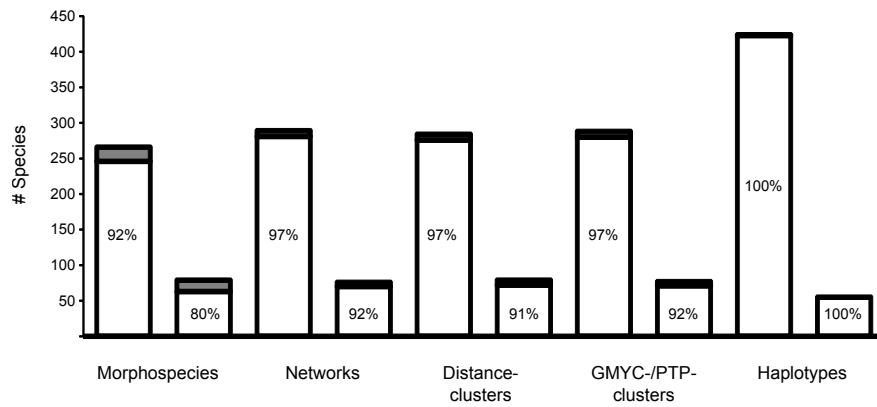


Figure 6.4: **Barplots illustrating occurrence of species at elevational levels.** Morphospecies, networks, distance- and GMYC-/PTP-clusters, and haplotypes found at one (white) or two (grey) elevational levels. Complete data set (left column) for each method and data set without singletons and doubletons (right column).

of networks and GMYC-/PTP-clusters).

The difference in species composition between the different elevations was also reflected in the species accumulation curve of specimens from Bombuscaro and ECSF (Fig. 6.5) which showed for neither of the elevations and none of the delimitation methods saturation. When species from Bombuscaro were added to the data, the slope of the curve steeply increased.

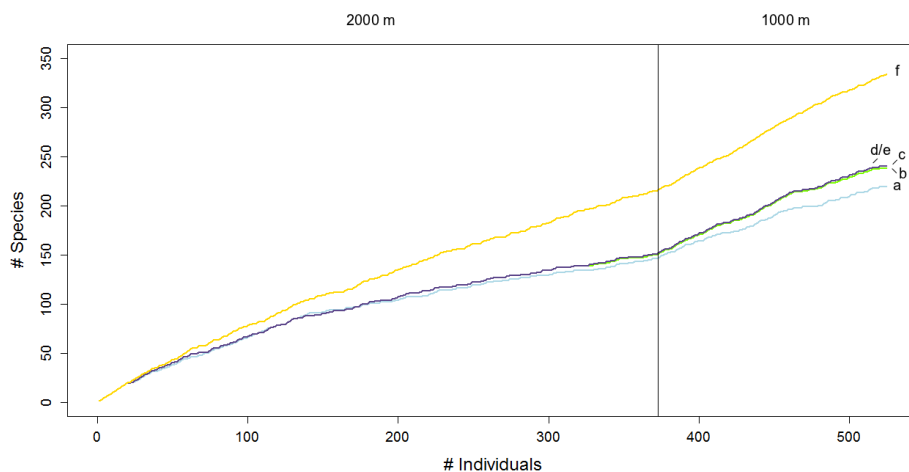


Figure 6.5: **Species accumulation curves for ECSF (2000 m a.s.l.) and Bombuscaro (1000 m a.s.l.).** Increase in the number of morphospecies (a), distance-clusters (b), networks (c), GMYC- and PTP-clusters (d/e), and haplotypes (f) with increasing number of analysed individuals.

## 6.4 Discussion

The study revealed a very diverse Chrysomelidae fauna with observed and estimated species numbers being higher for molecular species delimitation methods than for morphological species sorting. This indicates a significant amount of species that could not be discerned by the morphospecies approach and could contain potential cryptic diversity. Among the different DNA based species delimitation methods, there were only slight differences in observed and estimated species numbers. Species turnover in beetle communities seems to be high between the elevational levels. All findings revealed by MOTUs are similar to morphospecies data, confirming the qualification of DNA barcoding as a tool for assessing biodiversity of an unknown fauna, at least at a geographically restricted scale as in this study.

### Chrysomelid diversity

A high observed and estimated species number was expected as the study area is part of a biodiversity hotspot (Brummitt and Lughadha, 2003; Myers et al., 2000). Even when singletons and doubletons were excluded, most species are restricted to one elevational level indicating a high species turnover.

As frequently found for samples from rainforest communities of insects (e.g. Novotny and Basset (2000); Wagner (2000)), the species accumulation curve did not reach saturation indicating that further sampling would increase the species number. It also lets assume that most species are rather rare, and there is indeed a large proportion of singletons (53% and 55–56% of the morphospecies and MOTUs, respectively). These 'rare species' are an important part of rainforest communities of insect herbivores, often representing from 30% up to more than half of all species in tropical arthropod samples (Coddington et al., 2009; Novotny and Basset, 2000; Wagner, 2000). They may prevent the species accumulation curve from getting saturated even in very large sample series achieved with a huge sampling effort. As the number of specimens included in this study is rather small compared to many tropical arthropod surveys (see Coddington et al. (2009)), the percentage of singletons might decrease with additional sampling effort, but is expected to remain quite high.

It could be expected that the two locations harbour a different chrysomelid fauna, even though the turnover of communities might be overestimated due to undersampling. Mountains have different habitats close to each other as the elevational gradients result in differences in climate, soil, vegetation etc. Although the two sampling areas Bombuscaro and ECSF are as close as ~20 km, there are 1000 m elevation difference and the areas exhibit remarkable differences in climate and vegetation. The turnover of tropical insect communities along elevational gradients is generally rapid (Brühl et al., 1999; Ghalambor et al., 2006; Janzen, 1967) and there are often large differences in insect communities in considerably smaller ranges (e.g. Olson (1994); Smith et al. (2014)).

However, there is no significant difference in mean species richness. A difference could have been expected, as insect species richness often declines with increasing

elevation or shows a hump-shaped distribution (Olson, 1994; Rahbek, 1995, 2005). As *Cajanuma* was considerably undersampled in this study, it was excluded from the comparison of elevational levels. Species richness along the elevational gradient is analysed more detailed in Chapter 4.

The high species numbers found in this study illustrate how understudied Ecuador is when compared with the records listed by Blackwelder (1947): He lists ~450 species explicitly for Ecuador and in contrast 266 morphospecies were found in this preliminary survey that is far from being complete and restricted to a very small area. A comparison with species numbers found in other Neotropical countries (Charles and Bassett, 2005; Flowers and Hanson, 2003; Furth et al., 2003; Linzmeier and Ribeiro-Costa, 2009, 2011, 2012; Ødegaard, 2006; Sánchez-Reyes et al., 2014) or e.g. kept in the collection of the Museo del Instituto de Zoología Agrícola 'Francisco Fernández Yépez' (MIZA), Venezuela, also suggests a high discrepancy between recorded and actual species numbers. It should be noted that mainly one kind of habitat was sampled, the herbaceous and shrubby vegetation in a local forest including only small trees up to a height of ca. 2.5 m. Malaise- and light-trapping was performed additionally, adding on specimens with different ecology (flying and/or nocturnal species). So, besides further sampling with the same methods, the inclusion of other habitats would add more species. For example, a thorough sampling of the canopy would probably increase species numbers by far as a considerable part, perhaps the majority, of the arthropod diversity of tropical rainforests lives in the canopy (Basset et al., 2007; Didham, 2002; Didham and Fagan, 2003; Erwin, 1982) with communities distinct from understorey (Charles and Bassett, 2005). In Chapter 3 the leaf beetle species richness of the study area is compared more detailed with existing taxonomic information and with studies in other regions.

### **Implications of DNA barcoding on species richness estimates**

The successful application of DNA based species delimitation to the studied leaf beetle fauna is not surprising, as it has been proven a reliable method for identification, detection and delimitation of species for a broad variety of taxa, including beetles, in numerous studies (e.g. Astrin et al. (2012); Kubisz et al. (2012); Papadopoulou et al. (2013); Raupach et al. (2010); Tänzler et al. (2012)). It was able to indicate distinct clusters of sequences across all subfamilies of Chrysomelidae of this study, which is an important premise if a large assemblage of unknown species is to be studied. In all analyses the species numbers inferred by molecular methods were considerably higher than morphospecies numbers. Therefore, molecular methods of species delimitation should be included in biodiversity studies, as the morphospecies approach alone may considerably underestimate species richness. These molecular data are a very effective tool for taxonomists for species delimitation and descriptions.

Statistical parsimony analysis, distance-based clustering, and GMYC- and PTP modelling were compared and validated empirically. The high congruence among these different DNA based species delimitation methods indicates a minor relevance of the choice of the particular delimitation methods, at least when sampling as in this study is geographically restricted (but see Bergsten et al. (2012)). A geo-

graphically complete sampling of a species is usually very time and labour-intensive and, therefore, beyond the scope of most ecological studies at the species community level. Often, populations or locations are isolated, either naturally or induced by the progressive fragmentation of habitats, preventing a comprehensive covering of the complete diversity. This is even more valid for tropical insects, where a complete inventory of a certain area is, even if desirable, unachievable, as tropical species in general are high in numbers, but rare and often very localized (Kricher, 1999; Novotny and Basset, 2000). While Lim et al. (2012) argue that this bias may hamper semi-automated DNA based species delimitation, however, the congruence of results of the different delimitation methods used, seems to demonstrate the opposite. Despite a high percentage of singletons and doubletons the species richness estimates remain robust.

### **Haplotypes as a measure for diversity**

Although biodiversity is usually measured in species, the entire genetic diversity of a species, including the diversity of haplotypes, is crucial for conservation. The use of haplotype diversity seems to be an even more objective measure for biodiversity as it is completely independent from species concepts or delimitation methods including their assumptions (García-Lopez et al., 2013; Monaghan et al., 2009; Papadopoulou et al., 2011). Therefore haplotypes are in these analyses an independent estimator and a proxy for diversity in concert with DNA based species delimitation. It has been shown that mtDNA barcode accumulation curves lead to similar results as curves generated using morphology or nuclear genetic markers (Smith et al., 2009). Likewise, in the present study the haplotype accumulation curve was similar in shape to those based on morphospecies and MOTUs and differed only in scale. Therefore, 'haplotype diversity' can be a valuable tool for comparing diversity at a finer scale, which also allows the analysis of diversity of taxonomically unknown organisms, being transparent and reproducible and can be compared among sites (Smith et al., 2009). The distribution of haplotypes within species at different elevations with a strict restriction of haplotypes to a single elevation suggests a separation of populations that are occurring at different elevational zones. Thus the barcodes contain additional information compared to MOTUs or morphospecies. The barcodes are unique identifiers that allow the discovery of a specific haplotype in different samples. However, it should be expected that they will be more informative if applied in a wider geographical scale with much more extended intraspecific sampling (Papadopoulou et al., 2011).

## **6.5 Conclusions**

This study provides a rapid biodiversity assessment of the hitherto unstudied leaf beetle fauna of the understorey vegetation of a tropical montane rainforest in Ecuador. Based on a comparatively small data set compiled in only five months of field work, it revealed a remarkable diversity of Chrysomelidae in the study area and is a good

starting point for future, more detailed research on this fauna. Both morphospecies and DNA barcode data suggest a high turnover along the elevational gradient that is studied more detailed in Chapter 4.

Whereas the integration of different DNA based approaches for estimating species richness is strongly recommended (Carstens et al., 2013), the choice of the molecular species delimitation method seems at least with this data of minor relevance. All results illustrate the high potential of DNA barcoding for exploring communities of hyperdiverse taxa even before being taxonomically identified and formally described (Pons et al., 2006). It can be a useful complement to morphological approaches due to its capability of revealing cryptic diversity, and an effective tool for taxonomic species delimitation and description. Also in cases where experts are not available, applications of DNA barcoding are a suitable method. Nevertheless, an accurate taxonomic description with binary Linnaean names is highly desirable. An advantage of biodiversity assessment with DNA barcoding is that the results are verifiable and comparable among studies and sites. This is an important requirement if barcoding is used as a tool for direct biodiversity measurement. However, it should be established how far the results can be extrapolated also for other organismic groups and larger spatio-temporal scales.



# Comparison of sampling methods

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## 7.1 Introduction

The current biodiversity crisis is considered as a global mass extinction event (Myers (1993) and references therein; Brook et al. (2006); Dirzo et al. (2014); Myers (2003)). Biodiversity loss proceeds at a worrisome rapid pace. Most species are not described yet and there is even no ultimate consensus on total species numbers on Earth. Therefore, exact values of extinction should be considered with caution, but rates seem to be several hundred times their pre-human levels (Pimm et al., 1995). Tens of thousands of species are likely to go extinct per year (Myers, 2003). Probably alone from tropical forests, two to five species disappear per hour (Singh, 2002).

As most biodiversity is found in the tropics, also with regard to biodiversity loss these regions can be considered hotspots (Bradshaw et al., 2009; Dirzo and Raven, 2003; Laurance, 1999). The main cause of tropical biodiversity loss is habitat destruction (Bradshaw et al., 2009; Pimm and Raven, 2000). Especially tropical forests that are extraordinarily species-rich and ecologically complex are subjected to a multitude of threats and are disappearing at alarming rates (Dirzo and Raven, 2003; Laurance, 1999; Laurance and Peres, 2006; Pimm and Raven, 2000). Tropical cloud forests face many of the same threats as other tropical forests. However, their unique ecology and their location on mountain slopes make them particularly susceptible to climate change (Bubb et al., 2004).

In view of those circumstances, there is a necessity for faster methods of biodiversity assessments. One way is to postpone the time-intensive taxonomic identification and description of species. Instead of valid species, often morphospecies are used that are identified by external morphology but without dissection or use of identification literature. They have established in tropical arthropod research to handle the huge amount of accruing specimens and species (detailed information about the morphospecies approach is given in Chapters 2.5 and 5).

In addition, DNA barcoding can help to identify and discover species and has become a valuable method for discovering cryptic diversity (e.g. Hebert et al. (2004); Johnson et al. (2008); Lara et al. (2010); Witt et al. (2006)). Generation of DNA barcode data with standard laboratory protocols has become very efficient (Hajibabaei et al., 2005; Ivanova et al., 2006; Kneibelsberger and Stöger, 2012). With metabarcoding of mass samples even more sequences can be compiled with small time effort (Ji et al., 2013; Yu et al., 2012).

However, also the collection of study organisms in the field as a fundamental step of biodiversity research should be streamlined and must therefore be carefully

planned. Sampling methods and sampling design must be appropriate for the studied organisms, habitats, and the aims of the study. The volume of the studied organisms (number of individuals and species) depends directly on sampling effort. Often a sampling as comprehensive as possible is desired, e.g. for inventories. However, tropical arthropod samples usually remain incomplete, even in large-scale studies. This is typically due to the large number of rare species which can cause analytical problems (Coddington et al., 2009; Novotny and Basset, 2000). For certain questions a statistically relevant number of sampling replicates is needed and the choice of sampling time or period has to be considered.

This study focused on the leaf beetles of the low and medium understorey vegetation up to a height of ca. 2.5 m, including grasses, herbs, shrubs, and small trees in a montane rainforest in Ecuador. Although the leaf beetle fauna of the respective area has not been studied yet, it was not the aim to make a complete inventory. The focus was rather on the analysis and comparison of communities at different habitats and different elevational levels as a test for the application of rapid assessment methods in tropical habitats.

## 7.2 Methods

Chrysomelidae were sampled in RBSF and adjacent parts of Podocarpus National Park during November and December 2010 and from May 2011 to April 2012. The study area is described in Chapter 2.1.

Mainly three standardized sampling methods have been used for collecting leaf beetles from the lower vegetation on the study plots (*plot sampling*; for details of the study plots see Chapter 2.2): sweep netting, beating, and standardized hand-collection (picking up beetles from vegetation). All kind of vegetation within reach was sampled (up to ca. 2.5 m).

*Sweep netting*: Sweep netting was carried out using a standard insect net with 30 cm diameter on a pole. The net was emptied into a bottle containing ethanol. Sweep netting was made alongside two edges of a plot for 30 min (edge of a plot = 20 m).

*Beating*: Vegetation was jarred with a stick to dislodge insects alongside two edges of the plot for 30 min. Insects falling on a horizontal beating tray made of canvas were collected individually from the tray with an aspirator or funnelled into an ethanol-filled collecting jar.

*Standardized hand-collection*: Insects were individually picked from vegetation either with an aspirator or directly into an ethanol-filled jar. This was carried out within the plot for 15 min by a team of two collectors (= 30 sampling minutes).

Plot sampling has been supplemented with the following methods:

*Non-standardized hand-collection*: Additional picking up of beetles from vegetation outside the sampling plots was carried out, e.g. on the way to or from the plots.

*Malaise traps*: Malaise traps are open-sided tents with a collecting head filled with ethanol in which flying or crawling arthropods are trapped (Furth et al., 2003). Bi-directional Malaise traps have been used and collecting heads were usually emptied

after three to four weeks. In some cases the interval was shorter or longer.

*Light trapping:* For light trapping, a light tower with two 15 W tubes (one black light, one superactinic) was used.

Occasional *flight interception traps* with ~40 x 60 cm panels were placed in trees at ~5 m height and emptied after four weeks. On trial *pitfall traps* were emptied after two days.

Being killed and collected in 70% ethanol, beetles were subsequently transferred into 96% ethanol and stored at 4 – -20 °C. A total of 662 samples has been selected to be analysed. For each of these samples, Chrysomelidae were sorted into preliminary morphospecies. One specimen of each preliminary morphospecies was dry mounted, labeled, and sorted into morphospecies (more detailed description of morphospecies sorting is given in Chapters 2.5 and 5). The remaining specimens (*duplicate specimens*) were included in the comparison of sampling methods (data set 2a). The sorting of specimens of a single sample into preliminary morphospecies is quite reliable due to the small number of individuals and species per sample. Therefore, it is likely that similar specimens were correctly classified as the same morphospecies. Furthermore, the number of affected specimens is quite low as usually (~80%) only one individual per morphospecies was found in one sample. More detailed information about selection and processing of specimens and about the different data sets is given in Chapters 2.3 and 2.5.

## 7.3 Results

Within the present study a total of 1174 samples (refer to Chapter 2.2 for definition of sample) were taken. They varied considerably in size and effort as different sampling methods were used. A total of 4286 Chrysomelidae was collected. As the focus of the study was the community analysis of the study sites (plots), most samples have been taken on these plots: 306 samples of each standard collection method yielded a total of 2364 leaf beetles. Of these specimens, 1091 came from the 306 sweep netting samples, 980 from the 306 beating samples, and 293 from the 306 standardized hand-collection samples. Further 947 specimens were collected with 134 non-standardized hand-collection samples, 816 specimens with 45 Malaise trap samples, and 114 specimens with 27 light trap samples. Flight interception traps (32 samples) and pitfall traps (18 samples) yielded very few specimens (43, respectively two) and are disregarded hereafter (Fig. 7.1).

For the following comparison of sampling methods the same data set as for the general biodiversity analyses in Chapter 3 was used (data set 2a; for explanation of the data sets see Chapter 2.5): It contained 2227 specimens from 662 samples. The samples were biased towards sweep netting, beating, and standardized hand-collection samples (plot samples). Number of analysed samples, collected specimens, and identified morphospecies for the different methods is given in Tab. 7.1. Fig. 7.2 illustrates the proportion of samples and specimens. For morphospecies found by the respective method the composition is similar to the composition of collected

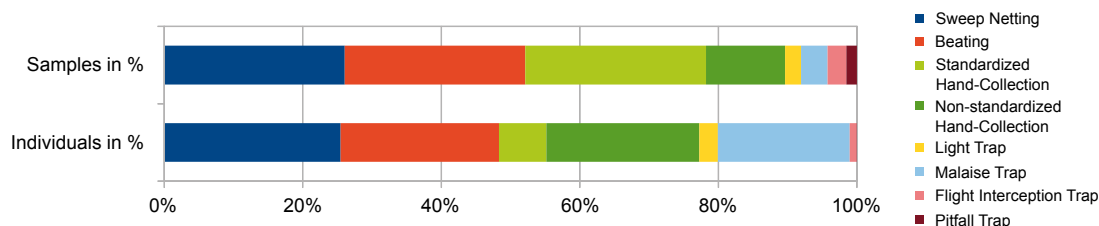


Figure 7.1: Proportion of all within the project taken samples and yielded individuals for the different sampling methods. Note the comparatively high number of individuals in non-standardized hand-collection and Malaise trap samples.

specimens: Sweep netting covered 54% of all 473 morphospecies, beating 45%, hand-collection (standardized and non-standardized hand-collection combined) 47%, light traps 6%, and Malaise traps 12%. Flight interception and pitfall traps covered  $\leq 1\%$ .

Table 7.1: Comparison of the number of analysed samples, collected specimens, and identified morphospecies for the different methods.

Method	# Samples	# Individuals	# Morphospecies
Sweep Netting	199	749	255
Beating	199	635	214
Hand-Collection (standardized & non-standardized combined)	233	597	222
Light Trap	20	54	27
Malaise Trap	6	187	58
Flight Interception Trap	3	3	3
Pitfall Trap	2	2	2
Total	662	2227	473

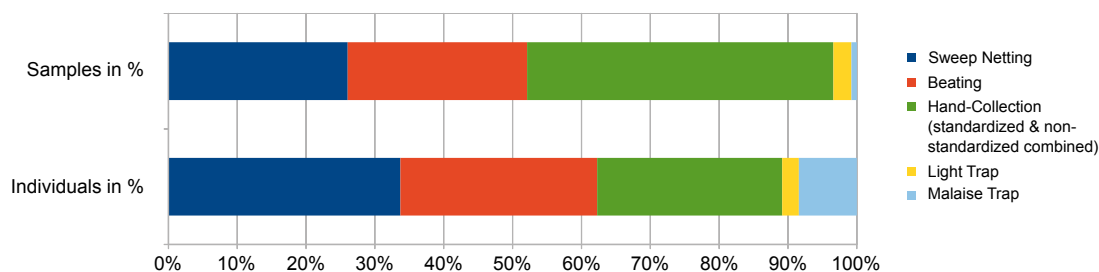


Figure 7.2: Proportion of taken samples and corresponding proportion of number of specimens collected with the different methods for data set 2a.

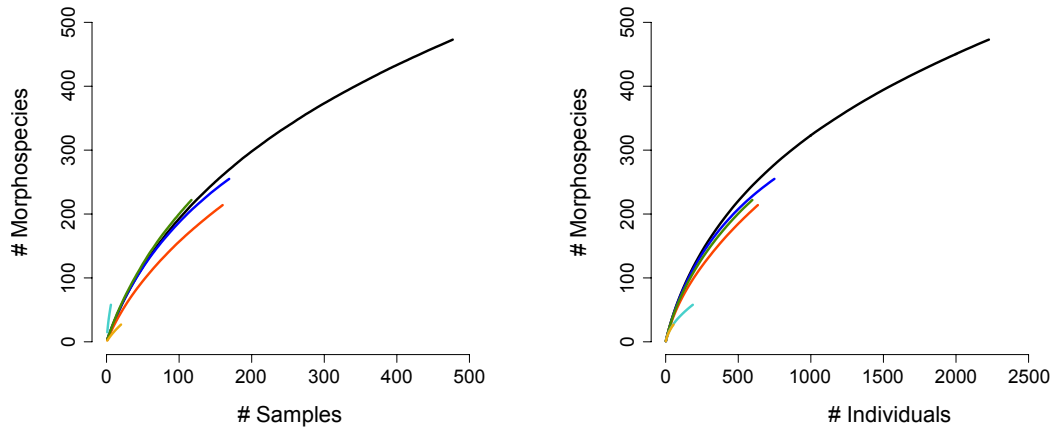


Figure 7.3: **Species accumulation curves showing the correlation between morphospecies number and number of samples (left) and sampled individuals (right) for all five sampling methods separately and for all methods combined.** Sweep netting (dark blue), beating (red), hand-collection standardized and non-standardized combined (green), light trap (yellow), and Malaise trap (light blue) samples, addition of all methods (black).

Species accumulation curves (Fig. 7.3) illustrate how the number of morphospecies grows with increasing number of samples (left), and individuals (right) for each sampling method and for all methods combined. In contrast to the sample based curve, the individual based curve shows for all sampling methods combined a higher efficiency in collecting the leaf beetle fauna of the studied area than each single method. The figures show that Malaise traps collected more morphospecies per sample than all other methods taken together whereas they were among the least efficient methods when based on number of individuals. None of the curves shows saturation.

The individual samples varied considerably in the number of sampled specimens and morphospecies (Fig. 7.4). In sweep netting, beating, hand-collection, and light trap samples, on average only few individuals and morphospecies were caught per single sample (less than five). The maximum in single standard samples was 28 specimens, in the very heterogeneous non-standardized hand-collection samples even up to 68. In contrast, the Malaise trap samples yielded a mean of 31.2 sampled individuals and 14.8 morphospecies per sample.

Regarding the sampling methods sweep netting, beating, hand-collection (standardized and non-standardized combined), light-, and Malaise trapping, more than half of all morphospecies (59%) were sampled by only one method. Twenty-three percent were sampled by two, 14% by three and only four percent by four methods. Only two morphospecies were sampled by all five methods. This is partly due to the high proportion of 'uniques': 49% of the morphospecies occurred in only one sample (were sampled only once).

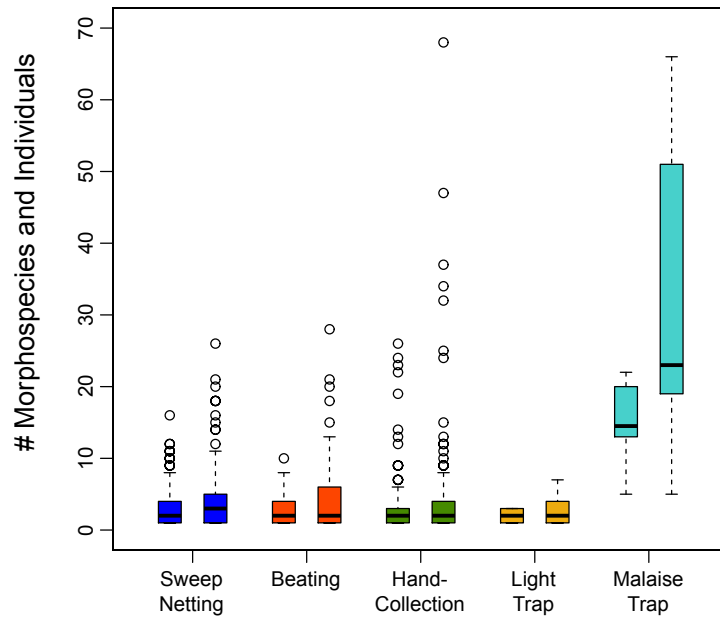


Figure 7.4: **Number of morphospecies and individuals per sample.** Found morphospecies (left column) and individuals (right column) per sample for sweep netting, beating, hand-collection, light trap, and Malaise trap samples. The boxplots show the median, the lower and upper hinge, the minimum, and the maximum.

The presence of the most frequent subfamilies accounting for 97% of all morphospecies revealed slight differences in the collection efficiency of different methods for certain subfamilies (Fig. 7.5). On average, most morphospecies belonged to Alticinae. Of the morphospecies found in Malaise trap samples, even 71% were Alticinae. In contrast, in light trap samples only 30% were Alticinae, whereas more than half of all morphospecies (52%) caught with light traps belonged to Galerucinae. Cassidinae that on average made up four percent of the morphospecies accounted for eight percent of non-standardized and standardized hand-collection samples.

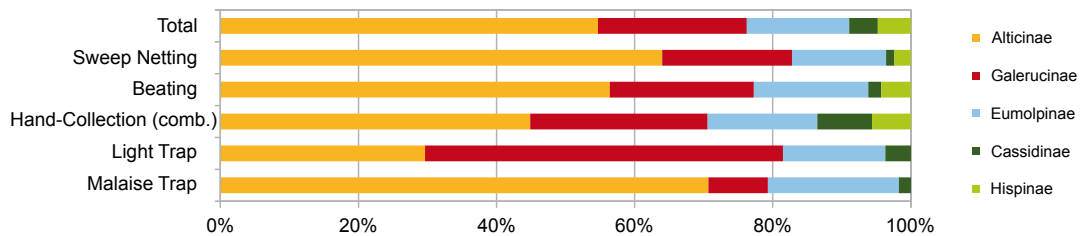


Figure 7.5: **Percentage of collected morphospecies according to subfamily for different sampling methods.**

### Standardized sampling methods

A total of 199 plot sample replicates has been analysed (199 sweep net, beating, and standardized hand-collection samples) resulting in 1578 specimens and 379 morphospecies (data set 3a). Sweep netting and beating resulted in highest morphospecies numbers (255 respectively 214), whereas only 106 morphospecies were found in hand-collection samples (Fig. 7.6). However, when corrected for the number of individuals (sweep netting: 749, beating: 635, hand-collection: 194), morphospecies richness was marginally higher for hand-collection than for beating samples. The combination of the three methods did not increase the efficiency in terms of collected morphospecies compared to sweep netting alone.

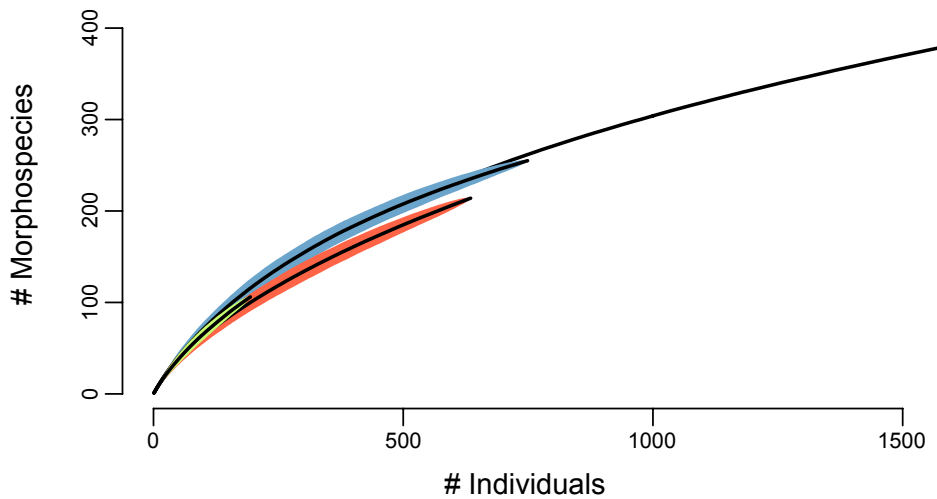


Figure 7.6: **Species accumulation curves for plot samples showing the correlation between the number of morphospecies and the number of specimens, based on 199 samples of each method: sweep netting (blue), beating (red), standardized hand-collection (green); all three methods combined (black line without confidence interval).** The trend for the standardized collection methods is the same as for all methods (see Fig. 7.3 left). Coloured polygons indicate 95% confidence intervals.

Most morphospecies of the plot data set were found by only one sampling method (61%). Twenty-seven percent were found by two methods and only 12% by all three methods. Sweep netting showed the highest percentage of morphospecies that occurred exclusively in samples of this method (45%). Thirty-nine percent of the morphospecies in beating samples and 29% of the morphospecies in standardized hand-collection samples were found only in samples of the respective sampling method. Of the morphospecies that were found by two methods, most occurred in sweep netting and beating samples (73%), 18% were shared by sweep netting and hand-collection, and ten percent by beating and hand-collection.

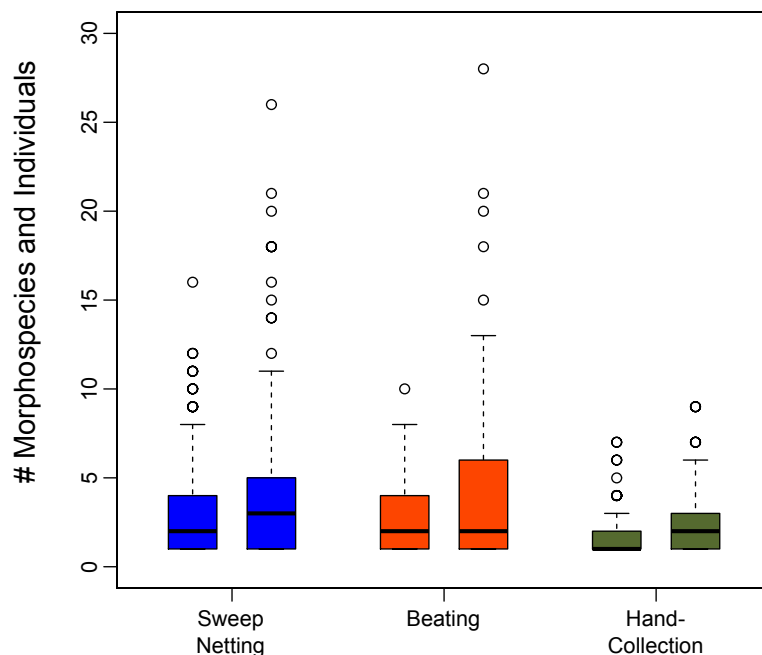


Figure 7.7: **Number of morphospecies and individuals per plot sample.** Number of morphospecies (left column) and individuals (right column) per sample for sweep netting, beating, and standardized hand-collection samples.

For the most part, a morphospecies was represented by a single individual per sample (826 times). In 137 cases, a morphospecies was represented by two specimens in a sample, in 58 cases by three or four specimens, and in 23 cases by five to eight specimens. Only occasionally more than eight specimens of the same morphospecies were found in one sample, maximally 26 specimens. Therefore, the mean number of morphospecies was similar to the mean number of individuals per sample (sweep netting: 3.3 morphospecies vs. 4.4 individuals, beating: 2.7 vs. 4.0, standardized hand-collection: 1.9 vs. 2.3). Generally, the number of individuals and morphospecies caught per sample is very low, however with a very large variance (Fig. 7.7). In sweep netting and beating samples up to 26, respectively 28 individuals could be found. In standardized hand-collection samples maximally nine specimens have been found. Also the maximum number of morphospecies per sample was considerably lower for hand-collection than for sweep netting or beating samples.

The species accumulation curve of the plot samples showed no saturation after 298.5 hours of sampling indicating that a further increase of morphospecies number is expected with further sampling (Fig. 7.8).



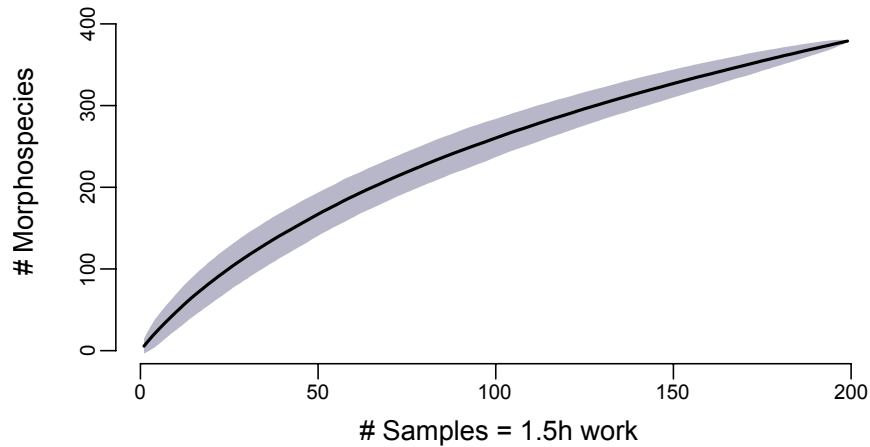


Figure 7.8: **Species accumulation curve (mean  $\pm$ 95% confidence interval) showing the number of morphospecies discovered with increasing number of plot samples.** One plot sample is equivalent to 1.5 hours of sampling: 30 min sweep netting, 30 min beating, and 30 min hand-collection.

## 7.4 Discussion

The main sampling methods (sweep netting, beating, hand-collection) are widely used for sampling of Chrysomelidae in the selected habitat, the lower vegetation of the study plots within the forest (Flowers and Chaboo, 2009; Furth, 2009; Sánchez-Reyes et al., 2014; Staines, 2011). As the comparison of communities at the study sites is an essential aspect of the project (see Chapter 4), an advantage is that the beetles are sampled more or less directly from the vegetation, so their provenance is known. In contrast, when Malaise-, light- or flight interception traps are used, the provenance of the flying insects is not known. These methods should not be used for a fine-scale sampling on sampling sites as close to each other as in this study, where they are sometimes only several meters apart, because samples will not be independent (Ozanne, 2005a). In those cases, especially light traps are not well suited because e.g. moths are attracted from a distance of up to 25 m or more (references in Brehm (2002)).

Comparing the three methods for standard sampling on the plots, sweep netting seems to be the most successful method in terms of collected specimens and species, slightly more efficient than beating (Fig. 7.6). The combination of all three methods did not result into a higher number of found species when corrected for the number of sampled individuals. In contrast, hand-collection on the plots was not very efficient, resulting in considerably lower specimen and morphospecies numbers than sweep netting and beating, in total (less than one third of the specimens and less than half of the morphospecies, Fig. 7.6) and on average (Fig. 7.7). However, when corrected for the number of sampled individuals, this method yields a comparable number of species (Fig. 7.6). The high numbers of individuals and morphospecies for the combined standardized and non-standardized hand-collection samples (Fig. 7.3,

Fig. 7.4) result from cases where extensive hand-collection was conducted under favorable weather conditions and at sites well suited for collection of Chrysomelidae. Although hand-collection on plots was less efficient, it was reasonable to be included to find additional species, as e.g. Cassidinae and Hispinae were mostly found by hand-collection.

Light trapping seems little appropriate for sampling Chrysomelidae as it was not very efficient in terms of specimens and morphospecies. This is probably because in tropical forests the majority of chrysomelids are active during daytime (Basset et al., 2001).

Malaise traps, on the contrary, show a favourable proportion between sample number and number of collected specimens and morphospecies. They seem to be an efficient method for sampling of tropical leaf beetles. They have been widely used, especially for sampling of Alticinae (Flowers and Hanson, 2003; Furth et al., 2003; Linzmeier and Ribeiro-Costa, 2008, 2009, 2012, 2013). Whereas Malaise trapping is less efficient than sweep netting, beating, and hand-collection, when based on the number of individuals, it is the most efficient method considering collected morphospecies per sampling effort (Fig. 7.3). However, compared to the other methods, Malaise trapping often generated larger numbers of a few common morphospecies, which resulted in a lower species richness per collected individuals (Fig. 7.4). The fact that Malaise traps collected the highest mean number of individuals and morphospecies per sample is not surprising because of the longer collecting time of up to ~30 days, meanwhile many insects can accrue in the collecting head of the trap.

Beside the numbers of collected specimens and species, the time effort required is a crucial factor to evaluate the effectiveness of different sampling methods. A disadvantage of the standard methods is the high workload and time requirement compared to the output: Although almost 300 hours have been spent on plot sampling, no saturation of species accumulation curves has been achieved. In addition to the pure sampling time of 298.5 hours, the time to arrive at the sampling plots can be considerable: It takes roughly one hour to arrive at the areas of Bombuscaro or Cajanuma by bus and car. The subsequent walk to the plots may require even more time. So, a team of two persons could accomplish sampling of not more than three plots per day. Although the required collection time for all of the three methods is equal, it is to note that in comparison with beating and hand-collection the processing of the sweep net samples takes a considerable amount of time: The samples contain lots of leaves and other parts of vegetation that must be carefully screened for insects, and sorting out of the specimens requires some effort.

Whereas it costs a considerable effort to carry the light trap equipment to the respective study sites, the Malaise traps are comparatively easy to handle: They can be readily placed at the sampling sites, even at remote sites, and just have to be cleared (preferentially after a few days to ensure good quality of the samples). Both light traps as well as Malaise traps could be used at a larger scale when sampling sites are further apart but are less suitable when sites are close to each other.

The methods seem differently efficient for certain subfamilies. Therefore, a mix of methods seems advisable if whole communities of leaf beetles are addressed, espe-

cially for inventories as it has been shown for ant fauna (Longino et al., 2002). However, the incomplete sampling impedes an exact comparison between the methods: The finding that most morphospecies were found by only one sampling method may be due to the high amount of morphospecies that were found only once (uniques). Therefore, differences in taxon-specific efficiency of collection methods can hardly be documented with the present data. A complete coverage of all species and a saturation of species accumulation curves was not attempted and expected because no complete inventory of the study area was intended.

The data indicate that the area is species-rich, but species usually occur with a low abundance, as it is typical for tropical rainforests. Therefore, it seems advisable to sample at several sites with a set of methods and with a statistically relevant number of sampling replicates over a certain time period. If the number of specimens that can be analysed (morphologically and molecular) is restricted, the decision to analyse only one specimen per morphospecies per sample seems reasonable: As most morphospecies were found only once per sample, the number of omitted specimens is rather low, and leaf beetle diversity of the region still can be characterized well.

Some habitats that have not been addressed explicitly in this study require special sampling methods: Schmidl et al. (2007–2008, unpublished) studied the arthropod fauna living under tree bark in parts of RBSF using an insecticide. There was no opportunity to sample canopy beetles for this project. Especially the canopy of tropical rainforests is known to harbour an extraordinarily high richness of insects in general and Chrysomelidae in particular (Basset et al., 2001; Charles and Bassett, 2005; Farrell and Erwin, 1988; Wagner, 1999, 2000, 2003). However, for Alticinae a redundancy between canopy fogging and Malaise trapping has been shown (Furth et al., 2003). Canopy fauna can be accessed directly from platforms, walkways, canopy rafts, sledges, balloons, towers, or cranes (Basset et al., 2001; Charles and Bassett, 2005; Ozanne, 2005b; Samways et al., 2010). Those structures that allow the application of different collecting methods in the canopy are very sophisticated and costly and are usually operated as part of large projects, e.g. IBISCA ([www.ibisca.net](http://www.ibisca.net); Basset et al. (2007)), or by major research institutes, e.g. the Smithsonian Tropical Research Institute in Panama ([www.stri.si.edu](http://www.stri.si.edu)). Other common methods involve climbing into the canopy or chemical knockdown (fogging or mist-blowing). For knockdown sampling, usually pyrethrum or related substances are used and fallen insects are captured on collecting trays, hoops, or mats (Farrell and Erwin, 1988; Ozanne, 2005b; Samways et al., 2010; Wagner, 2000).

## 7.5 Conclusions

Facing the rapid advance of biodiversity loss in tropical rainforests, an acceleration of biodiversity assessment is indispensable. Whereas molecular methods such as DNA barcoding, with laboratory protocols becoming more and more efficient, have been propagated to accelerate analysis and understanding of biodiversity, the sampling of specimens in the field is still the most important and basal step and can constitute

a bottleneck in the workflow.

In this study, especially the standardized sampling on the plots has been proven to be quite time and work intensive, also because the considerable way to the plots must be included. However, there seems to be no alternative method of more efficiently collecting the leaf beetles from the low vegetation of the plots. Especially sweep netting is an excellent method, however, with the disadvantage that the sorting of the samples takes time. Beating yielded only slightly fewer specimens and species with sorting of the samples being less time-consuming. Because there is no great difference in efficiency between the methods, a combination of both can be recommended. Hand-collection yielded comparatively few specimens, but raises the probability for also catching Cassidinae and Hispinae and therefore should be included as well if the study focuses on a broad variety of subfamilies. To further analyse leaf beetle diversity in the studied area, further sampling with the same methods could be performed in order to approach a saturation of the species accumulation curve.

For qualitative sampling, also hand-collection at selected spots can be recommended: Especially in sunny weather at dry days or after rainfall, various leaf beetles appear quite abundant at forest edges or gaps sun-basking or feeding on leaves where they can easily be collected.

A further implementation of Malaise traps can be highly recommended, however only for qualitative sampling (e.g. inventories) or when study sites at a certain distance are used (e.g. to compare the communities between the different elevational levels Bombuscaro, ECSF, and Cajanuma). In this case they seem to be very well suited for flying leaf beetle species. They can be used to obtain a high number of specimens and species with comparatively low workload and time-effort. Sampling can be even more facilitated if the changing of collection bottles is automated. This method would also allow a fine-scale study of temporal turnover, e.g. by sampling nocturnal and diurnal species separately.

Finally, it would be interesting to include the canopy, probably the most diverse habitat, into the study. Canopy fogging provides an efficient sampling method yielding very large numbers of individuals and species per sample.

# General discussion and future prospects

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This study is exceptional in two aspects: It is the first detailed, site-specific study about the diversity of leaf beetles (Chrysomelidae) in Ecuador, a hyperdiverse taxon that is severely understudied in the Neotropical region. Furthermore, the implementation of DNA barcoding makes it an important contribution towards the integration of DNA based methods into exploring and understanding the diversity and ecology of tropical insect assemblages.

Therefore, the lessons learned from this study cover those two aspects: On the one hand, it provides first data and insights of the hitherto unstudied leaf beetle fauna of the study area that can serve as a starting point for future research. On the other hand, it demonstrates the value of DNA barcoding for hyperdiverse arthropod assemblages, showing that even complex ecological questions can be analysed relying on DNA barcode data alone. The study provides baseline-data that can be used for future research, e.g. monitoring effects of climate change or anthropogenic disturbance on leaf beetle diversity, as well as the necessary tools.

## **Perspectives for future research**

Although the study area has not been completely inventoried and only a certain habitat was sampled, a considerable number of morphospecies, respectively MOTUs was found. This has been expected as the study area within the Tropical Andes is known as a biodiversity hotspot for different taxa. Besides a high turnover of communities between the three elevational levels, also a microhabitat-differentiation between ridge and valley forests was observed.

The standardized plot-based sampling design allows a detailed research on leaf beetle community ecology. In future, intermediate plots situated on the slopes between ridge and valley plots could be included in the study to allow finer-grained analyses. Furthermore, the gradient should be complemented with intermediate elevational levels and also its range should be extended: Whereas the tree line is almost reached in the present study, lower elevations should be included as well. The integration of lowland forest would allow more complex comparisons between low and high altitude fauna and could help to understand general diversity patterns along elevational gradients and their underlying causes.

As this study is the first of its kind in Ecuador and locally very restricted, nothing is known about how unique the leaf beetle fauna is in comparison with other regions even close by. Elevational gradients nearby or in other parts of Ecuador

should be investigated to interpret the observed patterns in a broader context. Apart from diversity changes along elevational gradients, also the response of leaf beetle communities to anthropogenic disturbance could be studied.

Another direction in which further research could lead concerns the role of leaf beetles as a model group of tropical herbivorous insects. A detailed analysis of host-specificity and species ranges along altitudinal gradients is necessary to understand general mechanisms that maintain high biodiversity in tropical mountain forests. The sampling design allows studying correlations of leaf beetle diversity with their potential host-plants on the sampling plots.

Another possibility for future research on Chrysomelidae in Podocarpus National Park and RBSF would be the continuation and intensification of the sampling towards a more complete inventory. This would allow an easier comparison with the fauna of other Neotropical regions. For an inventory also the canopy should be included that is supposed to harbour a largely different and very diverse fauna. Therefore, the comparison between understorey and canopy leaf beetle communities would be interesting as well.

In addition, once the beetles are investigated taxonomically by experts, the efforts made in this study are likely to lead to the description of a number of new species.

### **Integration of DNA barcoding into biodiversity assessments**

Facing the rapid advance of biodiversity loss in tropical rainforests, an acceleration of biodiversity assessment is indispensable. The traditional taxonomic approach is extremely time-intensive and for most tropical arthropods not possible at all: There is a severe lack of taxonomic expertise (taxonomic impediment), and if it is available, it is quite expensive.

Usually studies on tropical arthropod diversity rely on morphospecies instead of valid taxonomic species. Although this is a more superficial morphological approach, a precise morphospecies sorting is in many taxa quite challenging still and requires taxonomic expertise, too. Therefore, especially for individual- and species rich samples even a 'mere' morphospecies approach can be time- and cost-intensive if a high level of precision is attempted. In addition, even if precisely sorted, morphospecies always contain a certain degree of subjectivity and are hardly comparable among studies.

During the last years, molecular methods such as DNA barcoding have been propagated to accelerate analysis and understanding of biodiversity, with the time-intensive taxonomic identification and description of species being postponed or even completely relinquished. In this study, it could be shown that indeed DNA barcodes can be used for analysing diversity and ecology of leaf beetle communities even without valid taxonomic species information.

The use of DNA barcoding brings along plenty advantages: Whereas even the morphospecies sorting requires a certain degree of taxonomic expertise, in contrast DNA barcode sequences can be easily generated in any molecular laboratory with standard equipment and without any taxonomic knowledge. Furthermore, the

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method is open to scrutiny and replicable by third persons. The barcode-based MOTUs can be re-identified and allow e.g. a comparison with leaf beetle fauna of other regions. DNA barcoding can help to delimit species by facilitating and enhancing morphospecies sorting and therefore might prevent underestimation of species richness. Moreover, the method allows interesting follow-up studies as e.g. the exploration of phylogenetic diversity along the elevational gradient.

In the present study, a voucher-based approach of Sanger-sequencing of individuals was applied. Compared to a metabarcoding approach, this is more time-intensive and probably more costly, too. Usually not all collected specimens of a large sample but only a selection can be barcoded. In contrast, recent advances in metabarcoding allow compilation of large data sets with comparatively small time effort. On the other hand, the voucher-based approach can provide abundance data and it facilitates the connection of the specimen with all information connected to the individual. Especially when working with an unknown fauna it is advisable that as much information as possible stays connected to the sampled specimens: Specimens should be archived in collections together with comprehensive sampling information, DNA sequences, and photos that can be accessed via collection databases. This facilitates further research on the samples. In the long term, it is also desirable that the collection is complemented with formal Linnean species names. Although tropical samples contain many unknown species, in those cases where it is possible the specimens should be taxonomically identified. The Linnean names allow the linkage with existing knowledge of species' biology and ecology and are relevant for conservation issues as the compilation of red lists. The beetles from this study will be sent to taxonomic specialists for the respective subtaxon to be identified as far as possible and to formally describe new species. Whereas DNA barcoding certainly should be integrated into biodiversity assessments, it should create a feedback-loop with taxonomy and not replace it.

An essential factor influencing the efficacy of biodiversity assessment, regardless of whether morphology or DNA barcoding is used, is the sampling. Whereas, indeed, laboratory protocols for DNA sequencing have become more and more efficient during the last decades, the sampling that provides the specimens can constitute a bottleneck in the workflow because it can be quite effortful, as shown in this study. Although quite an effort was made, leaf beetle diversity is so rich that it could not be assessed exhaustively with the used methods, a circumstance implicating problems with the analyses. In this study, especially the plot-based sampling design is quite time-consuming as some plots are difficult to access and several replicates temporally scattered over the whole sampling period were sampled. However, it is exactly this sampling design that allows many possibilities of community analyses on a small-grained level. Sampling methods as Malaise trapping or canopy fogging could accelerate the sampling.

### **Résumé**

In the present study it could be shown that important knowledge about a very diverse unknown leaf beetle fauna can be derived using DNA barcodes instead of

taxonomic identifications. Beside information about species richness also more complex ecological issues such as species turnover and microhabitat specialization could be addressed. DNA barcoding allows an identification of species-like units without taxonomic expertise that is required even for a reliable morphospecies sorting when dealing with species- and individual rich assemblages. However, in the long term a taxonomic identification and the description of new species should be attempted. The chosen approach allows a variety of insights and offers many possibilities for subsequent research in both, the ecological and the methodological aspect.

This study shows once more how diverse and complex the insect fauna of tropical forests is and how much there is still to discover. Especially for tropical mountain forests there are large knowledge gaps although they are probably more diverse and more vulnerable than lowland forests. Although a considerable proportion of the country is protected area, Ecuador suffers the highest annual deforestation rate in South-America (-1.9%) and the Andean mountain forests are converted rapidly. To preserve Ecuador's outstanding and precious biodiversity, protection efforts should be intensified especially turning attention to the highly endangered mountain ecosystems.



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# List of abbreviations

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Table A.1: List of abbreviations used in this thesis.

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ANOVA	analysis of variance
a.s.l.	above sea level
B	Bombuscaro
BOLD	Barcode of Life Data Systems
bp	base pair
C	Cajanuma
CBOL	Consortium for the Barcode of Life
COI	cytochrome <i>c</i> oxidase I
E, ECSF	Estación Científica San Francisco
Fig.	Figure
GMYC	Generalized mixed Yule-coalescent
iBOL	international Barcode of Life project
MANOVA	multivariate analysis of variance
ML	Maximum Likelihood
MOTU	molecular operational taxonomic unit
mtDNA	mitochondrial DNA
NJ-Tree	Neighbor-Joining-Tree
NMDS	non-metric multidimensional scaling
PCR	Polymerase Chain Reaction
Podocarpus NP	Podocarpus National Park
PTP	Poisson tree processes
RBSF	Reserva Biológica San Francisco
rRNA	ribosomal RNA
SD	standard deviation
SE	standard error
Tab.	Table
UTPL	Universidad Técnica Particular de Loja
ZFMK	Zoologisches Forschungsmuseum Alexander Koenig

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# Additional information: Sampling site information

Table B.1: Sampling site information.

Site	Elevation (a.s.l.)	Latitude (S)	Longitude (W)
C-U1	2891	4° 6' 31.194"	79° 10' 44.184"
C-U2	2885	4° 6' 30.3804"	79° 10' 44.022"
C-U3	2869	4° 6' 30.7692"	79° 10' 42.9852"
C-U4	2886	4° 6' 30.5568"	79° 10' 49.6632"
C-U5	2890	4° 6' 30.618"	79° 10' 48.8208"
C-U6	2893	4° 6' 29.4804"	79° 10' 48.6948"
C-L1	2818	4° 6' 31.212"	79° 10' 37.3116"
C-L2	2805	4° 6' 31.896"	79° 10' 37.0488"
C-L3	2798	4° 6' 30.3048"	79° 10' 39.2268"
C-L4	2865	4° 6' 32.5404"	79° 10' 48.6552"
C-L5	2878	4° 6' 30.9492"	79° 10' 50.4084"
C-L6	2880	4° 6' 31.698"	79° 10' 50.3112"
B-U1	1075	4° 6' 49.8528"	78° 58' 1.0128"
B-U2	1066	4° 6' 50.8608"	78° 58' 0.7824"
B-U3	1072	4° 6' 50.2092"	78° 57' 59.94"
B-U4	1268	4° 7' 15.7008"	78° 58' 40.5588"
B-U5	1257	4° 7' 15.8592"	78° 58' 40.008"
B-U6	1266	4° 7' 16.8672"	78° 58' 39.648"
B-L1	1020	4° 6' 59.238"	78° 58' 5.2356"
B-L2	1026	4° 6' 59.5008"	78° 58' 5.916"
B-L3	1046	4° 6' 58.5648"	78° 58' 8.7384"
B-L4	1054	4° 7' 17.7888"	78° 58' 29.856"
B-L5	1056	4° 7' 18.0516"	78° 58' 31.1844"
B-L6	1044	4° 7' 19.614"	78° 58' 30.8568"
E-U1	2002	3° 58' 27.7896"	79° 4' 30.378"
E-U2	2026	3° 58' 29.8704"	79° 4' 30.2124"
E-U3	2089	3° 58' 37.9488"	79° 4' 32.1384"
E-U4	2063	3° 58' 32.1276"	79° 4' 20.0316"
E-U5	2054	3° 58' 31.0512"	79° 4' 20.4204"
E-U6	2039	3° 58' 30.108"	79° 4' 20.3916"
E-L1	2039	3° 58' 34.9536"	79° 4' 31.4328"
E-L2	1993	3° 58' 32.61"	79° 4' 32.5056"
E-L3	2030	3° 58' 34.63"	79° 4' 31.21"
E-L4	1913	3° 58' 26.13"	79° 4' 15.83"
E-L5	1954	3° 58' 27.3576"	79° 4' 12.8784"
E-L6	1933	3° 58' 28.7904"	79° 4' 12.972"
Cajanuma	Cajanuma area, unspecified		
Bombuscaro	Bombuscaro area, unspecified		
ECSF	ECSF area, unspecified		
E-Station	1826	3° 58' 17.19"	79° 4' 44.06"
E-Q2 (Quebrada 2)	1990	3° 58' 36"	79° 4' 32"
E-Q3 (Quebrada 3)	1990	3° 58' 27"	79° 4' 23"
E-Q5 (Quebrada 5)	1990	3° 58' 28"	79° 4' 13"
E-Lichtung	1900	3° 58' 24"	79° 4' 33"
El Tiro	2590	3° 59.55'	79° 07.30'



# Additional information: Specimen list

Table C.1: Specimen list with sampling information and GenBank accession numbers. All sampling sites are within Podocarpus National Park/Reserva Biológica San Francisco, Ecuador. Specimens 0227–0237, 0246, 0679–0737 were collected by G. Brehm, M. Adams, and L. Lehner, specimens 4572 and 4581 were collected by F. Bodner. All other specimens were sampled by B. Thormann, D. Sotomayor, J. Castillo, T. Klug, P. Schwalb, and J. Struwe. Coordinates of the sampling sites are provided in Tab. B.1. Hand-Coll.(N) = Non-standardized Hand-Collection, Hand-Coll.(S) = Standardized Hand-Collection, Flight-Intercept. = Flight Interception Trap.

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method	
BT_0001	Eumolpinae_sp_001	N1	KJ677921	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0002	Alticinae_sp_042	N1	KJ677411	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0003	Alticinae_sp_042	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0003a	Alticinae_sp_042	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0004	Eumolpinae_sp_042	N1	KJ677862	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0005	Galerucinae_sp_040	N1	KJ677774	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0006	Galerucinae_sp_040	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0006a	Galerucinae_sp_040	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0007	Galerucinae_sp_038	N1	KJ677526	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0008	Alticinae_sp_243	N1	KJ677417	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0009	Eumolpinae_sp_022	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0010	Eumolpinae_sp_022	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0011	Eumolpinae_sp_022	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0012	Eumolpinae_sp_021	N1	KJ677897	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0013	Eumolpinae_sp_021	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0014	Eumolpinae_sp_021	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0015	Galerucinae_sp_076	N1	KJ677559	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0016	Galerucinae_sp_076	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0017	Alticinae_sp_043	N1	KJ677407	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0018	Alticinae_sp_043	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0019	Alticinae_sp_043	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0020	Alticinae_sp_043	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0021	Alticinae_sp_007	N1	KJ677705	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0022	Alticinae_sp_219	N1	KJ677711	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0024	Galerucinae_sp_001	N1	KJ677550	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0025	Galerucinae_sp_001	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0026	Galerucinae_sp_001	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0027	Galerucinae_sp_001	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0028	Galerucinae_sp_001	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0029	Galerucinae_sp_001	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0030	Galerucinae_sp_001	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0031	Galerucinae_sp_001	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0032	Galerucinae_sp_001	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0033	Galerucinae_sp_037	N1	KJ677555	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0034	Eumolpinae_sp_014	N1	KJ677931	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0035	Eumolpinae_sp_006	N1	KJ677907	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0036	Galerucinae_sp_011	N1	KJ677532	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0043	Galerucinae_sp_005	N1	KJ677545	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0044	Galerucinae_sp_005	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0045	Galerucinae_sp_005	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0045a	Galerucinae_sp_005	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0046	Alticinae_sp_243	N1	KJ677415	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0047	Alticinae_sp_042	N1	KJ677412	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0048	Galerucinae_sp_039	N1	KJ677523	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0049	Galerucinae_sp_041	N1	KJ677775	ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0050	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0051	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0052	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0053	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0054	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0055	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0056	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0057	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0058	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0059	Galerucinae_sp_002	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0060	Galerucinae_sp_004	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0061	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0062	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0063	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0064	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0065	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0066	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0067	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0068	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0069	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0070	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)
BT_0071	Galerucinae_sp_007	N1		ECSF	Station	11/12 2010	Hand-Coll.(N)

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT_0072	Galerucinae sp_007	N1	ECSF	Station	11/12 2010	Hand-Coll.(N)	
BT_0085	Eumolpinae sp_048	N2	Bombuscaro	L1	20.11.2010	Hand-Coll.(N)	
BT_0088	Galerucinae sp_007	N4	ECSF	Station	20.11.2010	Light Trap	
BT_0089	Eumolpinae sp_001	N4	KJ677922	ECSF	Station	20.11.2010	Light Trap
BT_0090	Galerucinae sp_076	N5	KJ677558	ECSF	Station	21.11.2010	Light Trap
BT_0091	Eumolpinae sp_001	N5	KJ677923	ECSF	Station	21.11.2010	Light Trap
BT_0092	Eumolpinae sp_001	N5	ECSF	Station	21.11.2010	Light Trap	
BT_0093	Eumolpinae sp_001	N5	ECSF	Station	21.11.2010	Light Trap	
BT_0094	Galerucinae sp_011	N6	KJ677533	ECSF	U1	22.11.2010	Hand-Coll.(N)
BT_0095	Cassidinae sp_001	N7	KJ677873	ECSF		22.11.2010	Hand-Coll.(N)
BT_0096	Alticinae sp_010	N7	KJ677729	ECSF		22.11.2010	Hand-Coll.(N)
BT_0097	Alticinae sp_010	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0098	Galerucinae sp_002	N7	KJ677547	ECSF		22.11.2010	Hand-Coll.(N)
BT_0099	Galerucinae sp_007	N7	KJ677512	ECSF		22.11.2010	Hand-Coll.(N)
BT_0100	Galerucinae sp_007	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0101	Galerucinae sp_007	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0102	Alticinae sp_044	N7	KJ677374	ECSF		22.11.2010	Hand-Coll.(N)
BT_0103	Eumolpinae sp_038	N7	KJ677927	ECSF		22.11.2010	Hand-Coll.(N)
BT_0104	Eumolpinae sp_038	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0105	Eumolpinae sp_038	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0106	Eumolpinae sp_038	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0107	Galerucinae sp_046	N7	KJ677632	ECSF		22.11.2010	Hand-Coll.(N)
BT_0108	Galerucinae sp_046	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0109	Alticinae sp_251	N7	KJ677459	ECSF		22.11.2010	Hand-Coll.(N)
BT_0110	Alticinae sp_087	N7	KJ677497	ECSF		22.11.2010	Hand-Coll.(N)
BT_0111	Alticinae sp_087	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0112	Alticinae sp_087	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0113	Alticinae sp_087	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0114	Galerucinae sp_062	N7	KJ677756	ECSF		22.11.2010	Hand-Coll.(N)
BT_0115	Alticinae sp_098	N7	KJ677286	ECSF		22.11.2010	Hand-Coll.(N)
BT_0116	Alticinae sp_098	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0117	Eumolpinae sp_036	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0118	Eumolpinae sp_019	N7	KJ677877	ECSF		22.11.2010	Hand-Coll.(N)
BT_0119	Alticinae sp_124	N7	KJ677494	ECSF		22.11.2010	Hand-Coll.(N)
BT_0120	Alticinae sp_064	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0121	Alticinae sp_107	N7	KJ677776	ECSF		22.11.2010	Hand-Coll.(N)
BT_0122	Alticinae sp_107	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0123	Alticinae sp_129	N7	KJ677769	ECSF		22.11.2010	Hand-Coll.(N)
BT_0124	Alticinae sp_129	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0125	Alticinae sp_097	N7	KJ677311	ECSF		22.11.2010	Hand-Coll.(N)
BT_0126	Alticinae sp_123	N7	KJ677618	ECSF		22.11.2010	Hand-Coll.(N)
BT_0127	Hispaninae sp_001	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0128	Eumolpinae sp_009	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0129	Eumolpinae sp_009	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0130	Galerucinae sp_034	N7	KJ677691	ECSF		22.11.2010	Hand-Coll.(N)
BT_0131	Galerucinae sp_034	N7	ECSF			22.11.2010	Hand-Coll.(N)
BT_0133	Cassidinae sp_003	N8	ECSF	L6	24.11.2010	Hand-Coll.(N)	
BT_0134	Galerucinae sp_007	N8	KJ677513	ECSF	L6	24.11.2010	Hand-Coll.(N)
BT_0135	Eumolpinae sp_019	N9	KJ677878	ECSF		24.11.2010	Hand-Coll.(N)
BT_0136	Eumolpinae sp_019	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0137	Cassidinae sp_004	N9	KJ677850	ECSF		24.11.2010	Hand-Coll.(N)
BT_0138	Cassidinae sp_004	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0139	Alticinae sp_010	N9	KJ677730	ECSF		24.11.2010	Hand-Coll.(N)
BT_0140	Alticinae sp_028	N9	KJ677346	ECSF		24.11.2010	Hand-Coll.(N)
BT_0144	Eumolpinae sp_038	N9	KJ677926	ECSF		24.11.2010	Hand-Coll.(N)
BT_0145	Galerucinae sp_061	N9	KJ677514	ECSF		24.11.2010	Hand-Coll.(N)
BT_0146	Alticinae sp_029	N9	KJ677442	ECSF		24.11.2010	Hand-Coll.(N)
BT_0147	Alticinae sp_062	N9	KJ677421	ECSF		24.11.2010	Hand-Coll.(N)
BT_0148	Alticinae sp_066	N9	KJ677468	ECSF		24.11.2010	Hand-Coll.(N)
BT_0149	Alticinae sp_249	N9	KJ677456	ECSF		24.11.2010	Hand-Coll.(N)
BT_0150	Alticinae sp_249	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0151	Alticinae sp_249	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0152	Alticinae sp_249	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0153	Alticinae sp_109	N9	KJ677669	ECSF		24.11.2010	Hand-Coll.(N)
BT_0154	Alticinae sp_115	N9	KJ677287	ECSF		24.11.2010	Hand-Coll.(N)
BT_0155	Alticinae sp_193	N9	KJ677671	ECSF		24.11.2010	Hand-Coll.(N)
BT_0156	Alticinae sp_019	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0157	Alticinae sp_097	N9	KJ677300	ECSF		24.11.2010	Hand-Coll.(N)
BT_0158	Eumolpinae sp_002	N9	KJ677932	ECSF		24.11.2010	Hand-Coll.(N)
BT_0159	Galerucinae sp_096	N9	KJ677683	ECSF		24.11.2010	Hand-Coll.(N)
BT_0160	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0161	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0162	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0163	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0164	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0165	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0166	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0167	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0168	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0169	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0170	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0171	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0172	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0173	Galerucinae sp_096	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0174	Galerucinae sp_046	N9	KJ677633	ECSF		24.11.2010	Hand-Coll.(N)
BT_0175	Galerucinae sp_046	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0176	Galerucinae sp_046	N9	KJ677636	ECSF		24.11.2010	Hand-Coll.(N)
BT_0177	Galerucinae sp_046	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0178	Galerucinae sp_046	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0179	Galerucinae sp_046	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0180	Galerucinae sp_046	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0181	Galerucinae sp_046	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0182	Galerucinae sp_046	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0183	Galerucinae sp_034	N9	KJ677692	ECSF		24.11.2010	Hand-Coll.(N)
BT_0184	Galerucinae sp_034	N9	ECSF			24.11.2010	Hand-Coll.(N)
BT_0188	Galerucinae sp_011	N10	KJ677535	ECSF	Station	24.11.2010	Light Trap
BT_0189	Alticinae sp_161	N10	KJ677361	ECSF	Station	24.11.2010	Light Trap
BT_0190	Eumolpinae sp_001	N10	KJ677924	ECSF	Station	24.11.2010	Light Trap
BT_0191	Eumolpinae sp_001	N10	ECSF	Station		24.11.2010	Light Trap
BT_0195	Galerucinae sp_045	N11	ECSF			25.11.2010	Hand-Coll.(N)
BT_0196	Galerucinae sp_010	N11	KJ677807	Bombuscaro		25.11.2010	Hand-Coll.(N)
BT_0197	Galerucinae sp_033	N11	Bombuscaro			25.11.2010	Hand-Coll.(N)
BT_0198	Alticinae sp_137	N14	ECSF	U1	27.11.2010	Pitfall Trap	
BT_0199	Alticinae sp_118	N16	KJ677667	ECSF		27.11.2010	Hand-Coll.(N)
BT_0201	Chrysomelinae sp_002	N18	KJ677759	Bombuscaro	L3	29.11.2010	Hand-Coll.(N)
BT_0202	Galerucinae sp_032	N19	KJ677273	Bombuscaro		29.11.2010	Hand-Coll.(N)
BT_0203	Galerucinae sp_032	N19	Bombuscaro			29.11.2010	Hand-Coll.(N)
BT_0204	Hispaninae sp_002	N19	KJ677856	Bombuscaro		29.11.2010	Hand-Coll.(N)
BT_0206	Alticinae sp_085	N24	ECSF	U1	01.12.2010	Pitfall Trap	
BT_0207	Galerucinae sp_069	N25	KJ677778	ECSF		01.12.2010	Hand-Coll.(N)

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SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method	
BT 0208	Eumolpinae_sp_019	N26	KJ677879	ECSF	L2	02.12.2010	Hand-Coll.(N)
BT 0209	Cassidinae_sp_005	N27	KJ677822	ECSF		02.12.2010	Hand-Coll.(N)
BT 0210	Cassidinae_sp_006	N27		ECSF		02.12.2010	Hand-Coll.(N)
BT 0211	Alticinae_sp_087	N27	KJ677495	ECSF		02.12.2010	Hand-Coll.(N)
BT 0212	Galerucinae_sp_066	N27	KJ677794	ECSF		02.12.2010	Hand-Coll.(N)
BT 0213	Galerucinae_sp_024	N27	KJ677733	ECSF		02.12.2010	Hand-Coll.(N)
BT 0214	Alticinae_sp_028	N27	KJ677349	ECSF		02.12.2010	Hand-Coll.(N)
BT 0215	Alticinae_sp_028	N27		ECSF		02.12.2010	Hand-Coll.(N)
BT 0216	Eumolpinae_sp_038	N27		ECSF		02.12.2010	Hand-Coll.(N)
BT 0217	Eumolpinae_sp_038	N27		ECSF		02.12.2010	Hand-Coll.(N)
BT 0218	Galerucinae_sp_031	N27	KJ677751	ECSF		02.12.2010	Hand-Coll.(N)
BT 0219	Eumolpinae_sp_073	N27	KJ677831	ECSF		02.12.2010	Hand-Coll.(N)
BT 0220	Alticinae_sp_115	N27	KJ677288	ECSF		02.12.2010	Hand-Coll.(N)
BT 0221	Eumolpinae_sp_010	N27	KJ677906	ECSF		02.12.2010	Hand-Coll.(N)
BT 0223	Galerucinae_sp_034	N27	KJ677689	ECSF		02.12.2010	Hand-Coll.(N)
BT 0224	Galerucinae_sp_034	N27		ECSF		02.12.2010	Hand-Coll.(N)
BT 0227	Cassidinae_sp_014	N28	KJ677874	ECSF		02.12.2010	Hand-Coll.(N)
BT 0228	Cassidinae_sp_007	N28	KJ677837	ECSF		02.12.2010	Hand-Coll.(N)
BT 0229	Cassidinae_sp_007	N28		ECSF		02.12.2010	Hand-Coll.(N)
BT 0230	Eumolpinae_sp_022	N28		ECSF		02.12.2010	Hand-Coll.(N)
BT 0231	Hispinae_sp_003	N28	KJ677842	ECSF		02.12.2010	Hand-Coll.(N)
BT 0232	Hispinae_sp_004	N28	KJ677272	ECSF		02.12.2010	Hand-Coll.(N)
BT 0233	Alticinae_sp_061	N28	KJ677282	ECSF		02.12.2010	Hand-Coll.(N)
BT 0234	Alticinae_sp_097	N28	KJ677308	ECSF		02.12.2010	Hand-Coll.(N)
BT 0235	Alticinae_sp_010	N28	KJ677728	ECSF		02.12.2010	Hand-Coll.(N)
BT 0236	Alticinae_sp_156	N28	KJ677727	ECSF		02.12.2010	Hand-Coll.(N)
BT 0237	Alticinae_sp_156	N28		ECSF		02.12.2010	Hand-Coll.(N)
BT 0239	Galerucinae_sp_002	N29	KJ677548	ECSF		03.12.2010	Hand-Coll.(N)
BT 0240	Eumolpinae_sp_021	N29	KJ677898	ECSF		03.12.2010	Hand-Coll.(N)
BT 0241	Eumolpinae_sp_038	N29		ECSF		03.12.2010	Hand-Coll.(N)
BT 0242	Eumolpinae_sp_038	N29		ECSF		03.12.2010	Hand-Coll.(N)
BT 0243	Alticinae_sp_118	N29	KJ677666	ECSF		03.12.2010	Hand-Coll.(N)
BT 0244	Alticinae_sp_097	N29	KJ677309	ECSF		03.12.2010	Hand-Coll.(N)
BT 0245	Galerucinae_sp_034	N29	KJ677693	ECSF		03.12.2010	Hand-Coll.(N)
BT 0246	Galerucinae_sp_030	N30	KJ677701	ECSF	Q3	03.12.2010	Hand-Coll.(N)
BT 0247	Criocerinae_sp_001	N31	KJ677813	ECSF		05.12.2010	Hand-Coll.(N)
BT 0249	Criocerinae_sp_001	N31	KJ677814	ECSF		05.12.2010	Hand-Coll.(N)
BT 0250	Alticinae_sp_115	N31		ECSF		05.12.2010	Hand-Coll.(N)
BT 0251	Alticinae_sp_115	N31		ECSF		05.12.2010	Hand-Coll.(N)
BT 0252	Galerucinae_sp_034	N31	KJ677694	ECSF		05.12.2010	Hand-Coll.(N)
BT 0253	Galerucinae_sp_034	N31		ECSF		05.12.2010	Hand-Coll.(N)
BT 0254	Eumolpinae_sp_023	N31	KJ677899	ECSF		05.12.2010	Hand-Coll.(N)
BT 0255	Eumolpinae_sp_023	N31		ECSF		05.12.2010	Hand-Coll.(N)
BT 0256	Alticinae_sp_051	N32	KJ677625	ECSF	Station	03.12.2010	Light Trap
BT 0257	Galerucinae_sp_001	N32	KJ677551	ECSF	Station	03.12.2010	Light Trap
BT 0258	Galerucinae_sp_005	N32	KJ677546	ECSF	Station	03.12.2010	Light Trap
BT 0259	Alticinae_sp_096	N36	KJ677469	ECSF		08.12.2010	Hand-Coll.(N)
BT 0260	Eumolpinae_sp_038	N36		ECSF		08.12.2010	Hand-Coll.(N)
BT 0261	Eumolpinae_sp_038	N36		ECSF		08.12.2010	Hand-Coll.(N)
BT 0266	Alticinae_sp_145	N37		ECSF	Lichtung		Flight-Intercept.
BT 0267	Alticinae_sp_029	N39	KJ677443	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0268	Alticinae_sp_158	N39	KJ677582	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0269	Alticinae_sp_086	N39	KJ677396	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0270	Alticinae_sp_086	N39		ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0271	Alticinae_sp_064	N39	KJ677447	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0272	Alticinae_sp_064	N39		ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0273	Alticinae_sp_141	N39	KJ677585	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0274	Alticinae_sp_141	N39		ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0275	Alticinae_sp_141	N39		ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0276	Alticinae_sp_122	N39	KJ677777	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0277	Alticinae_sp_122	N39		ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0278	Alticinae_sp_124	N39	KJ677491	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0279	Alticinae_sp_115	N39	KJ677289	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0280	Eumolpinae_sp_022	N39		ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0281	Eumolpinae_sp_038	N39		ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0282	Eumolpinae_sp_038	N39		ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0283	Eumolpinae_sp_020	N39	KJ677941	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0284	Alticinae_sp_087	N39	KJ677496	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0285	Galerucinae_sp_022	N39	KJ677556	ECSF	Lichtung	08.12.2010	Malaise Trap
BT 0286	Alticinae_sp_012	N41		ECSF			Hand-Coll.(N)
BT 0287	Eumolpinae_sp_038	N41		ECSF			Hand-Coll.(N)
BT 0288	Eumolpinae_sp_017	N42	KJ677909	ECSF	Station	12.12.2010	Light Trap
BT 0289	Eumolpinae_sp_001	N42	KJ677925	ECSF	Station	12.12.2010	Light Trap
BT 0290	Eumolpinae_sp_001	N42		ECSF	Station	12.12.2010	Light Trap
BT 0291	Eumolpinae_sp_001	N42		ECSF	Station	12.12.2010	Light Trap
BT 0292	Galerucinae_sp_076	N42	KJ677557	ECSF	Station	12.12.2010	Light Trap
BT 0293	Galerucinae_sp_076	N42		ECSF	Station	12.12.2010	Light Trap
BT 0294	Galerucinae_sp_076	N42		ECSF	Station	12.12.2010	Light Trap
BT 0295	Galerucinae_sp_031	N43	KJ677752	ECSF		12.12.2010	Hand-Coll.(N)
BT 0296	Alticinae_sp_005	N44	KJ677652	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0297	Alticinae_sp_142	N44	KJ677593	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0298	Alticinae_sp_063	N44	KJ677342	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0299	Alticinae_sp_064	N44		ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0300	Alticinae_sp_064	N44		ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0301	Alticinae_sp_083	N44	KJ677334	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0302	Alticinae_sp_086	N44	KJ677398	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0303	Alticinae_sp_086	N44		ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0304	Alticinae_sp_086	N44		ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0305	Alticinae_sp_081	N44	KJ677765	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0306	Alticinae_sp_019	N44		ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0307	Alticinae_sp_115	N44	KJ677297	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0308	Alticinae_sp_018	N44		ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0309	Alticinae_sp_256	N44	KJ677301	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0310	Eumolpinae_sp_017	N44		ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0311	Eumolpinae_sp_042	N44	KJ677863	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0312	Eumolpinae_sp_021	N44	KJ677900	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0313	Alticinae_sp_122	N44		ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0314	Eumolpinae_sp_043	N44	KJ677809	ECSF	Lichtung	12.12.2010	Malaise Trap
BT 0318	Alticinae_sp_124	N50		ECSF	L6	14.12.2010	Malaise Trap
BT 0319	Galerucinae_sp_026	N50		ECSF	L6	14.12.2010	Malaise Trap
BT 0320	Galerucinae_sp_061	N50		ECSF	L6	14.12.2010	Malaise Trap
BT 0322	Alticinae_sp_143	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0323	Eumolpinae_sp_011	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0336	Galerucinae_sp_056	S1	KJ677740	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0337	Galerucinae_sp_056	S1	KJ677741	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0338	Galerucinae_sp_055	S1	KJ677742	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0339	Alticinae_sp_071	S1	KJ677330	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0340	Alticinae_sp_064	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0341	Alticinae_sp_073	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0342	Galerucinae_sp_051	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0343	Galerucinae_sp_052	S1	KJ677743	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0344	Galerucinae_sp_052	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0345	Alticinae_sp_051	S1		Bombuscaro	L1	20.11.2010	Sweep Netting

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SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT 0346	Alticinae sp 051	S1	KJ677366	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0347	Alticinae sp 051	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0348	Alticinae sp 051	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0349	Eumolpinae sp 020	S1	KJ677940	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0350	Eumolpinae sp 045	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0351	Eumolpinae sp 045	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0352	Alticinae sp 130	S1	KJ677771	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0353	Alticinae sp 132	S1	KJ677732	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0354	Alticinae sp 125	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0355	Alticinae sp 125	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0356	Alticinae sp 128	S1	KJ677509	Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0357	Alticinae sp 128	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0358	Alticinae sp 128	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0359	Alticinae sp 128	S1		Bombuscaro	L1	20.11.2010	Sweep Netting
BT 0361	Alticinae sp 050	S2	KJ677381	Bombuscaro	L1	20.11.2010	Beating
BT 0362	Alticinae sp 114	S2		Bombuscaro	L1	20.11.2010	Beating
BT 0363	Alticinae sp 051	S2	KJ677373	Bombuscaro	L1	20.11.2010	Beating
BT 0364	Eumolpinae sp 018	S2		Bombuscaro	L1	20.11.2010	Beating
BT 0365	Galerucinae sp 049	S2	KJ677645	Bombuscaro	L1	20.11.2010	Beating
BT 0366	Galerucinae sp 049	S2		Bombuscaro	L1	20.11.2010	Beating
BT 0367	Galerucinae sp 049	S2		Bombuscaro	L1	20.11.2010	Beating
BT 0368	Galerucinae sp 049	S2		Bombuscaro	L1	20.11.2010	Beating
BT 0369	Galerucinae sp 049	S2		Bombuscaro	L1	20.11.2010	Beating
BT 0370	Galerucinae sp 049	S2		Bombuscaro	L1	20.11.2010	Beating
BT 0371	Eumolpinae sp 006	S3		Bombuscaro	L1	20.11.2010	Hand-Coll.(S)
BT 0372	Alticinae sp 009	S3	KJ677723	Bombuscaro	L1	20.11.2010	Hand-Coll.(S)
BT 0373	Alticinae sp 009	S3		Bombuscaro	L1	20.11.2010	Hand-Coll.(S)
BT 0374	Alticinae sp 024	S4	KR424908	Bombuscaro	U2	20.11.2010	Sweep Netting
BT 0375	Eumolpinae sp 032	S4	KJ677904	Bombuscaro	U2	20.11.2010	Sweep Netting
BT 0376	Alticinae sp 051	S4	KJ677367	Bombuscaro	U2	20.11.2010	Sweep Netting
BT 0377	Eumolpinae sp 024	S4	KJ677912	Bombuscaro	U2	20.11.2010	Sweep Netting
BT 0378	Eumolpinae sp 024	S4		Bombuscaro	U2	20.11.2010	Sweep Netting
BT 0380	Eumolpinae sp 024	S5	KJ677918	Bombuscaro	U2	20.11.2010	Beating
BT 0381	Galerucinae sp 089	S5		Bombuscaro	U2	20.11.2010	Beating
BT 0382	Galerucinae sp 013	S6	KJ677537	Bombuscaro	U2	20.11.2010	Hand-Coll.(S)
BT 0383	Alticinae sp 087	S7	KJ677501	ECSF	U1	22.11.2010	Sweep Netting
BT 0384	Eumolpinae sp 039	S7	KJ677883	ECSF	U1	22.11.2010	Sweep Netting
BT 0385	Alticinae sp 064	S7		ECSF	U1	22.11.2010	Sweep Netting
BT 0386	Alticinae sp 064	S7		ECSF	U1	22.11.2010	Sweep Netting
BT 0387	Alticinae sp 104	S7	KJ677324	ECSF	U1	22.11.2010	Sweep Netting
BT 0388	Alticinae sp 104	S7		ECSF	U1	22.11.2010	Sweep Netting
BT 0389	Alticinae sp 105	S7		ECSF	U1	22.11.2010	Sweep Netting
BT 0390	Alticinae sp 090	S7	KJ677453	ECSF	U1	22.11.2010	Sweep Netting
BT 0391	Alticinae sp 090	S7	KJ677454	ECSF	U1	22.11.2010	Sweep Netting
BT 0392	Eumolpinae sp 074	S7	KJ677832	ECSF	U1	22.11.2010	Sweep Netting
BT 0395	Eumolpinae sp 074	S7	KJ677833	ECSF	U1	22.11.2010	Sweep Netting
BT 0396	Cassidinae sp 004	S8	KJ677851	ECSF	U1	22.11.2010	Beating
BT 0397	Alticinae sp I31	S8	KJ677654	ECSF	U1	22.11.2010	Beating
BT 0398	Cassidinae sp 003	S9		ECSF	U1	22.11.2010	Hand-Coll.(S)
BT 0399	Galerucinae sp 028	S10	KJ677437	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0400	Galerucinae sp 028	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0401	Galerucinae sp 054	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0402	Galerucinae sp 044	S10	KJ677603	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0403	Galerucinae sp 044	S10	KJ677604	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0404	Galerucinae sp 044	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0405	Galerucinae sp 045	S10	KJ677630	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0406	Galerucinae sp 045	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0407	Galerucinae sp 049	S10	KJ677642	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0408	Eumolpinae sp 024	S10	KJ677913	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0409	Eumolpinae sp 024	S10	KJ677914	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0410	Alticinae sp 141	S10	KJ677772	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0411	Alticinae sp 058	S10	KJ677350	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0412	Alticinae sp 058	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0413	Alticinae sp 058	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0414	Alticinae sp 058	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0415	Alticinae sp 242	S10	KJ677611	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0416	Alticinae sp 242	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0417	Galerucinae sp 074	S10	KJ677799	Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0418	Eumolpinae sp 045	S10		Bombuscaro	U4	25.11.2010	Sweep Netting
BT 0419	Alticinae sp 141	S11		Bombuscaro	U4	25.11.2010	Beating
BT 0420	Alticinae sp 128	S11	KJ677506	Bombuscaro	U4	25.11.2010	Beating
BT 0421	Alticinae sp 073	S11		Bombuscaro	U4	25.11.2010	Beating
BT 0422	Galerucinae sp 028	S11		Bombuscaro	U4	25.11.2010	Beating
BT 0423	Eumolpinae sp 024	S11	KJ677915	Bombuscaro	U4	25.11.2010	Beating
BT 0425	Eumolpinae sp 007	S12	KJ677876	Bombuscaro	U4	25.11.2010	Hand-Coll.(S)
BT 0426	Alticinae sp 150	S13	KJ677715	ECSF	U2	27.11.2010	Sweep Netting
BT 0427	Alticinae sp 104	S13	KJ677317	ECSF	U2	27.11.2010	Sweep Netting
BT 0428	Alticinae sp 105	S13	KJ677315	ECSF	U2	27.11.2010	Sweep Netting
BT 0429	Eumolpinae sp 039	S13	KJ677884	ECSF	U2	27.11.2010	Sweep Netting
BT 0430	Eumolpinae sp 039	S13		ECSF	U2	27.11.2010	Sweep Netting
BT 0431	Eumolpinae sp 039	S13		ECSF	U2	27.11.2010	Sweep Netting
BT 0432	Eumolpinae sp 030	S13	KJ677902	ECSF	U2	27.11.2010	Sweep Netting
BT 0433	Galerucinae sp 069	S14	KJ677780	ECSF	U2	27.11.2010	Beating
BT 0434	Galerucinae sp 072	S14	KJ677805	ECSF	U2	27.11.2010	Beating
BT 0436	Alticinae sp 149	S16	KJ677708	Bombuscaro	L2	29.11.2010	Sweep Netting
BT 0437	Alticinae sp 149	S16		Bombuscaro	L2	29.11.2010	Sweep Netting
BT 0438	Galerucinae sp 015	S16	KJ677747	Bombuscaro	L2	29.11.2010	Sweep Netting
BT 0439	Galerucinae sp 015	S16		Bombuscaro	L2	29.11.2010	Sweep Netting
BT 0440	Alticinae sp 025	S16	KR424909	Bombuscaro	L2	29.11.2010	Sweep Netting
BT 0441	Galerucinae sp 082	S16	KJ677687	Bombuscaro	L2	29.11.2010	Sweep Netting
BT 0442	Galerucinae sp 049	S16	KJ677647	Bombuscaro	L2	29.11.2010	Sweep Netting
BT 0443	Alticinae sp 051	S17	KJ677368	Bombuscaro	L2	29.11.2010	Beating
BT 0444	Alticinae sp 143	S17	KJ677574	Bombuscaro	L2	29.11.2010	Beating
BT 0445	Alticinae sp 153	S17		Bombuscaro	L2	29.11.2010	Beating
BT 0446	Alticinae sp 153	S17		Bombuscaro	L2	29.11.2010	Beating
BT 0447	Galerucinae sp 007	S17	KJ677541	Bombuscaro	L2	29.11.2010	Beating
BT 0448	Galerucinae sp 049	S17	KJ677648	Bombuscaro	L2	29.11.2010	Beating
BT 0449	Galerucinae sp 049	S17	KJ677646	Bombuscaro	L2	29.11.2010	Beating
BT 0451	Galerucinae sp 015	S18	KJ677748	Bombuscaro	L2	29.11.2010	Hand-Coll.(S)
BT 0452	Alticinae sp 009	S18	KJ677722	Bombuscaro	L2	29.11.2010	Hand-Coll.(S)
BT 0453	Alticinae sp 135	S19		Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0454	Alticinae sp 026	S19	KJ677408	Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0455	Alticinae sp 026	S19		Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0456	Alticinae sp 026	S19		Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0457	Alticinae sp 181	S19	KJ677279	Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0458	Alticinae sp 181	S19		Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0459	Galerucinae sp 050	S19	KJ677432	Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0460	Galerucinae sp 045	S19	KJ677629	Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0461	Galerucinae sp 045	S19	KJ677628	Bombuscaro	L3	29.11.2010	Sweep Netting
BT 0462	Eumolpinae sp 042	S20	KJ677858	Bombuscaro	L3	29.11.2010	Beating
BT 0463	Galerucinae sp 070	S20	KJ677803	Bombuscaro	L3	29.11.2010	Beating
BT 0464	Eumolpinae sp 045	S20		Bombuscaro	L3	29.11.2010	Beating
BT 0465	Alticinae sp 050	S20	KJ677382	Bombuscaro	L3	29.11.2010	Beating
BT 0467	Eumolpinae sp 018	S20		Bombuscaro	L3	29.11.2010	Beating
BT 0468	Alticinae sp 072	S20	KJ677620	Bombuscaro	L3	29.11.2010	Beating

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT 0469	Galerucinae sp 045	S20	KJ677631	Bombuscaro	L3	29.11.2010	Beating
BT 0470	Galerucinae sp 045	S20		Bombuscaro	L3	29.11.2010	Beating
BT 0471	Galerucinae sp 045	S20		Bombuscaro	L3	29.11.2010	Beating
BT 0472	Galerucinae sp 045	S20		Bombuscaro	L3	29.11.2010	Beating
BT 0473	Galerucinae sp 069	S22	KJ677779	ECSF	U1	01.12.2010	Sweep Netting
BT 0474	Alticinae sp 080	S22	KJ677615	ECSF	U1	01.12.2010	Sweep Netting
BT 0475	Eumolpinae sp 039	S22	KJ677885	ECSF	U1	01.12.2010	Sweep Netting
BT 0476	Eumolpinae sp 039	S22		ECSF	U1	01.12.2010	Sweep Netting
BT 0477	Alticinae sp 104	S22		ECSF	U1	01.12.2010	Sweep Netting
BT 0478	Alticinae sp 104	S22		ECSF	U1	01.12.2010	Sweep Netting
BT 0480	Hispiniae sp 005	S22		ECSF	U1	01.12.2010	Sweep Netting
BT 0489	Galerucinae sp 067	S23	KJ677802	ECSF	U1	01.12.2010	Beating
BT 0490	Hispiniae sp 005	S23	KJ677839	ECSF	U1	01.12.2010	Beating
BT 0491	Alticinae sp 157	S25	KJ677597	ECSF	L2	02.12.2010	Sweep Netting
BT 0492	Galerucinae sp 061	S25	KJ677515	ECSF	L2	02.12.2010	Sweep Netting
BT 0493	Eumolpinae sp 037	S25		ECSF	L2	02.12.2010	Sweep Netting
BT 0494	Alticinae sp 250	S25	KJ677450	ECSF	L2	02.12.2010	Sweep Netting
BT 0495	Alticinae sp 148	S25		ECSF	L2	02.12.2010	Sweep Netting
BT 0495a	Alticinae sp 148	S25		ECSF	L2	02.12.2010	Sweep Netting
BT 0496	Alticinae sp 112	S25	KJ677782	ECSF	L2	02.12.2010	Sweep Netting
BT 0497	Alticinae sp 111	S25		ECSF	L2	02.12.2010	Sweep Netting
BT 0498	Alticinae sp 113	S25		ECSF	L2	02.12.2010	Sweep Netting
BT 0499	Alticinae sp 044	S26	KJ677375	ECSF	L2	02.12.2010	Sweep Netting
BT 0500	Alticinae sp 044	S26		ECSF	L2	02.12.2010	Beating
BT 0501	Galerucinae sp 066	S26	KJ677795	ECSF	L2	02.12.2010	Beating
BT 0502	Eumolpinae sp 042	S26	KJ677864	ECSF	L2	02.12.2010	Beating
BT 0503	Alticinae sp 111	S26	KJ677601	ECSF	L2	02.12.2010	Beating
BT 0504	Alticinae sp 111	S26		ECSF	L2	02.12.2010	Beating
BT 0505	Alticinae sp 096	S27	KJ677470	ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0506	Alticinae sp 092	S27	KJ677737	ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0507	Alticinae sp 092	S27		ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0508	Galerucinae sp 053	S27	KJ677757	ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0509	Cassidinae sp 003	S27		ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0510	Cassidinae sp 004	S27	KJ677852	ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0511	Cassidinae sp 008	S27	KJ677821	ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0512	Hispiniae sp 006	S27	KJ677847	ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0513	Hispiniae sp 006	S27		ECSF	L2	02.12.2010	Hand-Coll.(S)
BT 0514	Eumolpinae sp 074	S28	KJ677834	ECSF	U3	03.12.2010	Sweep Netting
BT 0515	Eumolpinae sp 074	S28		ECSF	U3	03.12.2010	Sweep Netting
BT 0516	Alticinae sp 126	S28	KJ677783	ECSF	U3	03.12.2010	Sweep Netting
BT 0517	Alticinae sp 104	S28	KJ677325	ECSF	U3	03.12.2010	Sweep Netting
BT 0518	Alticinae sp 085	S28	KJ677388	ECSF	U3	03.12.2010	Sweep Netting
BT 0519	Alticinae sp 068	S29	KJ677462	ECSF	U3	03.12.2010	Beating
BT 0520	Alticinae sp 159	S29	KJ677595	ECSF	U3	03.12.2010	Beating
BT 0524	Eumolpinae sp 039	S29	KJ677895	ECSF	U3	03.12.2010	Beating
BT 0525	Hispiniae sp 023	S29	KJ677844	ECSF	U3	03.12.2010	Sweep Netting
BT 0526	Eumolpinae sp 039	S30	KJ677886	ECSF	U3	03.12.2010	Hand-Coll.(S)
BT 0527	Galerucinae sp 036	S31	KJ677761	ECSF	L1	08.12.2010	Sweep Netting
BT 0528	Alticinae sp 150	S31	KJ677714	ECSF	L1	08.12.2010	Sweep Netting
BT 0529	Alticinae sp 113	S31	KJ677755	ECSF	L1	08.12.2010	Sweep Netting
BT 0530	Galerucinae sp 031	S32	KJ677750	ECSF	L1	08.12.2010	Beating
BT 0531	Galerucinae sp 062	S32	KJ677758	ECSF	L1	08.12.2010	Beating
BT 0532	Galerucinae sp 075	S32	KJ677800	ECSF	L1	08.12.2010	Beating
BT 0533	Alticinae sp 097	S32	KJ677302	ECSF	L1	08.12.2010	Beating
BT 0534	Alticinae sp 111	S32		ECSF	L1	08.12.2010	Beating
BT 0535	Alticinae sp 097	S33	KJ677310	ECSF	L1	08.12.2010	Hand-Coll.(S)
BT 0536	Cassidinae sp 003	S33		ECSF	L1	08.12.2010	Hand-Coll.(S)
BT 0537	Alticinae sp 142	S34	KJ677591	ECSF	L3	08.12.2010	Sweep Netting
BT 0538	Alticinae sp 238	S34	KJ677679	ECSF	L3	08.12.2010	Sweep Netting
BT 0538a	Alticinae sp 238	S34		ECSF	L3	08.12.2010	Sweep Netting
BT 0539	Alticinae sp 013	S35	KJ677676	ECSF	L3	08.12.2010	Beating
BT 0540	Galerucinae sp 064	S35	KJ677788	ECSF	L3	08.12.2010	Beating
BT 0541	Galerucinae sp 064	S35		ECSF	L3	08.12.2010	Beating
BT 0543	Cassidinae sp 009	S36		ECSF	L3	08.12.2010	Hand-Coll.(S)
BT 0544	Cassidinae sp 012	S36	KJ677824	ECSF	L3	08.12.2010	Hand-Coll.(S)
BT 0546	Alticinae sp 083	S37	KJ677335	ECSF	L6	09.12.2010	Sweep Netting
BT 0547	Alticinae sp 096	S37	KJ677471	ECSF	L6	09.12.2010	Sweep Netting
BT 0548	Alticinae sp 112	S37		ECSF	L6	09.12.2010	Sweep Netting
BT 0549	Alticinae sp 150	S37	KJ677716	ECSF	L6	09.12.2010	Sweep Netting
BT 0550	Alticinae sp 265	S37	KJ677429	ECSF	L6	09.12.2010	Sweep Netting
BT 0551	Galerucinae sp 064	S38	KJ677789	ECSF	L6	09.12.2010	Beating
BT 0552	Alticinae sp 117	S38	KJ677492	ECSF	L6	09.12.2010	Beating
BT 0553	Eumolpinae sp 042	S38	KJ677865	ECSF	L6	09.12.2010	Beating
BT 0554	Galerucinae sp 071	S38	KJ677808	ECSF	L6	09.12.2010	Beating
BT 0555	Alticinae sp 086	S40	KJ677403	ECSF	L5	09.12.2010	Sweep Netting
BT 0556	Alticinae sp 096	S40	KJ677472	ECSF	L5	09.12.2010	Sweep Netting
BT 0557	Alticinae sp 181	S40	KJ677784	ECSF	L5	09.12.2010	Sweep Netting
BT 0558	Alticinae sp 149	S41	KJ677718	ECSF	L5	09.12.2010	Beating
BT 0559	Eumolpinae sp 042	S41	KJ677866	ECSF	L5	09.12.2010	Beating
BT 0560	Galerucinae sp 064	S41	KJ677790	ECSF	L5	09.12.2010	Beating
BT 0561	Galerucinae sp 066	S41	KJ677797	ECSF	L5	09.12.2010	Beating
BT 0562	Cassidinae sp 009	S42		ECSF	L5	09.12.2010	Hand-Coll.(S)
BT 0563	Cassidinae sp 009	S42		ECSF	L5	09.12.2010	Hand-Coll.(S)
BT 0564	Cassidinae sp 009	S42		ECSF	L5	09.12.2010	Hand-Coll.(S)
BT 0565	Alticinae sp 150	S42	KJ677712	ECSF	L5	09.12.2010	Hand-Coll.(S)
BT 0566	Hispiniae sp 006	S42	KJ677848	ECSF	L5	09.12.2010	Hand-Coll.(S)
BT 0567	Alticinae sp 104	S43	KJ677326	ECSF	U5	11.12.2010	Sweep Netting
BT 0568	Alticinae sp 118	S43		ECSF	U5	11.12.2010	Sweep Netting
BT 0569	Alticinae sp 118	S43		ECSF	U5	11.12.2010	Sweep Netting
BT 0570	Alticinae sp 118	S43		ECSF	U5	11.12.2010	Sweep Netting
BT 0571	Alticinae sp 118	S43		ECSF	U5	11.12.2010	Sweep Netting
BT 0572	Alticinae sp 118	S43		ECSF	U5	11.12.2010	Sweep Netting
BT 0573	Alticinae sp 118	S43		ECSF	U5	11.12.2010	Sweep Netting
BT 0574	Eumolpinae sp 039	S44	KJ677887	ECSF	U5	11.12.2010	Beating
BT 0575	Alticinae sp 140	S45	KJ677570	ECSF	U5	11.12.2010	Hand-Coll.(S)
BT 0576	Alticinae sp 118	S45		ECSF	U5	11.12.2010	Hand-Coll.(S)
BT 0577	Eumolpinae sp 039	S46	KJ677888	ECSF	U4	11.12.2010	Sweep Netting
BT 0578	Eumolpinae sp 039	S46		ECSF	U4	11.12.2010	Sweep Netting
BT 0579	Alticinae sp 104	S46	KJ677321	ECSF	U4	11.12.2010	Sweep Netting
BT 0580	Alticinae sp 104	S46		ECSF	U4	11.12.2010	Sweep Netting
BT 0581	Alticinae sp 118	S46		ECSF	U4	11.12.2010	Sweep Netting
BT 0582	Alticinae sp 118	S46		ECSF	U4	11.12.2010	Sweep Netting
BT 0583	Alticinae sp 118	S46		ECSF	U4	11.12.2010	Sweep Netting
BT 0584	Alticinae sp 118	S46		ECSF	U4	11.12.2010	Sweep Netting
BT 0585	Eumolpinae sp 044	S46		ECSF	U4	11.12.2010	Sweep Netting
BT 0586	Alticinae sp 091	S46		ECSF	U4	11.12.2010	Sweep Netting
BT 0587	Alticinae sp 006	S47	KJ677583	ECSF	U4	11.12.2010	Beating
BT 0588	Galerucinae sp 069	S47	KJ677781	ECSF	U4	11.12.2010	Beating
BT 0589	Eumolpinae sp 017	S47	KJ677910	ECSF	U4	11.12.2010	Beating
BT 0590	Alticinae sp 130	S47	KJ677571	ECSF	U4	11.12.2010	Beating
BT 0592	Eumolpinae sp 039	S47	KJ677889	ECSF	U4	11.12.2010	Beating
BT 0593	Alticinae sp 118	S47		ECSF	U4	11.12.2010	Beating
BT 0594	Alticinae sp 118	S48	KJ677663	ECSF	U4	11.12.2010	Hand-Coll.(S)
BT 0595	Alticinae sp 118	S48		ECSF	U4	11.12.2010	Hand-Coll.(S)

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method	
BT_0596	Eumolpinae_sp_039	S49	KJ677890	ECSF	U6	14.12.2010	Sweep Netting
BT_0597	Alticinae_sp_104	S49	KJ677322	ECSF	U6	14.12.2010	Sweep Netting
BT_0598	Alticinae_sp_118	S49		ECSF	U6	14.12.2010	Sweep Netting
BT_0599	Alticinae_sp_118	S49		ECSF	U6	14.12.2010	Sweep Netting
BT_0600	Alticinae_sp_118	S49		ECSF	U6	14.12.2010	Sweep Netting
BT_0601	Alticinae_sp_118	S49		ECSF	U6	14.12.2010	Sweep Netting
BT_0602	Alticinae_sp_118	S49		ECSF	U6	14.12.2010	Sweep Netting
BT_0603	Alticinae_sp_118	S49		ECSF	U6	14.12.2010	Sweep Netting
BT_0604	Alticinae_sp_005	S49	KJ677840	ECSF	U6	14.12.2010	Sweep Netting
BT_0605	Alticinae_sp_087	S50	KJ677498	ECSF	U6	14.12.2010	Beating
BT_0606	Eumolpinae_sp_039	S50	KJ677891	ECSF	U6	14.12.2010	Beating
BT_0607	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0608	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0609	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0610	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0611	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0612	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0613	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0614	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0615	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0616	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0617	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0618	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0619	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0620	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0621	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0622	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0623	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0624	Alticinae_sp_118	S50		ECSF	U6	14.12.2010	Beating
BT_0625	Cassidinae_sp_003	S51		ECSF	U6	14.12.2010	Hand-Coll.(S)
BT_0626	Alticinae_sp_118	S51		ECSF	U6	14.12.2010	Hand-Coll.(S)
BT_0627	Alticinae_sp_118	S51		ECSF	U6	14.12.2010	Hand-Coll.(S)
BT_0628	Cassidinae_sp_012	S52	KJ677825	ECSF	L4	14.12.2010	Sweep Netting
BT_0629	Cassidinae_sp_012	S52		ECSF	L4	14.12.2010	Sweep Netting
BT_0630	Alticinae_sp_036	S52	KJ677610	ECSF	L4	14.12.2010	Sweep Netting
BT_0631	Alticinae_sp_064	S52	KJ677451	ECSF	L4	14.12.2010	Sweep Netting
BT_0632	Alticinae_sp_118	S52	KJ677392	ECSF	L4	14.12.2010	Sweep Netting
BT_0633	Alticinae_sp_085	S52	KJ677404	ECSF	L4	14.12.2010	Sweep Netting
BT_0634	Alticinae_sp_066	S52	KJ677473	ECSF	L4	14.12.2010	Sweep Netting
BT_0635	Alticinae_sp_096	S52	KJ677474	ECSF	L4	14.12.2010	Sweep Netting
BT_0636	Alticinae_sp_096	S52		ECSF	L4	14.12.2010	Sweep Netting
BT_0637	Alticinae_sp_096	S52		ECSF	L4	14.12.2010	Sweep Netting
BT_0638	Alticinae_sp_084	S52		ECSF	L4	14.12.2010	Sweep Netting
BT_0639	Alticinae_sp_096	S53		ECSF	L4	14.12.2010	Beating
BT_0640	Alticinae_sp_018	S53	KJ677425	ECSF	L4	14.12.2010	Beating
BT_0641	Alticinae_sp_013	S53	KJ677674	ECSF	L4	14.12.2010	Beating
BT_0642	Eumolpinae_sp_042	S53	KJ677859	ECSF	L4	14.12.2010	Beating
BT_0643	Eumolpinae_sp_042	S53		ECSF	L4	14.12.2010	Beating
BT_0644	Galerucinae_sp_066	S53	KJ677796	ECSF	L4	14.12.2010	Beating
BT_0645	Galerucinae_sp_011	S54	KJ677552	ECSF	L4	14.12.2010	Hand-Coll.(S)
BT_0646	Alticinae_sp_071	S54	KJ677379	ECSF	L4	14.12.2010	Hand-Coll.(S)
BT_0647	Galerucinae_sp_035	S54	KJ677660	ECSF	L4	14.12.2010	Hand-Coll.(S)
BT_0648	Alticinae_sp_140	S55	KJ677568	ECSF	U1	15.12.2010	Sweep Netting
BT_0649	Alticinae_sp_140	S55		ECSF	U1	15.12.2010	Sweep Netting
BT_0650	Eumolpinae_sp_039	S55	KJ677892	ECSF	U1	15.12.2010	Sweep Netting
BT_0651	Eumolpinae_sp_039	S55		ECSF	U1	15.12.2010	Sweep Netting
BT_0652	Alticinae_sp_104	S55	KJ677318	ECSF	U1	15.12.2010	Sweep Netting
BT_0653	Alticinae_sp_104	S55		ECSF	U1	15.12.2010	Sweep Netting
BT_0654	Alticinae_sp_104	S55		ECSF	U1	15.12.2010	Sweep Netting
BT_0655	Alticinae_sp_257	S55	KJ677449	ECSF	U1	15.12.2010	Sweep Netting
BT_0656	Alticinae_sp_051	S56	KJ677351	ECSF	U1	15.12.2010	Sweep Netting
BT_0656a	Alticinae_sp_051	S56		ECSF	U1	15.12.2010	Beating
BT_0657	Eumolpinae_sp_039	S56	KJ677893	ECSF	U1	15.12.2010	Beating
BT_0658	Eumolpinae_sp_039	S56		ECSF	U1	15.12.2010	Beating
BT_0659	Eumolpinae_sp_039	S56		ECSF	U1	15.12.2010	Beating
BT_0660	Alticinae_sp_104	S56	KJ677327	ECSF	U1	15.12.2010	Beating
BT_0661	Alticinae_sp_104	S56		ECSF	U1	15.12.2010	Beating
BT_0662	Alticinae_sp_126	S56	KJ677313	ECSF	U1	15.12.2010	Beating
BT_0663	Alticinae_sp_150	S56	KJ677713	ECSF	U1	15.12.2010	Beating
BT_0664	Alticinae_sp_150	S56	KJ677720	ECSF	U1	15.12.2010	Beating
BT_0665	Alticinae_sp_150	S56	KJ677717	ECSF	U1	15.12.2010	Beating
BT_0666	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0667	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0668	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0669	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0670	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0671	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0672	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0673	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0674	Alticinae_sp_150	S56		ECSF	U1	15.12.2010	Beating
BT_0675	Alticinae_sp_008	S56	KJ677710	ECSF	U1	15.12.2010	Beating
BT_0676	Alticinae_sp_064	S57	KJ677457	ECSF	U1	15.12.2010	Beating
BT_0677	Eumolpinae_sp_039	S57	KJ677894	ECSF	U1	15.12.2010	Hand-Coll.(S)
BT_0678	Eumolpinae_sp_044	S57		ECSF	U1	15.12.2010	Hand-Coll.(S)
BT_0679	Eumolpinae_sp_001	N60		ECSF	Station	21.03.2011	Hand-Coll.(S)
BT_0680	Galerucinae_sp_076	N60	KJ677560	ECSF	Station	21.03.2011	Light Trap
BT_0681	Galerucinae_sp_076	N60		ECSF	Station	21.03.2011	Light Trap
BT_0682	Galerucinae_sp_076	N60		ECSF	Station	21.03.2011	Light Trap
BT_0683	Alticinae_sp_115	N60	KJ677298	ECSF	Station	21.03.2011	Light Trap
BT_0684	Alticinae_sp_115	N60		ECSF	Station	21.03.2011	Light Trap
BT_0685	Alticinae_sp_115	N60		ECSF	Station	21.03.2011	Light Trap
BT_0686	Eumolpinae_sp_023	N61	KJ677901	ECSF	Station	21.03.2011	Light Trap
BT_0687	Alticinae_sp_087	N61	KJ677499	ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0688	Criocerinae_sp_001	N61	KJ677815	ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0689	Criocerinae_sp_001	N61		ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0690	Alticinae_sp_124	N61	KJ677493	ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0691	Galerucinae_sp_031	N61	KJ677753	ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0692	Alticinae_sp_014	N61	KJ677588	ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0693	Alticinae_sp_014	N61		ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0694	Alticinae_sp_014	N61		ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0695	Alticinae_sp_014	N61		ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0696	Alticinae_sp_014	N61		ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0697	Alticinae_sp_014	N61		ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0698	Alticinae_sp_013	N62	KJ677675	ECSF	Station	22.03.2011	Hand-Coll.(N)
BT_0699	Alticinae_sp_083	N62	KJ677336	ECSF	Station	22.03.2011	Light Trap
BT_0700	Alticinae_sp_083	N62		ECSF	Station	22.03.2011	Light Trap
BT_0701	Alticinae_sp_083	N62		ECSF	Station	22.03.2011	Light Trap
BT_0705	Galerucinae_sp_008	N63	KJ677539	Bombuscaro	Station	23.03.2011	Light Trap
BT_0706	Eumolpinae_sp_038	N64		ECSF	Station	24.3.2011	Light Trap
BT_0708	Alticinae_sp_135	N64		ECSF	Station	24.3.2011	Light Trap
BT_0709	Galerucinae_sp_064	N65	KJ677791	ECSF	Station	24.03.2011	Light Trap
BT_0710	Alticinae_sp_013	N66	KJ677678	ECSF	Station	25.03.2011	Light Trap
BT_0711	Galerucinae_sp_073	N67	KJ677785	Cajanuma	Station	26.03.2011	Light Trap
BT_0712	Galerucinae_sp_073	N67		Cajanuma	Station	26.03.2011	Light Trap

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method	
BT 0713	Galerucinae sp 073	N68	KJ677786	Cajanuma		26.03.2011	Light Trap
BT 0714	Galerucinae sp 042	N70		ECSF	Station	28.03.2011	Hand-Coll.(N)
BT 0715	Galerucinae sp 076	N70	KJ677561	ECSF	Station	28.03.2011	Hand-Coll.(N)
BT 0716	Galerucinae sp 011	N70	KJ677534	ECSF	Station	28.03.2011	Hand-Coll.(N)
BT 0717	Galerucinae sp 002	N70	KJ677553	ECSF	Station	28.03.2011	Hand-Coll.(N)
BT 0718	Galerucinae sp 002	N70		ECSF	Station	28.03.2011	Hand-Coll.(N)
BT 0719	Eumolpinae sp 043	N72	KJ677810	ECSF		29.03.2011	Light Trap
BT 0720	Eumolpinae sp 043	N72		ECSF		29.03.2011	Light Trap
BT 0721	Eumolpinae sp 020	N72	KJ677945	ECSF		29.03.2011	Light Trap
BT 0722	Cassidinae sp 013	N73	KJ677811	Bombuscaro		30.03.2011	Light Trap
BT 0725	Alticinae sp 243	N75	KJ677416	ECSF		31.03.2011	Light Trap
BT 0726	Alticinae sp 243	N75		ECSF		31.03.2011	Light Trap
BT 0727	Galerucinae sp 073	N75	KJ677787	ECSF		31.03.2011	Light Trap
BT 0728	Galerucinae sp 064	N75	KJ677792	ECSF		31.03.2011	Light Trap
BT 0729	Galerucinae sp 015	N76	KJ677749	Bombuscaro		01.04.2011	Light Trap
BT 0730	Alticinae sp 136	N76	KJ677658	Bombuscaro		01.04.2011	Light Trap
BT 0731	Alticinae sp 004	N77		Bombuscaro		01.04.2011	Light Trap
BT 0732	Galerucinae sp 004	N77	KJ677538	ECSF		01.04.2011	Light Trap
BT 0733	Galerucinae sp 097	N77	KJ677540	Bombuscaro		01.04.2011	Light Trap
BT 0734	Galerucinae sp 039	N78	KJ677524	ECSF		02.04.2011	Light Trap
BT 0735	Galerucinae sp 038	N79	KJ677527	El Tiro	El Tiro	03.04.2011	Light Trap
BT 0736	Galerucinae sp 026	N80	KJ677768	ECSF		04.04.2011	Light Trap
BT 0737	Galerucinae sp 026	N80		ECSF		04.04.2011	Light Trap
BT 0738	Eumolpinae sp 004	N81	KJ677905	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0739	Galerucinae sp 018	N81	KJ677522	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0740	Galerucinae sp 076	N81	KJ677562	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0741	Galerucinae sp 002	N81	KJ677549	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0742	Galerucinae sp 019	N81	KJ677565	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0743	Galerucinae sp 014	N81	KJ677531	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0744	Galerucinae sp 007	N81	KJ677544	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0745	Galerucinae sp 007	N81		ECSF	Station	May 2011	Hand-Coll.(N)
BT 0746	Galerucinae sp 007	N81		ECSF	Station	May 2011	Hand-Coll.(N)
BT 0747	Galerucinae sp 038	N81	KJ677528	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0748	Galerucinae sp 038	N81		ECSF	Station	May 2011	Hand-Coll.(N)
BT 0749	Galerucinae sp 039	N81	KJ677525	ECSF	Station	May 2011	Hand-Coll.(N)
BT 0750	Cassidinae sp 005	N82	KJ677823	ECSF		08.05.2011	Hand-Coll.(N)
BT 0751	Galerucinae sp 062	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0752	Alticinae sp 039	N82	KJ677653	ECSF		08.05.2011	Hand-Coll.(N)
BT 0753	Alticinae sp 010	N82	KJ677731	ECSF		08.05.2011	Hand-Coll.(N)
BT 0754	Eumolpinae sp 008	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0755	Galerucinae sp 036	N82	KJ677760	ECSF		08.05.2011	Hand-Coll.(N)
BT 0756	Eumolpinae sp 022	N82	KJ677896	ECSF		08.05.2011	Hand-Coll.(N)
BT 0757	Eumolpinae sp 022	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0758	Eumolpinae sp 022	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0759	Eumolpinae sp 038	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0760	Eumolpinae sp 038	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0761	Eumolpinae sp 038	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0762	Eumolpinae sp 038	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0763	Eumolpinae sp 038	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0764	Eumolpinae sp 038	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0765	Eumolpinae sp 038	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0766	Alticinae sp 096	N82	KJ677475	ECSF		08.05.2011	Hand-Coll.(N)
BT 0767	Alticinae sp 096	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0768	Alticinae sp 096	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0770	Eumolpinae sp 016	N82	KJ677937	ECSF		08.05.2011	Hand-Coll.(N)
BT 0771	Eumolpinae sp 016	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0772	Eumolpinae sp 016	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0773	Eumolpinae sp 016	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0774	Eumolpinae sp 016	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0775	Eumolpinae sp 016	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0776	Alticinae sp 018	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0777	Alticinae sp 018	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0778	Alticinae sp 018	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0779	Alticinae sp 115	N82	KJ677290	ECSF		08.05.2011	Hand-Coll.(N)
BT 0780	Alticinae sp 115	N82		ECSF		08.05.2011	Hand-Coll.(N)
BT 0781	Eumolpinae sp 020	N82	KJ677942	ECSF		08.05.2011	Hand-Coll.(N)
BT 0782	Galerucinae sp 034	N82	KJ677695	ECSF		08.05.2011	Hand-Coll.(N)
BT 0788	Alticinae sp 006	N84	KJ677580	ECSF		10.05.2011	Hand-Coll.(N)
BT 0789	Alticinae sp 014	N84	KJ677589	ECSF		10.05.2011	Hand-Coll.(N)
BT 0790	Alticinae sp 014	N84		ECSF		10.05.2011	Hand-Coll.(N)
BT 0791	Eumolpinae sp 016	N84	KJ677935	ECSF		10.05.2011	Hand-Coll.(N)
BT 0792	Alticinae sp 115	N84	KJ677291	ECSF		10.05.2011	Hand-Coll.(N)
BT 0793	Alticinae sp 115	N84		ECSF		10.05.2011	Hand-Coll.(N)
BT 0794	Alticinae sp 129	N84	KJ677770	ECSF		10.05.2011	Hand-Coll.(N)
BT 0795	Alticinae sp 096	N84	KJ677463	ECSF		10.05.2011	Hand-Coll.(N)
BT 0796	Alticinae sp 097	N84	KJ677305	ECSF		10.05.2011	Hand-Coll.(N)
BT 0799	Cassidinae sp 014	N86	KJ677875	ECSF		12.05.2011	Hand-Coll.(N)
BT 0800	Galerucinae sp 030	N86	KJ677702	ECSF		12.05.2011	Hand-Coll.(N)
BT 0801	Cassidinae sp 006	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0802	Cassidinae sp 006	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0803	Galerucinae sp 076	N86	KJ677563	ECSF		12.05.2011	Hand-Coll.(N)
BT 0804	Alticinae sp 028	N86	KJ677347	ECSF		12.05.2011	Hand-Coll.(N)
BT 0805	Alticinae sp 014	N86	KJ677590	ECSF		12.05.2011	Hand-Coll.(N)
BT 0806	Alticinae sp 013	N86	KJ677677	ECSF		12.05.2011	Hand-Coll.(N)
BT 0807	Alticinae sp 054	N86	KJ677378	ECSF		12.05.2011	Hand-Coll.(N)
BT 0808	Alticinae sp 064	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0809	Alticinae sp 012	N86	KJ677622	ECSF		12.05.2011	Hand-Coll.(N)
BT 0810	Galerucinae sp 046	N86	KJ677637	ECSF		12.05.2011	Hand-Coll.(N)
BT 0811	Criocerinae sp 006	N86	KJ677817	ECSF		12.05.2011	Hand-Coll.(N)
BT 0812	Criocerinae sp 006	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0813	Criocerinae sp 004	N86	KJ677818	ECSF		12.05.2011	Hand-Coll.(N)
BT 0814	Alticinae sp 086	N86	KJ677405	ECSF		12.05.2011	Hand-Coll.(N)
BT 0815	Alticinae sp 032	N86	KJ677623	ECSF		12.05.2011	Hand-Coll.(N)
BT 0816	Galerucinae sp 019	N86	KJ677566	ECSF		12.05.2011	Hand-Coll.(N)
BT 0817	Alticinae sp 087	N86	KJ677592	ECSF		12.05.2011	Hand-Coll.(N)
BT 0818	Galerucinae sp 061	N86	KJ677516	ECSF		12.05.2011	Hand-Coll.(N)
BT 0819	Eumolpinae sp 074	N86	KJ677836	ECSF		12.05.2011	Hand-Coll.(N)
BT 0820	Eumolpinae sp 038	N86	KJ677928	ECSF		12.05.2011	Hand-Coll.(N)
BT 0821	Eumolpinae sp 038	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0822	Galerucinae sp 033	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0823	Galerucinae sp 034	N86	KJ677690	ECSF		12.05.2011	Hand-Coll.(N)
BT 0824	Galerucinae sp 034	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0825	Galerucinae sp 034	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0826	Galerucinae sp 034	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0827	Eumolpinae sp 020	N86	KJ677943	ECSF		12.05.2011	Hand-Coll.(N)
BT 0828	Alticinae sp 118	N86	KJ677664	ECSF		12.05.2011	Hand-Coll.(N)
BT 0829	Alticinae sp 115	N86	KJ677292	ECSF		12.05.2011	Hand-Coll.(N)
BT 0830	Alticinae sp 115	N86		ECSF		12.05.2011	Hand-Coll.(N)
BT 0831	Alticinae sp 147	N86	KJ677439	ECSF		12.05.2011	Hand-Coll.(N)
BT 0832	Alticinae sp 147	N86		ECSF		12.05.2011	Hand-Coll.(N)

Continued on next page(s)

Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT 0835	Galerucinae sp 029	N87	KJ677519	Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0836	Alticinae sp 003	N87	KJ677703	Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0837	Alticinae sp 009	N87	KJ677724	Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0838	Alticinae sp 059	N87		Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0839	Alticinae sp 040	N87	KJ677420	Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0840	Alticinae sp 140	N87	KJ677575	Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0841	Alticinae sp 127	N87	KJ677277	Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0842	Alticinae sp 127	N87		Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0843	Alticinae sp 136	N87	KJ677659	Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0844	Alticinae sp 138	N87		Bombuscaro		14.05.2011	Hand-Coll.(N)
BT 0847	Eumolpinae sp 005	N89	KJ677933	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0848	Alticinae sp 045	N89	KJ677419	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0849	Alticinae sp 045	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0850	Alticinae sp 049	N89	KJ677607	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0851	Alticinae sp 065	N89	KJ677476	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0852	Alticinae sp 133	N89	KJ677586	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0853	Alticinae sp 133	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0854	Alticinae sp 133	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0855	Alticinae sp 018	N89	KJ677426	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0856	Alticinae sp 018	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0857	Alticinae sp 018	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0858	Alticinae sp 018	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0859	Alticinae sp 018	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0860	Alticinae sp 018	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0861	Eumolpinae sp 020	N89	KJ677944	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0862	Alticinae sp 115	N89	KJ677293	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0863	Alticinae sp 115	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0864	Alticinae sp 253	N89	KJ677306	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0865	Alticinae sp 096	N89	KJ677477	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0866	Alticinae sp 096	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0867	Alticinae sp 096	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0868	Alticinae sp 122	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0869	Alticinae sp 122	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0870	Alticinae sp 122	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0871	Alticinae sp 086	N89	KJ677391	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0872	Alticinae sp 086	N89		ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0873	Alticinae sp 085	N89	KJ677397	ECSF	Lichtung	15.05.2011	Malaise Trap
BT 0877	Cassidinae sp 005	N90		ECSF		15.05.2011	Hand-Coll.(N)
BT 0879	Eumolpinae sp 043	N90		ECSF		15.05.2011	Hand-Coll.(N)
BT 0880	Alticinae sp 043	N90		ECSF		15.05.2011	Hand-Coll.(N)
BT 0892	Galerucinae sp 035	N90		ECSF		15.05.2011	Hand-Coll.(N)
BT 0893	Alticinae sp 083	N90		ECSF		15.05.2011	Hand-Coll.(N)
BT 0895	Cassidinae sp 006	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0899	Chrysomelinae sp 001	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0900	Alticinae sp 033	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0902	Alticinae sp 062	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0903	Alticinae sp 087	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0904	Alticinae sp 046	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0905	Galerucinae sp 060	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0906	Galerucinae sp 046	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0907	Galerucinae sp 046	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0908	Criocerinae sp 005	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0909	Criocerinae sp 001	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0924	Cryptocephalinae sp 001	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0925	Eumolpinae sp 029	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0926	Lamprosomatinae sp 002	N91		ECSF		17.05.2011	Hand-Coll.(N)
BT 0930	Galerucinae sp 046	N92		ECSF		18.05.2011	Hand-Coll.(N)
BT 0932	Alticinae sp 015	N92		ECSF		18.05.2011	Hand-Coll.(N)
BT 0934	Alticinae sp 116	N92		ECSF		18.05.2011	Hand-Coll.(N)
BT 0935	Alticinae sp 082	N92		ECSF		18.05.2011	Hand-Coll.(N)
BT 0936	Eumolpinae sp 026	N92		ECSF		18.05.2011	Hand-Coll.(N)
BT 0937	Alticinae sp 101	N92		ECSF		18.05.2011	Hand-Coll.(N)
BT 0943	Galerucinae sp 033	N92		ECSF		18.05.2011	Hand-Coll.(N)
BT 0949	Cassidinae sp 002	N93	KJ677871	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0950	Cassidinae sp 010	N93	KJ677849	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0951	Cassidinae sp 011	N93	KJ677872	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0952	Alticinae sp 269	N93	KJ677331	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0953	Galerucinae sp 028	N93	KJ677434	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0954	Galerucinae sp 029	N93	KJ677520	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0955	Galerucinae sp 020	N93	KJ677564	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0956	Alticinae sp 011	N93		Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0957	Galerucinae sp 036	N93	KJ677764	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0958	Alticinae sp 154	N93	KJ677376	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0959	Galerucinae sp 052	N93	KJ677746	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0960	Galerucinae sp 047	N93	KJ677639	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0961	Alticinae sp 077	N93	KJ677489	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0962	Alticinae sp 076	N93	KJ677343	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0963	Eumolpinae sp 033	N93		Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0964	Alticinae sp 030	N93	KJ677339	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0965	Criocerinae sp 002	N93	KJ677812	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0966	Criocerinae sp 003	N93	KJ677274	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0967	Eumolpinae sp 046	N93	KJ677920	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0968	Alticinae sp 099	N93	KJ677440	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0969	Alticinae sp 099	N93		Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0970	Alticinae sp 023	N93		Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0971	Alticinae sp 194	N93	KJ677341	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0972	Alticinae sp 127	N93	KJ677276	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0973	Alticinae sp 097	N93	KJ677285	Bombuscaro		20.05.2011	Hand-Coll.(N)
BT 0979	Eumolpinae sp 042	N94		ECSF	Lichtung	23.05.2011	Malaise Trap
BT 0980	Alticinae sp 005	N94		ECSF	Lichtung	23.05.2011	Malaise Trap
BT 0981	Eumolpinae sp 006	N94		ECSF	Lichtung	23.05.2011	Malaise Trap
BT 0982	Alticinae sp 022	N94		ECSF	Lichtung	23.05.2011	Malaise Trap
BT 0983	Alticinae sp 052	N94		ECSF	Lichtung	23.05.2011	Malaise Trap
BT 1021	Galerucinae sp 027	N97		ECSF	L6	27.05.2011	Flight-Intercept.
BT 1025	Eumolpinae sp 038	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1026	Eumolpinae sp 038	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1027	Eumolpinae sp 038	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1028	Alticinae sp 052	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1029	Alticinae sp 052	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1030	Alticinae sp 240	N99	KJ677332	ECSF	L6	27.05.2011	Malaise Trap
BT 1031	Alticinae sp 002	N99	KJ677707	ECSF	L6	27.05.2011	Malaise Trap
BT 1032	Alticinae sp 002	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1033	Alticinae sp 149	N99	KJ677719	ECSF	L6	27.05.2011	Malaise Trap
BT 1034	Alticinae sp 048	N99	KJ677377	ECSF	L6	27.05.2011	Malaise Trap
BT 1035	Alticinae sp 150	N99	KJ677721	ECSF	L6	27.05.2011	Malaise Trap
BT 1036	Alticinae sp 062	N99	KJ677422	ECSF	L6	27.05.2011	Malaise Trap
BT 1037	Alticinae sp 062	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1038	Alticinae sp 065	N99	KJ677478	ECSF	L6	27.05.2011	Malaise Trap
BT 1039	Alticinae sp 065	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1040	Alticinae sp 065	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1041	Alticinae sp 065	N99		ECSF	L6	27.05.2011	Malaise Trap
BT 1042	Alticinae sp 065	N99		ECSF	L6	27.05.2011	Malaise Trap

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method
BT_1043_Alticinae_sp_066	N99	KJ677479	ECSF	L6	27.05.2011	Malaise Trap
BT_1044_Alticinae_sp_081	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1045_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1046_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1047_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1048_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1049_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1050_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1051_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1052_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1053_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1054_Alticinae_sp_086	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1055_Alticinae_sp_085	N99	KJ677393	ECSF	L6	27.05.2011	Malaise Trap
BT_1056_Alticinae_sp_085	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1057_Alticinae_sp_085	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1058_Alticinae_sp_085	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1059_Alticinae_sp_081	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1060_Alticinae_sp_085	N99	KJ677399	ECSF	L6	27.05.2011	Malaise Trap
BT_1061_Galerucinae_sp_096	N99	KJ677684	ECSF	L6	27.05.2011	Malaise Trap
BT_1062_Galerucinae_sp_096	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1064_Alticinae_sp_142	N99	KJ677592	ECSF	L6	27.05.2011	Malaise Trap
BT_1065_Alticinae_sp_020	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1066_Alticinae_sp_018	N99	KJ677681	ECSF	L6	27.05.2011	Malaise Trap
BT_1067_Alticinae_sp_018	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1068_Alticinae_sp_018	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1069_Alticinae_sp_123	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1070_Alticinae_sp_097	N99	KJ677307	ECSF	L6	27.05.2011	Malaise Trap
BT_1071_Alticinae_sp_096	N99	KJ677480	ECSF	L6	27.05.2011	Malaise Trap
BT_1072_Alticinae_sp_096	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1073_Alticinae_sp_096	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1074_Alticinae_sp_096	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1075_Alticinae_sp_096	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1076_Alticinae_sp_096	N99		ECSF	L6	27.05.2011	Malaise Trap
BT_1079_Alticinae_sp_104	S58	KJ677328	ECSF	U1	07.05.2011	Sweep Netting
BT_1080_Eumolpinae_sp_074	S58	KJ677835	ECSF	U1	07.05.2011	Sweep Netting
BT_1081_Alticinae_sp_027	S59	KJ677409	ECSF	U1	07.05.2011	Beating
BT_1082_Alticinae_sp_104	S59	KJ677319	ECSF	U1	07.05.2011	Beating
BT_1083_Alticinae_sp_096	S61	KJ677481	ECSF	L1	08.05.2011	Sweep Netting
BT_1084_Alticinae_sp_096	S61		ECSF	L1	08.05.2011	Sweep Netting
BT_1085_Alticinae_sp_061	S61	KJ677283	ECSF	L1	08.05.2011	Sweep Netting
BT_1086_Alticinae_sp_083	S61	KJ677337	ECSF	L1	08.05.2011	Sweep Netting
BT_1087_Alticinae_sp_109	S61	KJ677670	ECSF	L1	08.05.2011	Sweep Netting
BT_1088_Galerucinae_sp_031	S61	KJ677754	ECSF	L1	08.05.2011	Sweep Netting
BT_1089_Galerucinae_sp_031	S61		ECSF	L1	08.05.2011	Sweep Netting
BT_1090_Alticinae_sp_019	S61	KJ677423	ECSF	L1	08.05.2011	Sweep Netting
BT_1091_Cassidinae_sp_012	S62	KJ677826	ECSF	L1	08.05.2011	Beating
BT_1092_Cassidinae_sp_003	S62	KJ677854	ECSF	L1	08.05.2011	Beating
BT_1093_Hispinae_sp_007	S62	KJ677869	ECSF	L1	08.05.2011	Beating
BT_1094_Eumolpinae_sp_038	S62	KJ677929	ECSF	L1	08.05.2011	Beating
BT_1095_Alticinae_sp_096	S62	KJ677464	ECSF	L1	08.05.2011	Beating
BT_1096_Galerucinae_sp_036	S62	KJ677762	ECSF	L1	08.05.2011	Beating
BT_1098_Alticinae_sp_097	S63	KJ677303	ECSF	L1	08.05.2011	Hand-Coll.(S)
BT_1101_Alticinae_sp_104	S64	KR424910	ECSF	U2	09.05.2011	Sweep Netting
BT_1104_Alticinae_sp_140	S68	KJ677569	ECSF	U3	10.05.2011	Beating
BT_1105_Alticinae_sp_074	S68	KJ677680	ECSF	U3	10.05.2011	Beating
BT_1106_Galerucinae_sp_064	S68	KJ677734	ECSF	U3	10.05.2011	Beating
BT_1107_Alticinae_sp_099	S70	KJ677672	ECSF	L5	12.05.2011	Sweep Netting
BT_1108_Alticinae_sp_092	S70	KJ677738	ECSF	L5	12.05.2011	Sweep Netting
BT_1109_Alticinae_sp_041	S70	KJ677598	ECSF	L5	12.05.2011	Sweep Netting
BT_1110_Galerucinae_sp_046	S70	KJ677638	ECSF	L5	12.05.2011	Sweep Netting
BT_1111_Galerucinae_sp_034	S70		ECSF	L5	12.05.2011	Sweep Netting
BT_1112_Alticinae_sp_112	S70	KJ677613	ECSF	L5	12.05.2011	Sweep Netting
BT_1114_Alticinae_sp_041	S71	KJ677599	ECSF	L5	12.05.2011	Beating
BT_1116_Cassidinae_sp_006	S72		ECSF	L5	12.05.2011	Hand-Coll.(S)
BT_1117_Alticinae_sp_049	S72	KJ677609	ECSF	L5	12.05.2011	Hand-Coll.(S)
BT_1118_Alticinae_sp_123	S72	KJ677619	ECSF	L5	12.05.2011	Hand-Coll.(S)
BT_1119_Alticinae_sp_096	S72	KJ677482	ECSF	L5	12.05.2011	Hand-Coll.(S)
BT_1120_Alticinae_sp_181	S72		ECSF	L5	12.05.2011	Hand-Coll.(S)
BT_1121_Alticinae_sp_124	S72	KJ677490	ECSF	L5	12.05.2011	Hand-Coll.(S)
BT_1122_Alticinae_sp_265	S72	KJ677430	ECSF	L5	12.05.2011	Hand-Coll.(S)
BT_1124_Cassidinae_sp_007	S73	KJ677430	ECSF	L6	12.05.2011	Sweep Netting
BT_1125_Galerucinae_sp_059	S73	KJ677838	ECSF	L6	12.05.2011	Sweep Netting
BT_1126_Hispinae_sp_003	S73	KJ677843	ECSF	L6	12.05.2011	Sweep Netting
BT_1127_Alticinae_sp_066	S73	KJ677483	ECSF	L6	12.05.2011	Sweep Netting
BT_1128_Alticinae_sp_018	S73	KJ677280	ECSF	L6	12.05.2011	Sweep Netting
BT_1129_Alticinae_sp_123	S73	KJ677617	ECSF	L6	12.05.2011	Sweep Netting
BT_1130_Alticinae_sp_140	S73	KR424911	ECSF	L6	12.05.2011	Sweep Netting
BT_1131_Alticinae_sp_081	S73	KJ677767	ECSF	L6	12.05.2011	Sweep Netting
BT_1132_Alticinae_sp_086	S73	KJ677400	ECSF	L6	12.05.2011	Sweep Netting
BT_1133_Alticinae_sp_086	S73		ECSF	L6	12.05.2011	Sweep Netting
BT_1134_Alticinae_sp_086	S73		ECSF	L6	12.05.2011	Sweep Netting
BT_1135_Alticinae_sp_086	S73		ECSF	L6	12.05.2011	Sweep Netting
BT_1136_Alticinae_sp_086	S73		ECSF	L6	12.05.2011	Sweep Netting
BT_1137_Alticinae_sp_086	S73		ECSF	L6	12.05.2011	Sweep Netting
BT_1138_Alticinae_sp_086	S73		ECSF	L6	12.05.2011	Sweep Netting
BT_1142_Alticinae_sp_096	S74	KJ677394	ECSF	L6	12.05.2011	Beating
BT_1143_Hispinae_sp_007	S74	KJ677870	ECSF	L6	12.05.2011	Beating
BT_1145_Cassidinae_sp_003	S75	KJ677855	ECSF	L6	12.05.2011	Hand-Coll.(S)
BT_1146_Eumolpinae_sp_014	S75	KJ677934	ECSF	L6	12.05.2011	Hand-Coll.(S)
BT_1147_Alticinae_sp_152	S75	KJ677355	ECSF	L6	12.05.2011	Hand-Coll.(S)
BT_1148_Alticinae_sp_096	S75	KJ677485	ECSF	L6	12.05.2011	Hand-Coll.(S)
BT_1151_Alticinae_sp_057	S76	KJ677445	Bombuscaro	U2	14.05.2011	Sweep Netting
BT_1152_Alticinae_sp_141	S79	KJ677773	Bombuscaro	L1	14.05.2011	Sweep Netting
BT_1154_Alticinae_sp_121	S80	KR424912	Bombuscaro	L1	14.05.2011	Beating
BT_1155_Alticinae_sp_121	S80		Bombuscaro	L1	14.05.2011	Beating
BT_1157_Alticinae_sp_149	S82	KJ677709	Bombuscaro	L2	14.05.2011	Sweep Netting
BT_1158_Alticinae_sp_146	S82	KJ677567	Bombuscaro	L2	14.05.2011	Sweep Netting
BT_1159_Alticinae_sp_146	S82		Bombuscaro	L2	14.05.2011	Sweep Netting
BT_1160_Alticinae_sp_055	S82	KJ677602	Bombuscaro	L2	14.05.2011	Sweep Netting
BT_1161_Alticinae_sp_143	S82	KJ677578	Bombuscaro	L2	14.05.2011	Sweep Netting
BT_1164_Alticinae_sp_087	S85		ECSF	U4	17.05.2011	Sweep Netting
BT_1165_Alticinae_sp_061	S85	KJ677284	ECSF	U4	17.05.2011	Sweep Netting
BT_1166_Alticinae_sp_140	S85	KJ677572	ECSF	U4	17.05.2011	Sweep Netting
BT_1167_Alticinae_sp_140	S85		ECSF	U4	17.05.2011	Sweep Netting
BT_1168_Alticinae_sp_140	S85		ECSF	U4	17.05.2011	Sweep Netting
BT_1169_Alticinae_sp_100	S85	KR424913	ECSF	U4	17.05.2011	Sweep Netting
BT_1170_Alticinae_sp_105	S85	KJ677316	ECSF	U4	17.05.2011	Sweep Netting
BT_1171_Alticinae_sp_085	S85	KJ677386	ECSF	U4	17.05.2011	Sweep Netting
BT_1175_Alticinae_sp_087	S86	KJ677503	ECSF	U4	17.05.2011	Beating
BT_1176_Alticinae_sp_052	S86	KJ677353	ECSF	U4	17.05.2011	Beating
BT_1178_Alticinae_sp_087	S87	KJ677504	ECSF	U4	17.05.2011	Hand-Coll.(S)
BT_1179_Alticinae_sp_118	S88	KJ677665	ECSF	U5	17.05.2011	Sweep Netting
BT_1180_Alticinae_sp_118	S88		ECSF	U5	17.05.2011	Sweep Netting
BT_1181_Alticinae_sp_118	S88		ECSF	U5	17.05.2011	Sweep Netting
BT_1182_Alticinae_sp_118	S88		ECSF	U5	17.05.2011	Sweep Netting

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SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT 1183	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1184	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1185	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1186	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1187	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1188	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1189	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1190	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1191	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1192	Alticinae sp 118	S88	ECSF	U5	17.05.2011	Sweep Netting	
BT 1194	Alticinae sp 087	S89	KJ677500	ECSF	U5	17.05.2011	Beating
BT 1195	Alticinae sp 087	S89	ECSF	U5	17.05.2011	Beating	
BT 1196	Alticinae sp 108	S89	KJ677455	ECSF	U5	17.05.2011	Beating
BT 1197	Alticinae sp 104	S89	KJ677320	ECSF	U5	17.05.2011	Beating
BT 1198	Alticinae sp 131	S89	KJ677655	ECSF	U5	17.05.2011	Beating
BT 1199	Alticinae sp 118	S90	KJ677661	ECSF	U5	17.05.2011	Hand-Coll.(S)
BT 1200	Alticinae sp 118	S90	ECSF	U5	17.05.2011	Hand-Coll.(S)	
BT 1201	Alticinae sp 118	S90	ECSF	U5	17.05.2011	Hand-Coll.(S)	
BT 1202	Alticinae sp 118	S90	ECSF	U5	17.05.2011	Hand-Coll.(S)	
BT 1203	Alticinae sp 118	S90	ECSF	U5	17.05.2011	Hand-Coll.(S)	
BT 1204	Alticinae sp 111	S90	ECSF	U5	17.05.2011	Hand-Coll.(S)	
BT 1205	Alticinae sp 052	S91	KJ677357	ECSF	L4	18.05.2011	Sweep Netting
BT 1206	Alticinae sp 052	S91	ECSF	L4	18.05.2011	Sweep Netting	
BT 1207	Alticinae sp 052	S91	ECSF	L4	18.05.2011	Sweep Netting	
BT 1208	Galerucinae sp 046	S91	KJ677634	ECSF	L4	18.05.2011	Sweep Netting
BT 1209	Galerucinae sp 046	S91	ECSF	L4	18.05.2011	Sweep Netting	
BT 1210	Alticinae sp 064	S91	KJ677452	ECSF	L4	18.05.2011	Sweep Netting
BT 1211	Alticinae sp 018	S91	KJ677427	ECSF	L4	18.05.2011	Sweep Netting
BT 1212	Alticinae sp 017	S91	KJ677380	ECSF	L4	18.05.2011	Sweep Netting
BT 1213	Alticinae sp 096	S91	KJ677486	ECSF	L4	18.05.2011	Sweep Netting
BT 1214	Alticinae sp 118	S91	KJ677649	ECSF	L4	18.05.2011	Sweep Netting
BT 1215	Alticinae sp 144	S91	KJ677584	ECSF	L4	18.05.2011	Sweep Netting
BT 1216	Alticinae sp 049	S91	KJ677606	ECSF	L4	18.05.2011	Sweep Netting
BT 1217	Alticinae sp 145	S91	KJ677579	ECSF	L4	18.05.2011	Sweep Netting
BT 1219	Alticinae sp 086	S91	KJ677406	ECSF	L4	18.05.2011	Sweep Netting
BT 1220	Galerucinae sp 034	S91	KJ677696	ECSF	L4	18.05.2011	Sweep Netting
BT 1221	Galerucinae sp 034	S91	ECSF	L4	18.05.2011	Sweep Netting	
BT 1222	Alticinae sp 052	S92	KJ677358	ECSF	L4	18.05.2011	Beating
BT 1223	Alticinae sp 028	S92	KJ677348	ECSF	L4	18.05.2011	Beating
BT 1224	Galerucinae sp 075	S92	KJ677801	ECSF	L4	18.05.2011	Beating
BT 1225	Eumolpinae sp 040	S92	KJ677867	ECSF	L4	18.05.2011	Beating
BT 1226	Galerucinae sp 034	S92	KJ677697	ECSF	L4	18.05.2011	Beating
BT 1227	Galerucinae sp 034	S92	ECSF	L4	18.05.2011	Beating	
BT 1228	Alticinae sp 096	S92	KJ677487	ECSF	L4	18.05.2011	Beating
BT 1229	Alticinae sp 096	S92	ECSF	L4	18.05.2011	Beating	
BT 1230	Alticinae sp 115	S92	KJ677295	ECSF	L4	18.05.2011	Beating
BT 1233	Alticinae sp 052	S93	KJ677356	ECSF	L4	18.05.2011	Hand-Coll.(S)
BT 1234	Alticinae sp 052	S93	ECSF	L4	18.05.2011	Hand-Coll.(S)	
BT 1235	Alticinae sp 096	S93	KJ677465	ECSF	L4	18.05.2011	Hand-Coll.(S)
BT 1236	Alticinae sp 096	S93	ECSF	L4	18.05.2011	Hand-Coll.(S)	
BT 1237	Galerucinae sp 034	S93	ECSF	L4	18.05.2011	Hand-Coll.(S)	
BT 1238	Galerucinae sp 034	S93	ECSF	L4	18.05.2011	Hand-Coll.(S)	
BT 1239	Galerucinae sp 034	S93	ECSF	L4	18.05.2011	Hand-Coll.(S)	
BT 1240	Eumolpinae sp 041	S93	KJ677857	ECSF	L4	18.05.2011	Hand-Coll.(S)
BT 1242	Lamprosomatinae sp 003	S93	ECSF	L4	18.05.2011	Hand-Coll.(S)	
BT 1244	Alticinae sp 118	S94	ECSF	U6	18.05.2011	Sweep Netting	
BT 1245	Alticinae sp 085	S94	KJ677389	ECSF	U6	18.05.2011	Sweep Netting
BT 1248	Alticinae sp 118	S95	ECSF	U6	18.05.2011	Beating	
BT 1249	Alticinae sp 104	S95	KJ677323	ECSF	U6	18.05.2011	Beating
BT 1250	Alticinae sp 104	S95	ECSF	U6	18.05.2011	Beating	
BT 1251	Alticinae sp 131	S95	KJ677656	ECSF	U6	18.05.2011	Beating
BT 1252	Alticinae sp 086	S95	KJ677401	ECSF	U6	18.05.2011	Beating
BT 1258	Galerucinae sp 028	S97	KJ677438	Bombuscaro	U4	20.05.2011	Sweep Netting
BT 1259	Galerucinae sp 049	S97	KJ677643	Bombuscaro	U4	20.05.2011	Sweep Netting
BT 1260	Eumolpinae sp 024	S97	KJ677916	Bombuscaro	U4	20.05.2011	Sweep Netting
BT 1261	Alticinae sp 103	S97	KR424914	Bombuscaro	U4	20.05.2011	Sweep Netting
BT 1262	Alticinae sp 111	S97	ECSF	U4	20.05.2011	Sweep Netting	
BT 1263	Alticinae sp 051	S97	KJ677369	Bombuscaro	U4	20.05.2011	Sweep Netting
BT 1264	Alticinae sp 078	S97	KJ677616	Bombuscaro	U4	20.05.2011	Sweep Netting
BT 1265	Alticinae sp 093	S97	ECSF	U4	20.05.2011	Sweep Netting	
BT 1266	Alticinae sp 093	S97	ECSF	U4	20.05.2011	Sweep Netting	
BT 1267	Alticinae sp 128	S98	KJ677510	Bombuscaro	U4	20.05.2011	Beating
BT 1269	Alticinae sp 009	S99	KJ677725	Bombuscaro	U4	20.05.2011	Hand-Coll.(S)
BT 1270	Galerucinae sp 028	S99	KJ677435	Bombuscaro	U4	20.05.2011	Hand-Coll.(S)
BT 1271	Alticinae sp 057	S99	KJ677444	Bombuscaro	U4	20.05.2011	Hand-Coll.(S)
BT 1272	Alticinae sp 128	S99	KJ677507	Bombuscaro	U4	20.05.2011	Hand-Coll.(S)
BT 1273	Galerucinae sp 028	S100	KJ677436	Bombuscaro	U6	20.05.2011	Sweep Netting
BT 1274	Alticinae sp 042	S100	KJ677410	Bombuscaro	U6	20.05.2011	Sweep Netting
BT 1275	Alticinae sp 051	S100	KJ677363	Bombuscaro	U6	20.05.2011	Sweep Netting
BT 1276	Alticinae sp 093	S100	ECSF	U6	20.05.2011	Sweep Netting	
BT 1277	Alticinae sp 093	S100	ECSF	U6	20.05.2011	Sweep Netting	
BT 1278	Alticinae sp 143	S100	ECSF	U6	20.05.2011	Sweep Netting	
BT 1279	Alticinae sp 125	S100	ECSF	U6	20.05.2011	Sweep Netting	
BT 1280	Alticinae sp 069	S100	KJ677461	Bombuscaro	U6	20.05.2011	Sweep Netting
BT 1281	Galerucinae sp 047	S100	KJ677640	Bombuscaro	U6	20.05.2011	Sweep Netting
BT 1282	Galerucinae sp 052	S100	KJ677744	Bombuscaro	U6	20.05.2011	Sweep Netting
BT 1283	Galerucinae sp 006	S101	KJ677536	Bombuscaro	U6	20.05.2011	Beating
BT 1284	Galerucinae sp 063	S101	KJ677529	Bombuscaro	U6	20.05.2011	Beating
BT 1285	Galerucinae sp 047	S101	KR425305	Bombuscaro	U6	20.05.2011	Beating
BT 1286	Alticinae sp 021	S101	KJ677441	Bombuscaro	U6	20.05.2011	Beating
BT 1287	Alticinae sp 067	S101	KJ677458	Bombuscaro	U6	20.05.2011	Beating
BT 1288	Alticinae sp 103	S101	KJ677364	Bombuscaro	U6	20.05.2011	Beating
BT 1289	Alticinae sp 102	S101	ECSF	U6	20.05.2011	Beating	
BT 1293	Alticinae sp 050	S101	KJ677383	Bombuscaro	U6	20.05.2011	Beating
BT 1294	Galerucinae sp 072	S101	KJ677804	Bombuscaro	U6	20.05.2011	Beating
BT 1295	Alticinae sp 057	S101	KJ677446	Bombuscaro	U6	20.05.2011	Beating
BT 1296	Alticinae sp 057	S101	ECSF	U6	20.05.2011	Beating	
BT 1297	Alticinae sp 057	S101	ECSF	U6	20.05.2011	Beating	
BT 1298	Alticinae sp 057	S101	ECSF	U6	20.05.2011	Beating	
BT 1299	Alticinae sp 057	S101	ECSF	U6	20.05.2011	Beating	
BT 1300	Alticinae sp 057	S101	ECSF	U6	20.05.2011	Beating	
BT 1301	Alticinae sp 057	S101	ECSF	U6	20.05.2011	Beating	
BT 1302	Alticinae sp 057	S101	ECSF	U6	20.05.2011	Beating	
BT 1303	Alticinae sp 057	S101	ECSF	U6	20.05.2011	Beating	
BT 1304	Galerucinae sp 047	S102	KJ677641	Bombuscaro	U6	20.05.2011	Hand-Coll.(S)
BT 1305	Alticinae sp 093	S102	KJ677359	Bombuscaro	U6	20.05.2011	Hand-Coll.(S)
BT 1306	Alticinae sp 125	S102	ECSF	U6	20.05.2011	Hand-Coll.(S)	
BT 1307	Eumolpinae sp 016	S102	KJ677938	Bombuscaro	U6	20.05.2011	Hand-Coll.(S)
BT 1308	Alticinae sp 134	S103	KJ677596	Bombuscaro	U5	20.05.2011	Sweep Netting
BT 1309	Alticinae sp 118	S103	ECSF	U5	20.05.2011	Sweep Netting	
BT 1310	Alticinae sp 035	S103	KJ677612	Bombuscaro	U5	20.05.2011	Sweep Netting
BT 1311	Alticinae sp 069	S103	KJ677460	Bombuscaro	U5	20.05.2011	Sweep Netting
BT 1312	Alticinae sp 093	S103	KJ677360	Bombuscaro	U5	20.05.2011	Sweep Netting
BT 1313	Alticinae sp 093	S103	ECSF	U5	20.05.2011	Sweep Netting	

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method
BT 1314	Alticinae sp_093	S103	Bombuscaro	U5	20.05.2011	Sweep Netting
BT 1315	Alticinae sp_093	S103	Bombuscaro	U5	20.05.2011	Sweep Netting
BT 1318	Eumolpinae sp_024	S104	KJ677917	Bombuscaro	U5	20.05.2011
BT 1319	Alticinae sp_051	S104	KJ677370	Bombuscaro	U5	20.05.2011
BT 1321	Galerucinae sp_061	S106	KJ677517	ECSF	L2	23.05.2011
BT 1322	Alticinae sp_019	S106	KJ677424	ECSF	L2	23.05.2011
BT 1323	Alticinae sp_104	S106	KJ677651	ECSF	L2	23.05.2011
BT 1324	Alticinae sp_092	S106	KJ677739	ECSF	L2	23.05.2011
BT 1325	Alticinae sp_019	S107		ECSF	L2	23.05.2011
BT 1326	Eumolpinae sp_019	S108	KJ677880	ECSF	L2	23.05.2011
BT 1327	Alticinae sp_097	S109	KR424915	ECSF	L1	23.05.2011
BT 1328	Alticinae sp_106	S109		ECSF	L1	23.05.2011
BT 1329	Alticinae sp_106	S109		ECSF	L1	23.05.2011
BT 1330	Alticinae sp_140	S110		ECSF	L1	23.05.2011
BT 1331	Cassidinae sp_004	S111	KR424916	ECSF	L1	23.05.2011
BT 1332	Cassidinae sp_004	S111	KR424783	ECSF	L1	23.05.2011
BT 1334	Alticinae sp_245	S112		Cajunuma	L6	24.05.2011
BT 1335	Alticinae sp_118	S112	KJ677662	Cajunuma	L6	24.05.2011
BT 1336	Alticinae sp_244	S112	KR424917	Cajunuma	L6	24.05.2011
BT 1337	Lamprosomatinae sp_001	S112		Cajunuma	L6	24.05.2011
BT 1339	Alticinae sp_244	S118	KR424918	Cajunuma	L4	24.05.2011
BT 1340	Galerucinae sp_066	S119	KJ677798	Cajunuma	L4	24.05.2011
BT 1347	Alticinae sp_092	S123		ECSF	L3	27.05.2011
BT 1349	Alticinae sp_051	S125	KJ677352	ECSF	U2	27.05.2011
BT 1350	Alticinae sp_131	S126	KJ677657	ECSF	U2	27.05.2011
BT 1756	Alticinae sp_014	N101		ECSF	01.06.2011	Hand-Coll.(S)
BT 1758	Alticinae sp_014	N101		ECSF	01.06.2011	Hand-Coll.(N)
BT 1773	Galerucinae sp_030	N102		ECSF	03.06.2011	Hand-Coll.(N)
BT 1775	Alticinae sp_033	N102		ECSF	03.06.2011	Hand-Coll.(N)
BT 1776	Galerucinae sp_062	N102		ECSF	03.06.2011	Hand-Coll.(N)
BT 1784	Lamprosomatinae sp_001	N102		ECSF	03.06.2011	Hand-Coll.(N)
BT 1785	Chrysomelinae sp_003	N102		ECSF	03.06.2011	Hand-Coll.(N)
BT 1789	Galerucinae sp_022	N104		ECSF	09.06.2011	Hand-Coll.(N)
BT 1794	Alticinae sp_001	N104		ECSF	09.06.2011	Hand-Coll.(N)
BT 1795	Chrysomelinae sp_001	N104		ECSF	09.06.2011	Hand-Coll.(N)
BT 1809	Alticinae sp_001	N105		ECSF	09.06.2011	Malaise Trap
BT 1810	Eumolpinae sp_012	N105		ECSF	09.06.2011	Malaise Trap
BT 1811	Galerucinae sp_054	N105		ECSF	09.06.2011	Malaise Trap
BT 1812	Alticinae sp_002	N105		ECSF	09.06.2011	Malaise Trap
BT 1813	Alticinae sp_052	N105		ECSF	09.06.2011	Malaise Trap
BT 1820	Alticinae sp_142	N105		ECSF	09.06.2011	Malaise Trap
BT 1821	Alticinae sp_088	N105		ECSF	09.06.2011	Malaise Trap
BT 1928	Galerucinae sp_019	N109		ECSF	16.06.2011	Hand-Coll.(N)
BT 1932	Eumolpinae sp_042	N110		ECSF	16.06.2011	Malaise Trap
BT 1934	Eumolpinae sp_014	N110		ECSF	16.06.2011	Malaise Trap
BT 1935	Alticinae sp_074	N110		ECSF	16.06.2011	Malaise Trap
BT 1936	Alticinae sp_049	N110		ECSF	16.06.2011	Malaise Trap
BT 1954	Galerucinae sp_041	N113		ECSF	28.06.2011	Hand-Coll.(N)
BT 1963	Alticinae sp_037	N115		ECSF	05.07.2011	Malaise Trap
BT 1964	Galerucinae sp_022	N115		ECSF	05.07.2011	Malaise Trap
BT 1970	Eumolpinae sp_007	N115		ECSF	05.07.2011	Malaise Trap
BT 1971	Eumolpinae sp_015	N115		ECSF	05.07.2011	Malaise Trap
BT 1972	Eumolpinae sp_015	N115		ECSF	05.07.2011	Malaise Trap
BT 1973	Eumolpinae sp_015	N115		ECSF	05.07.2011	Malaise Trap
BT 1974	Alticinae sp_015	N115		ECSF	05.07.2011	Malaise Trap
BT 1975	Alticinae sp_015	N115		ECSF	05.07.2011	Malaise Trap
BT 1976	Galerucinae sp_102	N115		ECSF	05.07.2011	Malaise Trap
BT 1977	Alticinae sp_007	N115		ECSF	05.07.2011	Malaise Trap
BT 1978	Eumolpinae sp_040	N115		ECSF	05.07.2011	Malaise Trap
BT 1981	Galerucinae sp_065	N115		ECSF	05.07.2011	Malaise Trap
BT 1982	Alticinae sp_062	N115		ECSF	05.07.2011	Malaise Trap
BT 2050	Alticinae sp_123	N115		ECSF	05.07.2011	Malaise Trap
BT 2054	Alticinae sp_139	N115		ECSF	05.07.2011	Malaise Trap
BT 2055	Alticinae sp_145	N115		ECSF	05.07.2011	Malaise Trap
BT 2056	Alticinae sp_141	N115		ECSF	05.07.2011	Malaise Trap
BT 2059	Alticinae sp_142	N115		ECSF	05.07.2011	Malaise Trap
BT 2065	Alticinae sp_142	N115		ECSF	05.07.2011	Malaise Trap
BT 2067	Alticinae sp_142	N115		ECSF	05.07.2011	Malaise Trap
BT 2068	Alticinae sp_142	N115		ECSF	05.07.2011	Malaise Trap
BT 2073	Alticinae sp_022	N116	KJ677340	ECSF	Q2	05.07.2011
BT 2074	Galerucinae sp_034	N116	KJ677698	ECSF	Q2	05.07.2011
BT 2075	Galerucinae sp_034	N116		ECSF	Q2	05.07.2011
BT 2076	Alticinae sp_002	N116	KJ677706	ECSF	Q2	05.07.2011
BT 2077	Galerucinae sp_007	N116	KJ677542	ECSF	Q2	05.07.2011
BT 2078	Alticinae sp_265	N116	KJ677431	ECSF	Q2	05.07.2011
BT 2079	Eumolpinae sp_038	N116		ECSF	Q2	05.07.2011
BT 2080	Galerucinae sp_065	N116		ECSF	Q2	05.07.2011
BT 2081	Eumolpinae sp_042	N116	KJ677860	ECSF	Q2	05.07.2011
BT 2082	Alticinae sp_064	N116	KJ677448	ECSF	Q2	05.07.2011
BT 2083	Alticinae sp_064	N116		ECSF	Q2	05.07.2011
BT 2084	Alticinae sp_032	N116	KJ677624	ECSF	Q2	05.07.2011
BT 2085	Eumolpinae sp_042	N116	KJ677861	ECSF	Q2	05.07.2011
BT 2086	Eumolpinae sp_042	N116		ECSF	Q2	05.07.2011
BT 2087	Alticinae sp_049	N116	KJ677608	ECSF	Q2	05.07.2011
BT 2088	Alticinae sp_049	N116		ECSF	Q2	05.07.2011
BT 2089	Alticinae sp_049	N116		ECSF	Q2	05.07.2011
BT 2090	Alticinae sp_081	N116	KJ677766	ECSF	Q2	05.07.2011
BT 2091	Alticinae sp_081	N116		ECSF	Q2	05.07.2011
BT 2092	Alticinae sp_081	N116		ECSF	Q2	05.07.2011
BT 2093	Alticinae sp_096	N116	KJ677466	ECSF	Q2	05.07.2011
BT 2094	Alticinae sp_096	N116		ECSF	Q2	05.07.2011
BT 2095	Alticinae sp_096	N116		ECSF	Q2	05.07.2011
BT 2096	Alticinae sp_096	N116		ECSF	Q2	05.07.2011
BT 2097	Alticinae sp_122	N116	KJ677685	ECSF	Q2	05.07.2011
BT 2098	Alticinae sp_122	N116		ECSF	Q2	05.07.2011
BT 2099	Alticinae sp_122	N116		ECSF	Q2	05.07.2011
BT 2100	Alticinae sp_122	N116		ECSF	Q2	05.07.2011
BT 2101	Alticinae sp_086	N116	KJ677402	ECSF	Q2	05.07.2011
BT 2102	Alticinae sp_086	N116		ECSF	Q2	05.07.2011
BT 2103	Alticinae sp_086	N116		ECSF	Q2	05.07.2011
BT 2104	Alticinae sp_086	N116		ECSF	Q2	05.07.2011
BT 2105	Alticinae sp_086	N116		ECSF	Q2	05.07.2011
BT 2106	Alticinae sp_086	N116		ECSF	Q2	05.07.2011
BT 2107	Alticinae sp_085	N116	KJ677390	ECSF	Q2	05.07.2011
BT 2108	Alticinae sp_085	N116		ECSF	Q2	05.07.2011
BT 2109	Alticinae sp_133	N116	KJ677587	ECSF	Q2	05.07.2011
BT 2110	Alticinae sp_133	N116		ECSF	Q2	05.07.2011
BT 2111	Alticinae sp_133	N116		ECSF	Q2	05.07.2011
BT 2112	Alticinae sp_133	N116		ECSF	Q2	05.07.2011
BT 2113	Alticinae sp_142	N116	KJ677594	ECSF	Q2	05.07.2011
BT 2114	Alticinae sp_142	N116		ECSF	Q2	05.07.2011
BT 2115	Alticinae sp_142	N116		ECSF	Q2	05.07.2011
BT 2116	Alticinae sp_142	N116		ECSF	Q2	05.07.2011
BT 2117	Alticinae sp_142	N116		ECSF	Q2	05.07.2011
BT 2118	Alticinae sp_142	N116		ECSF	Q2	05.07.2011

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT_2119	Eumolpinae_sp_016	N116	KJ677936	ECSF	Q2	05.07.2011	Malaise Trap
BT_2120	Eumolpinae_sp_016	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2121	Eumolpinae_sp_016	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2122	Eumolpinae_sp_016	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2123	Alticinae_sp_018	N116	KJ677281	ECSF	Q2	05.07.2011	Malaise Trap
BT_2124	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2125	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2126	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2127	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2128	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2129	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2130	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2131	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2132	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2133	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2134	Alticinae_sp_018	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2135	Alticinae_sp_122	N116	KJ677686	ECSF	Q2	05.07.2011	Malaise Trap
BT_2136	Alticinae_sp_110	N116	KJ677673	ECSF	Q2	05.07.2011	Malaise Trap
BT_2137	Alticinae_sp_110	N116		ECSF	Q2	05.07.2011	Malaise Trap
BT_2138	Alticinae_sp_115	N116	KJ677294	ECSF	Q2	05.07.2011	Malaise Trap
BT_2143	Eumolpinae_sp_042	N117		ECSF	L6	05.07.2011	Flight-Intercept.
BT_2152	Eumolpinae_sp_038	N121	KJ677930	ECSF		14.07.2011	Hand-Coll.(N)
BT_2153	Eumolpinae_sp_038	N121		ECSF		14.07.2011	Hand-Coll.(N)
BT_2154	Galerucinae_sp_017	N121	KJ677554	ECSF		14.07.2011	Hand-Coll.(N)
BT_2155	Alticinae_sp_115	N121	KJ677296	ECSF		14.07.2011	Hand-Coll.(N)
BT_2156	Cassidinae_sp_012	N121	KJ677830	ECSF		14.07.2011	Hand-Coll.(N)
BT_2157	Eumolpinae_sp_024	N121	KJ677911	ECSF		14.07.2011	Hand-Coll.(N)
BT_2158	Cassidinae_sp_083	N121	KJ677338	ECSF		14.07.2011	Hand-Coll.(N)
BT_2161	Cassidinae_sp_012	N122	KJ677827	ECSF		19.07.2011	Hand-Coll.(N)
BT_2162	Cassidinae_sp_012	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2163	Cassidinae_sp_012	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2164	Cassidinae_sp_012	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2165	Cassidinae_sp_012	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2166	Cassidinae_sp_012	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2167	Cassidinae_sp_012	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2168	Galerucinae_sp_036	N122	KJ677763	ECSF		19.07.2011	Hand-Coll.(N)
BT_2169	Galerucinae_sp_033	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2170	Galerucinae_sp_034	N122	KJ677699	ECSF		19.07.2011	Hand-Coll.(N)
BT_2171	Galerucinae_sp_034	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2172	Galerucinae_sp_062	N122		ECSF		19.07.2011	Hand-Coll.(N)
BT_2173	Eumolpinae_sp_034	N122	KJ677882	ECSF		19.07.2011	Hand-Coll.(N)
BT_2176	Galerucinae_sp_043	N123	KJ677682	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2177	Galerucinae_sp_043	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2178	Galerucinae_sp_043	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2179	Alticinae_sp_011	N123	KJ677726	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2180	Alticinae_sp_075	N123	KJ677621	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2181	Alticinae_sp_251	N123	KJ677508	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2182	Galerucinae_sp_029	N123	KJ677521	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2183	Galerucinae_sp_029	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2184	Galerucinae_sp_029	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2185	Galerucinae_sp_029	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2186	Galerucinae_sp_029	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2187	Galerucinae_sp_029	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2188	Galerucinae_sp_029	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2189	Hispinae_sp_008	N123	KJ677845	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2190	Hispinae_sp_022	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2191	Criocerinae_sp_005	N123	KJ677820	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2192	Galerucinae_sp_028	N123	KJ677433	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2193	Alticinae_sp_076	N123	KJ677344	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2194	Alticinae_sp_127	N123	KJ677275	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2195	Alticinae_sp_127	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2196	Alticinae_sp_127	N123	KJ677278	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2197	Eumolpinae_sp_047	N123	KJ677919	Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2198	Eumolpinae_sp_047	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2199	Eumolpinae_sp_035	N123		Bombuscaro		21.07.2011	Hand-Coll.(N)
BT_2203	Alticinae_sp_016	N124		ECSF		26.07.2011	Hand-Coll.(N)
BT_2208	Alticinae_sp_112	N126	KJ677614	Bombuscaro		02.08.2011	Hand-Coll.(N)
BT_2218	Galerucinae_sp_012	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2219	Eumolpinae_sp_042	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2220	Galerucinae_sp_007	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2221	Galerucinae_sp_022	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2222	Galerucinae_sp_076	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2229	Alticinae_sp_063	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2231	Alticinae_sp_087	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2232	Alticinae_sp_087	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2233	Alticinae_sp_087	N128		ECSF	Q2	04.08.2011	Malaise Trap
BT_2310	Cassidinae_sp_004	N129		ECSF	U1	04.08.2011	Malaise Trap
BT_2311	Alticinae_sp_074	N129		ECSF	U1	04.08.2011	Malaise Trap
BT_2312	Alticinae_sp_042	N129	KJ677413	ECSF	U1	04.08.2011	Malaise Trap
BT_2313	Eumolpinae_sp_025	N129		ECSF	U1	04.08.2011	Malaise Trap
BT_2314	Eumolpinae_sp_030	N129		ECSF	U1	04.08.2011	Malaise Trap
BT_2317	Alticinae_sp_007	N130		ECSF	L6	04.08.2011	Flight-Intercept.
BT_2353	Hispinae_sp_006	N134		ECSF		09.08.2011	Hand-Coll.(N)
BT_2383	Galerucinae_sp_059	S130		ECSF	L6	01.06.2011	Sweep Netting
BT_2399	Alticinae_sp_038	S132		ECSF	L6	01.06.2011	Hand-Coll.(S)
BT_2405	Alticinae_sp_042	S133		ECSF	L5	01.06.2011	Sweep Netting
BT_2491	Alticinae_sp_085	S139	KJ677387	ECSF	U4	03.06.2011	Sweep Netting
BT_2492	Alticinae_sp_104	S139	KJ677329	ECSF	U4	03.06.2011	Sweep Netting
BT_2495	Alticinae_sp_243	S140	KJ677418	ECSF	U4	03.06.2011	Beating
BT_2496	Alticinae_sp_140	S140	KJ677573	ECSF	U4	03.06.2011	Beating
BT_2498	Alticinae_sp_052	S141	KJ677354	ECSF	U4	03.06.2011	Hand-Coll.(S)
BT_2499	Alticinae_sp_118	S143	KJ677668	Cajanuma	L2	07.06.2011	Beating
BT_2502	Alticinae_sp_160	S151	KJ677362	ECSF	L4	14.06.2011	Sweep Netting
BT_2503	Alticinae_sp_079	S151		ECSF	L4	14.06.2011	Sweep Netting
BT_2504	Galerucinae_sp_098	S151	KJ677627	ECSF	L4	14.06.2011	Sweep Netting
BT_2505	Galerucinae_sp_046	S151	KJ677635	ECSF	L4	14.06.2011	Sweep Netting
BT_2506	Alticinae_sp_047	S152	KJ677735	ECSF	L4	14.06.2011	Beating
BT_2507	Alticinae_sp_203	S152	KR424919	ECSF	L4	14.06.2011	Beating
BT_2508	Galerucinae_sp_034	S152		ECSF	L4	14.06.2011	Beating
BT_2509	Galerucinae_sp_034	S152		ECSF	L4	14.06.2011	Beating
BT_2510	Hispinae_sp_010	S154		ECSF	L2	16.06.2011	Sweep Netting
BT_2511	Hispinae_sp_010	S154		ECSF	L2	16.06.2011	Sweep Netting
BT_2516	Galerucinae_sp_009	S158	KJ677530	Bombuscaro	L1	21.06.2011	Beating
BT_2517	Alticinae_sp_201	S159	KJ677333	Bombuscaro	L1	21.06.2011	Hand-Coll.(S)
BT_2518	Alticinae_sp_070	S159	KJ677511	Bombuscaro	L1	21.06.2011	Hand-Coll.(S)
BT_2519	Alticinae_sp_153	S160	KJ677384	Bombuscaro	L2	21.06.2011	Sweep Netting
BT_2520	Alticinae_sp_073	S160		Bombuscaro	L2	21.06.2011	Sweep Netting
BT_2521	Galerucinae_sp_082	S161	KJ677688	Bombuscaro	L2	21.06.2011	Beating
BT_2522	Alticinae_sp_051	S161	KJ677365	Bombuscaro	L2	21.06.2011	Beating
BT_2523	Alticinae_sp_153	S161	KJ677385	Bombuscaro	L2	21.06.2011	Beating
BT_2524	Alticinae_sp_153	S161		Bombuscaro	L2	21.06.2011	Beating
BT_2525	Alticinae_sp_153	S161		Bombuscaro	L2	21.06.2011	Beating
BT_2526	Alticinae_sp_153	S161		Bombuscaro	L2	21.06.2011	Beating

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method	
BT 2528	Eumolpinae sp 045	S162	Bombuscaro	L2	21.06.2011	Hand-Coll.(S)	
BT 2529	Galerucinae sp 055	S163	Bombuscaro	L3	21.06.2011	Sweep Netting	
BT 2531	Alticinae sp 008	S166	ECSF	U1	23.06.2011	Sweep Netting	
BT 2544	Alticinae sp 097	S178	KJ677312	L3	28.06.2011	Sweep Netting	
BT 2546	Cassidinae sp 004	S180	KJ677853	L3	28.06.2011	Hand-Coll.(S)	
BT 2547	Cassidinae sp 004	S180	ECSF	L3	28.06.2011	Hand-Coll.(S)	
BT 2548	Cassidinae sp 012	S180	KJ677828	L3	28.06.2011	Hand-Coll.(S)	
BT 2549	Alticinae sp 094	S184	Cajanuma	U2	30.06.2011	Sweep Netting	
BT 2550	Alticinae sp 056	S185	KJ677314	U2	30.06.2011	Beating	
BT 2551	Alticinae sp 246	S185	KR424924	U2	30.06.2011	Beating	
BT 2552	Alticinae sp 246	S185	Cajanuma	U2	30.06.2011	Beating	
BT 2557	Galerucinae sp 007	S187	ECSF	U2	07.07.2011	Sweep Netting	
BT 2572	Alticinae sp 018	S202	KJ677428	L5	14.07.2011	Sweep Netting	
BT 2573	Cassidinae sp 012	S203	KJ677829	L5	14.07.2011	Beating	
BT 2574	Cassidinae sp 012	S203	ECSF	L5	14.07.2011	Beating	
BT 2575	Eumolpinae sp 031	S203	KJ677939	L5	14.07.2011	Beating	
BT 2576	Alticinae sp 096	S203	ECSF	L5	14.07.2011	Beating	
BT 2578	Galerucinae sp 034	S204	KJ677700	L5	14.07.2011	Hand-Coll.(S)	
BT 2579	Alticinae sp 115	S204	KJ677299	L5	14.07.2011	Hand-Coll.(S)	
BT 2594	Galerucinae sp 002	S207	ECSF	L6	14.07.2011	Hand-Coll.(S)	
BT 2600	Alticinae sp 008	S211	KR424922	U5	19.07.2011	Sweep Netting	
BT 2601	Galerucinae sp 007	S212	KR425306	U5	19.07.2011	Beating	
BT 2602	Alticinae sp 085	S212	KR424923	U5	19.07.2011	Beating	
BT 2603	Eumolpinae sp 039	S212	KR424811	U5	19.07.2011	Beating	
BT 2604	Alticinae sp 106	S212	ECSF	U5	19.07.2011	Beating	
BT 2605	Alticinae sp 118	S212	KR424925	U5	19.07.2011	Beating	
BT 2606	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2607	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2608	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2609	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2610	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2611	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2612	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2613	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2614	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2615	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2616	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2617	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2618	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2619	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2620	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2621	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2622	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2623	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2624	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2625	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2626	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2627	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2628	Alticinae sp 118	S212	ECSF	U5	19.07.2011	Beating	
BT 2629	Eumolpinae sp 029	S215	KJ677903	U6	19.07.2011	Beating	
BT 2631	Alticinae sp 135	S220	KJ677581	L6	21.07.2011	Sweep Netting	
BT 2632	Alticinae sp 143	S221	KJ677577	L6	21.07.2011	Beating	
BT 2634	Alticinae sp 196	S223	Bombuscaro	L5	21.07.2011	Sweep Netting	
BT 2637	Eumolpinae sp 040	S226	KJ677868	L2	26.07.2011	Sweep Netting	
BT 2638	Galerucinae sp 061	S226	KJ677518	L2	26.07.2011	Sweep Netting	
BT 2639	Alticinae sp 061	S226	KR424926	L2	26.07.2011	Sweep Netting	
BT 2640	Alticinae sp 104	S226	KJ677650	L2	26.07.2011	Sweep Netting	
BT 2641	Alticinae sp 097	S226	KJ677304	L2	26.07.2011	Sweep Netting	
BT 2642	Eumolpinae sp 019	S227	KJ677881	L2	26.07.2011	Beating	
BT 2643	Alticinae sp 042	S227	KJ677414	L2	26.07.2011	Beating	
BT 2644	Criocerinae sp 001	S227	KJ677816	L2	26.07.2011	Beating	
BT 2645	Eumolpinae sp 038	S227	ECSF	L2	26.07.2011	Beating	
BT 2646	Alticinae sp 096	S227	KJ677488	L2	26.07.2011	Beating	
BT 2657	Alticinae sp 001	S232	KJ677704	L1	26.07.2011	Sweep Netting	
BT 2658	Alticinae sp 041	S232	ECSF	L1	26.07.2011	Sweep Netting	
BT 2659	Alticinae sp 086	S232	KJ677395	L1	26.07.2011	Sweep Netting	
BT 2660	Alticinae sp 097	S232	KR424927	L1	26.07.2011	Sweep Netting	
BT 2661	Hispinae sp 009	S232	KJ677846	L1	26.07.2011	Sweep Netting	
BT 2662	Alticinae sp 087	S233	KJ677505	L1	26.07.2011	Beating	
BT 2663	Hispinae sp 005	S233	KJ677841	L1	26.07.2011	Beating	
BT 2664	Cassidinae sp 003	S234	ECSF	L1	26.07.2011	Hand-Coll.(S)	
BT 2665	Criocerinae sp 004	S234	KJ677819	ECSF	L1	26.07.2011	Hand-Coll.(S)
BT 2666	Alticinae sp 089	S235	KJ677736	Cajanuma	L4	28.07.2011	Sweep Netting
BT 2668	Eumolpinae sp 028	S236	Cajanuma	L4	28.07.2011	Beating	
BT 2670	Alticinae sp 051	S244	KJ677371	Bombuscaro	U1	02.08.2011	Sweep Netting
BT 2671	Galerucinae sp 072	S247	KJ677806	Bombuscaro	U2	02.08.2011	Sweep Netting
BT 2672	Alticinae sp 051	S247	KJ677372	Bombuscaro	U2	02.08.2011	Sweep Netting
BT 2673	Galerucinae sp 049	S248	KJ677644	Bombuscaro	U2	02.08.2011	Beating
BT 2675	Galerucinae sp 002	S252	KR425307	ECSF	L4	09.08.2011	Sweep Netting
BT 2676	Galerucinae sp 046	S254	KR425308	ECSF	L4	09.08.2011	Beating
BT 2677	Galerucinae sp 061	S254	KR425309	ECSF	L4	09.08.2011	Beating
BT 2678	Alticinae sp 111	S254	ECSF	L4	09.08.2011	Beating	
BT 2679	Alticinae sp 096	S254	KR424928	ECSF	L4	09.08.2011	Beating
BT 2681	Criocerinae sp 001	S255	KR425411	ECSF	L4	09.08.2011	Hand-Coll.(S)
BT 2684	Alticinae sp 049	S259	ECSF	L6	09.08.2011	Sweep Netting	
BT 2697	Alticinae sp 053	S265	KJ677345	Cajanuma	U1	11.08.2011	Sweep Netting
BT 2698	Eumolpinae sp 071	S266	KJ677908	Cajanuma	U1	11.08.2011	Beating
BT 2699	Alticinae sp 060	S266	KR424929	Cajanuma	U1	11.08.2011	Beating
BT 2701	Alticinae sp 056	S268	Cajanuma	L1	11.08.2011	Sweep Netting	
BT 2702	Alticinae sp 056	S268	Cajanuma	L1	11.08.2011	Sweep Netting	
BT 2703	Alticinae sp 056	S268	Cajanuma	L1	11.08.2011	Sweep Netting	
BT 2704	Alticinae sp 246	S268	Cajanuma	L1	11.08.2011	Sweep Netting	
BT 2705	Alticinae sp 094	S268	KJ677605	Cajanuma	L1	11.08.2011	Sweep Netting
BT 2706	Alticinae sp 120	S268	Cajanuma	L1	11.08.2011	Sweep Netting	
BT 2707	Alticinae sp 034	S269	KJ677626	Cajanuma	L1	11.08.2011	Beating
BT 2708	Alticinae sp 224	S269	KR424930	Cajanuma	L1	11.08.2011	Beating
BT 2709	Hispinae sp 024	S269	Cajanuma	L1	11.08.2011	Beating	
BT 2717	Alticinae sp 001	S274	KR424931	ECSF	L1	16.08.2011	Sweep Netting
BT 2718	Alticinae sp 096	S275	KR424932	ECSF	L1	16.08.2011	Beating
BT 2719	Criocerinae sp 001	S276	KR425412	ECSF	L1	16.08.2011	Hand-Coll.(S)
BT 2723	Alticinae sp 225	S278	ECSF	L2	16.08.2011	Beating	
BT 2771	Galerucinae sp 100	S282	ECSF	L4	18.08.2011	Hand-Coll.(S)	
BT 2787	Galerucinae sp 002	S285	ECSF	L5	18.08.2011	Hand-Coll.(S)	
BT 2795	Galerucinae sp 017	S286	ECSF	L6	18.08.2011	Sweep Netting	
BT 2805	Eumolpinae sp 006	S292	Bombuscaro	U5	23.08.2011	Sweep Netting	
BT 2806	Alticinae sp 050	S295	KR424933	Bombuscaro	U6	23.08.2011	Sweep Netting
BT 2807	Alticinae sp 103	S295	KR424934	Bombuscaro	U6	23.08.2011	Sweep Netting
BT 2809	Alticinae sp 259	S296	KR424935	Bombuscaro	U6	23.08.2011	Beating
BT 2810	Galerucinae sp 072	S298	KR425310	ECSF	U1	25.08.2011	Sweep Netting
BT 2812	Alticinae sp 104	S299	KR424936	ECSF	U1	25.08.2011	Beating
BT 2813	Alticinae sp 104	S299	ECSF	U1	25.08.2011	Beating	
BT 2814	Alticinae sp 104	S299	ECSF	U1	25.08.2011	Beating	
BT 2816	Alticinae sp 150	S301	KR424937	ECSF	U1	25.08.2011	Sweep Netting
BT 2817	Alticinae sp 150	S301	ECSF	U2	25.08.2011	Sweep Netting	
BT 2818	Alticinae sp 051	S301	KR424938	ECSF	U2	25.08.2011	Sweep Netting

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SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT 2819	Eumolpinae_sp_030	S302	KR424812	ECSF	U2	25.08.2011	Beating
BT 2820	Eumolpinae_sp_030	S302	KR424812	ECSF	U2	25.08.2011	Beating
BT 2821	Eumolpinae_sp_074	S302	KR424813	ECSF	U2	25.08.2011	Beating
BT 2822	Galerucinae_sp_048	S302	KR425311	ECSF	U2	25.08.2011	Beating
BT 2823	Alticinae_sp_068	S302	KR424939	ECSF	U2	25.08.2011	Beating
BT 2824	Alticinae_sp_129	S302	KR424940	ECSF	U2	25.08.2011	Beating
BT 2825	Alticinae_sp_126	S304	KR424941	ECSF	L3	25.08.2011	Sweep Netting
BT 2826	Alticinae_sp_104	S304	KR424942	ECSF	L3	25.08.2011	Sweep Netting
BT 2827	Eumolpinae_sp_067	S304	KR424814	ECSF	L3	25.08.2011	Sweep Netting
BT 2828	Eumolpinae_sp_029	S304	KR424815	ECSF	L3	25.08.2011	Sweep Netting
BT 2829	Eumolpinae_sp_030	S305	KR424815	ECSF	L3	25.08.2011	Beating
BT 2830	Alticinae_sp_051	S305	KR424943	ECSF	L3	25.08.2011	Beating
BT 2831	Cassidinae_sp_015	S307	KR424784	Bombuscaro	L1	30.08.2011	Sweep Netting
BT 2832	Alticinae_sp_072	S311	KR424944	Bombuscaro	L2	30.08.2011	Beating
BT 2833	Alticinae_sp_104	S313	KR424945	Bombuscaro	L3	30.08.2011	Sweep Netting
BT 2834	Alticinae_sp_118	S316	KR424946	ECSF	U4	01.09.2011	Beating
BT 2835	Alticinae_sp_118	S316		ECSF	U4	01.09.2011	Sweep Netting
BT 2837	Alticinae_sp_118	S317		ECSF	U4	01.09.2011	Beating
BT 2838	Alticinae_sp_118	S317		ECSF	U4	01.09.2011	Beating
BT 2839	Eumolpinae_sp_066	S317	KR424816	ECSF	U4	01.09.2011	Beating
BT 2840	Alticinae_sp_052	S317		ECSF	U4	01.09.2011	Beating
BT 2841	Galerucinae_sp_096	S317	KR425312	ECSF	U4	01.09.2011	Beating
BT 2842	Galerucinae_sp_098	S317	KR425313	ECSF	U4	01.09.2011	Beating
BT 2843	Hispininae_sp_005	S317		ECSF	U4	01.09.2011	Beating
BT 2844	Alticinae_sp_126	S319	KR424947	ECSF	U5	01.09.2011	Sweep Netting
BT 2845	Alticinae_sp_118	S319		ECSF	U5	01.09.2011	Sweep Netting
BT 2846	Alticinae_sp_152	S319	KR424948	ECSF	U5	01.09.2011	Sweep Netting
BT 2847	Alticinae_sp_152	S319		ECSF	U5	01.09.2011	Sweep Netting
BT 2848	Alticinae_sp_118	S320		ECSF	U5	01.09.2011	Beating
BT 2849	Alticinae_sp_118	S320		ECSF	U5	01.09.2011	Beating
BT 2850	Alticinae_sp_118	S320	KR424949	ECSF	U5	01.09.2011	Beating
BT 2851	Alticinae_sp_118	S320		ECSF	U5	01.09.2011	Beating
BT 2852	Alticinae_sp_052	S320	KR424950	ECSF	U5	01.09.2011	Beating
BT 2853	Alticinae_sp_052	S320		ECSF	U5	01.09.2011	Beating
BT 2854	Eumolpinae_sp_017	S320		ECSF	U5	01.09.2011	Beating
BT 2855	Galerucinae_sp_098	S320	KR425314	ECSF	U5	01.09.2011	Beating
BT 2856	Cryptocephalinae_sp_002	S320		ECSF	U5	01.09.2011	Beating
BT 2863	Alticinae_sp_104	S321	KR424951	ECSF	U5	01.09.2011	Hand-Coll.(S)
BT 2864	Eumolpinae_sp_039	S321	KR424817	ECSF	U5	01.09.2011	Hand-Coll.(S)
BT 2865	Alticinae_sp_118	S321		ECSF	U5	01.09.2011	Hand-Coll.(S)
BT 2866	Alticinae_sp_118	S321		ECSF	U5	01.09.2011	Hand-Coll.(S)
BT 2867	Alticinae_sp_118	S321		ECSF	U5	01.09.2011	Hand-Coll.(S)
BT 2868	Alticinae_sp_118	S321		ECSF	U5	01.09.2011	Hand-Coll.(S)
BT 2869	Alticinae_sp_118	S321		ECSF	U5	01.09.2011	Hand-Coll.(S)
BT 2870	Alticinae_sp_118	S322	KR424952	ECSF	U6	01.09.2011	Sweep Netting
BT 2871	Alticinae_sp_213	S322	KR424953	ECSF	U6	01.09.2011	Sweep Netting
BT 2872	Alticinae_sp_064	S323	KR424954	ECSF	U6	01.09.2011	Beating
BT 2873	Alticinae_sp_053	S326	KR424955	Cajanuma	U1	06.09.2011	Beating
BT 2874	Alticinae_sp_094	S328		Cajanuma	U2	06.09.2011	Sweep Netting
BT 2876	Alticinae_sp_087	S334	KR424956	ECSF	U3	08.09.2011	Sweep Netting
BT 2877	Eumolpinae_sp_024	S335	KR424818	ECSF	U3	08.09.2011	Beating
BT 2878	Eumolpinae_sp_024	S335		ECSF	U3	08.09.2011	Beating
BT 2879	Alticinae_sp_164	S335	KR424957	ECSF	U3	08.09.2011	Beating
BT 2880	Eumolpinae_sp_060	S335	KR424819	ECSF	U3	08.09.2011	Beating
BT 2881	Alticinae_sp_085	S335	KR424958	ECSF	U3	08.09.2011	Beating
BT 2900	Alticinae_sp_245	S343	KR424959	Cajanuma	U4	14.09.2012	Sweep Netting
BT 2901	Alticinae_sp_244	S343	KR424960	Cajanuma	U4	14.09.2012	Sweep Netting
BT 2902	Alticinae_sp_208	S343	KR424961	Cajanuma	U4	14.09.2012	Sweep Netting
BT 2903	Eumolpinae_sp_071	S344	KR424820	Cajanuma	U4	14.09.2012	Beating
BT 2905	Alticinae_sp_245	S347	KR424962	Cajanuma	U5	14.09.2012	Beating
BT 2907	Alticinae_sp_191	S350	KR424963	Cajanuma	U6	14.09.2012	Beating
BT 2908	Alticinae_sp_258	S350	KR424964	Cajanuma	U6	14.09.2012	Beating
BT 2912	Galerucinae_sp_014	S353		ECSF	U1	19.09.2012	Beating
BT 2935	Galerucinae_sp_106	S362		Bombuscaro	U1	21.09.2012	Beating
BT 2938	Alticinae_sp_234	S364		Cajanuma	L1	27.09.2012	Sweep Netting
BT 2939	Alticinae_sp_235	S364		Cajanuma	L1	27.09.2012	Sweep Netting
BT 2948	Galerucinae_sp_014	S373		ECSF	L4	29.09.2012	Sweep Netting
BT 2949	Hispininae_sp_007	S374		ECSF	L4	29.09.2012	Beating
BT 2954	Hispininae_sp_018	S374		ECSF	L4	29.09.2012	Beating
BT 2969	Eumolpinae_sp_013	S375		ECSF	L4	29.09.2012	Hand-Coll.(S)
BT 2999	Eumolpinae_sp_003	S379		ECSF	L6	29.09.2012	Sweep Netting
BT 3026	Galerucinae_sp_029	S382	KR425315	Bombuscaro	L4	04.10.2011	Sweep Netting
BT 3027	Alticinae_sp_173	S382	KR424966	Bombuscaro	L4	04.10.2011	Sweep Netting
BT 3028	Alticinae_sp_204	S382	KR424967	Bombuscaro	L4	04.10.2011	Sweep Netting
BT 3029	Eumolpinae_sp_017	S383		Bombuscaro	L4	04.10.2011	Beating
BT 3030	Alticinae_sp_193	S383	KR424968	Bombuscaro	L4	04.10.2011	Beating
BT 3031	Eumolpinae_sp_055	S385	KR424821	Bombuscaro	L5	04.10.2011	Sweep Netting
BT 3032	Galerucinae_sp_106	S385	KR425316	Bombuscaro	L5	04.10.2011	Sweep Netting
BT 3033	Alticinae_sp_254	S385	KR424969	Bombuscaro	L5	04.10.2011	Sweep Netting
BT 3034	Alticinae_sp_181	S385	KR424970	Bombuscaro	L5	04.10.2011	Sweep Netting
BT 3035	Galerucinae_sp_051	S386	KR425317	Bombuscaro	L5	04.10.2011	Beating
BT 3036	Alticinae_sp_042	S388	KR424971	Bombuscaro	L6	04.10.2011	Sweep Netting
BT 3037	Alticinae_sp_135	S388	KR424972	Bombuscaro	L6	04.10.2011	Sweep Netting
BT 3038	Galerucinae_sp_084	S389	KR425318	Bombuscaro	L6	04.10.2011	Beating
BT 3039	Eumolpinae_sp_040	S389	KR424822	Bombuscaro	L6	04.10.2011	Beating
BT 3040	Eumolpinae_sp_040	S389		Bombuscaro	L6	04.10.2011	Beating
BT 3041	Cassidinae_sp_017	S389	KR424785	Bombuscaro	L6	04.10.2011	Beating
BT 3060	Alticinae_sp_115	S394	KR424973	ECSF	U5	06.10.2011	Sweep Netting
BT 3061	Alticinae_sp_115	S394		ECSF	U5	06.10.2011	Sweep Netting
BT 3062	Alticinae_sp_249	S394	KR424974	ECSF	U5	06.10.2011	Sweep Netting
BT 3063	Alticinae_sp_249	S394		ECSF	U5	06.10.2011	Sweep Netting
BT 3064	Alticinae_sp_253	S394	KR424975	ECSF	U5	06.10.2011	Sweep Netting
BT 3065	Galerucinae_sp_034	S394	KR425319	ECSF	U5	06.10.2011	Sweep Netting
BT 3066	Alticinae_sp_147	S394	KR424976	ECSF	U5	06.10.2011	Sweep Netting
BT 3067	Alticinae_sp_147	S394		ECSF	U5	06.10.2011	Sweep Netting
BT 3068	Alticinae_sp_074	S394		ECSF	U5	06.10.2011	Sweep Netting
BT 3070	Eumolpinae_sp_038	S395	KR424823	ECSF	U5	06.10.2011	Beating
BT 3071	Eumolpinae_sp_038	S395		ECSF	U5	06.10.2011	Beating
BT 3072	Eumolpinae_sp_038	S395		ECSF	U5	06.10.2011	Beating
BT 3073	Eumolpinae_sp_038	S395		ECSF	U5	06.10.2011	Beating
BT 3074	Alticinae_sp_096	S395	KR424977	ECSF	U5	06.10.2011	Beating
BT 3075	Alticinae_sp_096	S395		ECSF	U5	06.10.2011	Beating
BT 3076	Criocerinae_sp_001	S395	KR425413	ECSF	U5	06.10.2011	Beating
BT 3077	Alticinae_sp_064	S395		ECSF	U5	06.10.2011	Beating
BT 3079	Alticinae_sp_104	S397	KR424978	ECSF	U6	06.10.2011	Sweep Netting
BT 3080	Alticinae_sp_112	S397	KR424979	ECSF	U6	06.10.2011	Sweep Netting
BT 3081	Alticinae_sp_205	S397	KR424980	ECSF	U6	06.10.2011	Sweep Netting
BT 3082	Galerucinae_sp_012	S397	KR425320	ECSF	U6	06.10.2011	Sweep Netting
BT 3083	Alticinae_sp_096	S397	KR424981	ECSF	U6	06.10.2011	Sweep Netting
BT 3084	Alticinae_sp_062	S397	KR424982	ECSF	U6	06.10.2011	Sweep Netting
BT 3085	Alticinae_sp_064	S397	KR424983	ECSF	U6	06.10.2011	Sweep Netting
BT 3086	Alticinae_sp_115	S397	KR424984	ECSF	U6	06.10.2011	Sweep Netting
BT 3087	Eumolpinae_sp_038	S397	KR424824	ECSF	U6	06.10.2011	Sweep Netting
BT 3088	Alticinae_sp_061	S397	KR424985	ECSF	U6	06.10.2011	Sweep Netting
BT 3089	Eumolpinae_sp_034	S397	KR424825	ECSF	U6	06.10.2011	Sweep Netting

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SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method	
BT 3090	Eumolpinae sp 065	S398	KR424826	ECSF	U6	06.10.2011	Beating
BT 3091	Eumolpinae sp 065	S398		ECSF	U6	06.10.2011	Beating
BT 3092	Eumolpinae sp 002	S398	KR424827	ECSF	U6	06.10.2011	Beating
BT 3093	Alticinae sp 051	S398	KR424986	ECSF	U6	06.10.2011	Beating
BT 3094	Alticinae sp 051	S398		ECSF	U6	06.10.2011	Beating
BT 3095	Alticinae sp 254	S398	KR424987	ECSF	U6	06.10.2011	Beating
BT 3097	Eumolpinae sp 038	S399	KR424828	ECSF	U6	06.10.2011	Hand-Coll.(S)
BT 3098	Eumolpinae sp 038	S399		ECSF	U6	06.10.2011	Hand-Coll.(S)
BT 3099	Eumolpinae sp 038	S399		ECSF	U6	06.10.2011	Hand-Coll.(S)
BT 3100	Alticinae sp 172	S400	KR424988	Cajanuma	L4	11.10.2011	Sweep Netting
BT 3101	Alticinae sp 245	S400	KR424989	Cajanuma	L4	11.10.2011	Sweep Netting
BT 3102	Alticinae sp 244	S403	KR424990	Cajanuma	L5	11.10.2011	Sweep Netting
BT 3103	Alticinae sp 121	S404		Cajanuma	L5	11.10.2011	Beating
BT 3104	Alticinae sp 244	S404	KR424991	Cajanuma	L5	11.10.2011	Beating
BT 3105	Alticinae sp 244	S404		Cajanuma	L5	11.10.2011	Beating
BT 3106	Alticinae sp 171	S407	KR424992	Cajanuma	L6	11.10.2011	Beating
BT 3107	Galerucinae sp 066	S407	KR425321	Cajanuma	L6	11.10.2011	Beating
BT 3108	Alticinae sp 001	S409	KR424993	ECSF	U1	13.10.2011	Sweep Netting
BT 3109	Galerucinae sp 109	S409		ECSF	U1	13.10.2011	Sweep Netting
BT 3110	Galerucinae sp 048	S409	KR425322	ECSF	U1	13.10.2011	Sweep Netting
BT 3111	Alticinae sp 065	S409	KR424994	ECSF	U1	13.10.2011	Sweep Netting
BT 3112	Alticinae sp 197	S409	KR424995	ECSF	U1	13.10.2011	Sweep Netting
BT 3113	Alticinae sp 150	S409	KR424996	ECSF	U1	13.10.2011	Sweep Netting
BT 3114	Alticinae sp 150	S409	KR424997	ECSF	U1	13.10.2011	Sweep Netting
BT 3115	Galerucinae sp 098	S409	KR425323	ECSF	U1	13.10.2011	Sweep Netting
BT 3116	Galerucinae sp 098	S409		ECSF	U1	13.10.2011	Sweep Netting
BT 3117	Alticinae sp 163	S409	KR424998	ECSF	U1	13.10.2011	Sweep Netting
BT 3118	Alticinae sp 178	S409	KR424999	ECSF	U1	13.10.2011	Sweep Netting
BT 3120	Galerucinae sp 098	S410		ECSF	U1	13.10.2011	Beating
BT 3121	Hispaniae sp 025	S410	KR424795	ECSF	U1	13.10.2011	Beating
BT 3122	Alticinae sp 005	S410		ECSF	U1	13.10.2011	Beating
BT 3123	Eumolpinae sp 065	S410		ECSF	U1	13.10.2011	Beating
BT 3124	Alticinae sp 087	S410	KR425000	ECSF	U1	13.10.2011	Beating
BT 3125	Alticinae sp 065	S410		ECSF	U1	13.10.2011	Beating
BT 3126	Eumolpinae sp 039	S410	KR424829	ECSF	U1	13.10.2011	Beating
BT 3127	Eumolpinae sp 039	S410		ECSF	U1	13.10.2011	Beating
BT 3128	Eumolpinae sp 039	S410		ECSF	U1	13.10.2011	Beating
BT 3143	Galerucinae sp 017	S415		ECSF	L1	18.10.2011	Sweep Netting
BT 3157	Galerucinae sp 024	S415		ECSF	L1	18.10.2011	Sweep Netting
BT 3177	Galerucinae sp 017	S418		ECSF	L3	18.10.2011	Sweep Netting
BT 3191	Galerucinae sp 017	S420		ECSF	L3	18.10.2011	Hand-Coll.(S)
BT 3197	Alticinae sp 228	S422		Cajanuma	U1	20.10.2011	Beating
BT 3198	Alticinae sp 119	S422		Cajanuma	U1	20.10.2011	Beating
BT 3200	Galerucinae sp 073	S424	KR425324	Cajanuma	U2	20.10.2011	Sweep Netting
BT 3201	Alticinae sp 255	S425	KR425001	Cajanuma	U2	20.10.2011	Beating
BT 3202	Alticinae sp 184	S425	KR425002	Cajanuma	U2	20.10.2011	Beating
BT 3203	Galerucinae sp 073	S427	KR425325	Cajanuma	U3	20.10.2011	Sweep Netting
BT 3204	Alticinae sp 172	S427	KR425003	Cajanuma	U3	20.10.2011	Sweep Netting
BT 3205	Eumolpinae sp 052	S427	KR424830	Cajanuma	U3	20.10.2011	Sweep Netting
BT 3206	Alticinae sp 245	S428		Cajanuma	U3	20.10.2011	Beating
BT 3207	Alticinae sp 197	S430	KR425004	Cajanuma	L4	24.10.2011	Sweep Netting
BT 3208	Alticinae sp 197	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3209	Alticinae sp 123	S430	KR425005	ECSF	L4	24.10.2011	Sweep Netting
BT 3210	Galerucinae sp 066	S430	KR425326	ECSF	L4	24.10.2011	Sweep Netting
BT 3211	Alticinae sp 052	S430	KR425006	ECSF	L4	24.10.2011	Sweep Netting
BT 3212	Criocerinae sp 001	S430	KR425414	ECSF	L4	24.10.2011	Sweep Netting
BT 3213	Eumolpinae sp 040	S430	KR424831	ECSF	L4	24.10.2011	Sweep Netting
BT 3214	Eumolpinae sp 014	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3215	Eumolpinae sp 014	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3216	Galerucinae sp 104	S430	KR425327	ECSF	L4	24.10.2011	Sweep Netting
BT 3217	Alticinae sp 087	S430	KR425007	ECSF	L4	24.10.2011	Sweep Netting
BT 3218	Alticinae sp 087	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3219	Alticinae sp 087	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3220	Galerucinae sp 034	S430	KR425328	ECSF	L4	24.10.2011	Sweep Netting
BT 3221	Galerucinae sp 034	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3222	Galerucinae sp 034	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3223	Galerucinae sp 034	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3224	Galerucinae sp 034	S430		ECSF	L4	24.10.2011	Sweep Netting
BT 3225	Galerucinae sp 025	S431	KR425329	ECSF	L4	24.10.2011	Beating
BT 3226	Galerucinae sp 025	S431	KR425330	ECSF	L4	24.10.2011	Beating
BT 3227	Galerucinae sp 025	S431		ECSF	L4	24.10.2011	Beating
BT 3228	Galerucinae sp 025	S431		ECSF	L4	24.10.2011	Beating
BT 3229	Galerucinae sp 025	S431		ECSF	L4	24.10.2011	Beating
BT 3230	Alticinae sp 052	S431	KR425008	ECSF	L4	24.10.2011	Beating
BT 3231	Alticinae sp 052	S431		ECSF	L4	24.10.2011	Beating
BT 3232	Alticinae sp 064	S431	KR425009	ECSF	L4	24.10.2011	Beating
BT 3233	Alticinae sp 097	S431	KR425010	ECSF	L4	24.10.2011	Beating
BT 3234	Eumolpinae sp 038	S431		ECSF	L4	24.10.2011	Beating
BT 3235	Alticinae sp 018	S431	KR425011	ECSF	L4	24.10.2011	Beating
BT 3236	Galerucinae sp 034	S431	KR425331	ECSF	L4	24.10.2011	Beating
BT 3237	Galerucinae sp 034	S431		ECSF	L4	24.10.2011	Beating
BT 3238	Galerucinae sp 034	S431		ECSF	L4	24.10.2011	Beating
BT 3239	Galerucinae sp 034	S431		ECSF	L4	24.10.2011	Beating
BT 3241	Alticinae sp 097	S432		ECSF	L4	24.10.2011	Beating
BT 3242	Galerucinae sp 025	S432	KR425332	ECSF	L4	24.10.2011	Hand-Coll.(S)
BT 3243	Alticinae sp 240	S432	KR425012	ECSF	L4	24.10.2011	Hand-Coll.(S)
BT 3244	Alticinae sp 052	S432	KR425013	ECSF	L4	24.10.2011	Hand-Coll.(S)
BT 3248	Alticinae sp 116	S432	KR425014	ECSF	L4	24.10.2011	Hand-Coll.(S)
BT 3249	Galerucinae sp 034	S432	KR425333	ECSF	L4	24.10.2011	Hand-Coll.(S)
BT 3250	Galerucinae sp 034	S432		ECSF	L4	24.10.2011	Hand-Coll.(S)
BT 3251	Galerucinae sp 034	S432		ECSF	L4	24.10.2011	Hand-Coll.(S)
BT 3252	Galerucinae sp 034	S432		ECSF	L4	24.10.2011	Hand-Coll.(S)
BT 3253	Alticinae sp 115	S433	KR425015	ECSF	L5	24.10.2011	Sweep Netting
BT 3254	Alticinae sp 115	S433		ECSF	L5	24.10.2011	Sweep Netting
BT 3255	Galerucinae sp 061	S433	KR425334	ECSF	L5	24.10.2011	Sweep Netting
BT 3256	Galerucinae sp 092	S433	KR425335	ECSF	L5	24.10.2011	Sweep Netting
BT 3257	Alticinae sp 212	S433	KR425016	ECSF	L5	24.10.2011	Sweep Netting
BT 3258	Galerucinae sp 066	S433	KR425336	ECSF	L5	24.10.2011	Sweep Netting
BT 3259	Galerucinae sp 066	S433		ECSF	L5	24.10.2011	Sweep Netting
BT 3260	Alticinae sp 109	S433	KR425017	ECSF	L5	24.10.2011	Sweep Netting
BT 3261	Alticinae sp 029	S433		ECSF	L5	24.10.2011	Sweep Netting
BT 3262	Alticinae sp 042	S434	KR425018	ECSF	L5	24.10.2011	Beating
BT 3263	Eumolpinae sp 040	S434	KR424832	ECSF	L5	24.10.2011	Beating
BT 3264	Galerucinae sp 026	S434	KR425337	ECSF	L5	24.10.2011	Beating
BT 3265	Eumolpinae sp 042	S434	KR424833	ECSF	L5	24.10.2011	Beating
BT 3266	Galerucinae sp 021	S434	KR425338	ECSF	L5	24.10.2011	Beating
BT 3266a	Alticinae sp 112	S434		ECSF	L5	24.10.2011	Beating
BT 3267	Galerucinae sp 098	S435	KR425339	ECSF	L5	24.10.2011	Hand-Coll.(S)
BT 3268	Alticinae sp 170	S435	KR425019	ECSF	L5	24.10.2011	Hand-Coll.(S)
BT 3269	Alticinae sp 111	S436		ECSF	L6	24.10.2011	Sweep Netting
BT 3270	Alticinae sp 092	S436	KR425020	ECSF	L6	24.10.2011	Sweep Netting
BT 3271	Alticinae sp 202	S436	KR425021	ECSF	L6	24.10.2011	Sweep Netting
BT 3272	Alticinae sp 096	S436		ECSF	L6	24.10.2011	Sweep Netting
BT 3273	Galerucinae sp 034	S436	KR425340	ECSF	L6	24.10.2011	Sweep Netting
BT 3274	Alticinae sp 087	S437	KR425022	ECSF	L6	24.10.2011	Beating

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT 3275	Alticinae sp 087	S437	ECSF	L6	24.10.2011	Beating	
BT 3276	Alticinae sp 087	S437	ECSF	L6	24.10.2011	Beating	
BT 3277	Eumolpinae sp 039	S437	KR424834	ECSF	L6	24.10.2011	Beating
BT 3278	Galerucinae sp 061	S437	KR425341	ECSF	L6	24.10.2011	Beating
BT 3279	Alticinae sp 061	S438	KR425023	ECSF	L6	24.10.2011	Hand-Coll.(S)
BT 3280	Alticinae sp 051	S439		Bombuscaro	U3	01.11.2011	Sweep Netting
BT 3281	Alticinae sp 051	S439		Bombuscaro	U3	01.11.2011	Sweep Netting
BT 3282	Eumolpinae sp 024	S439	KR424835	Bombuscaro	U3	01.11.2011	Sweep Netting
BT 3283	Eumolpinae sp 024	S440		Bombuscaro	U3	01.11.2011	Beating
BT 3284	Alticinae sp 260	S440	KR425024	Bombuscaro	U3	01.11.2011	Beating
BT 3299	Eumolpinae sp 039	S449	KR424836	ECSF	L2	03.11.2011	Beating
BT 3300	Eumolpinae sp 039	S449		ECSF	L2	03.11.2011	Beating
BT 3301	Alticinae sp 260	S449		ECSF	L2	03.11.2011	Beating
BT 3302	Alticinae sp 260	S449		ECSF	L2	03.11.2011	Beating
BT 3303	Hispiinae sp 005	S449	KR424796	ECSF	L2	03.11.2011	Beating
BT 3304	Alticinae sp 260	S449		ECSF	L2	03.11.2011	Beating
BT 3305	Galerucinae sp 028	S451	KR425342	Bombuscaro	U4	08.11.2011	Sweep Netting
BT 3306	Alticinae sp 254	S451	KR425026	Bombuscaro	U4	08.11.2011	Sweep Netting
BT 3307	Alticinae sp 181	S451	KR425027	Bombuscaro	U4	08.11.2011	Sweep Netting
BT 3308	Alticinae sp 051	S452	KR425028	Bombuscaro	U4	08.11.2011	Beating
BT 3309	Alticinae sp 051	S452	KR425029	Bombuscaro	U4	08.11.2011	Beating
BT 3310	Alticinae sp 063	S452	KR425030	Bombuscaro	U4	08.11.2011	Beating
BT 3311	Galerucinae sp 015	S452	KR425033	Bombuscaro	U4	08.11.2011	Beating
BT 3312	Eumolpinae sp 042	S452	KR424837	Bombuscaro	U4	08.11.2011	Beating
BT 3313	Eumolpinae sp 055	S452	KR424838	Bombuscaro	U4	08.11.2011	Beating
BT 3314	Eumolpinae sp 042	S452	KR424839	Bombuscaro	U4	08.11.2011	Beating
BT 3316	Alticinae sp 117	S453	KR425031	Bombuscaro	U4	08.11.2011	Hand-Coll.(S)
BT 3317	Eumolpinae sp 051	S454	KR424840	Bombuscaro	U5	08.11.2011	Sweep Netting
BT 3318	Alticinae sp 210	S454	KR425032	Bombuscaro	U5	08.11.2011	Sweep Netting
BT 3319	Alticinae sp 086	S454	KR425033	Bombuscaro	U5	08.11.2011	Beating
BT 3320	Galerucinae sp 078	S454	KR425344	Bombuscaro	U5	08.11.2011	Sweep Netting
BT 3321	Alticinae sp 035	S454	KR425034	Bombuscaro	U5	08.11.2011	Sweep Netting
BT 3322	Eumolpinae sp 042	S455	KR424841	Bombuscaro	U5	08.11.2011	Beating
BT 3323	Eumolpinae sp 042	S455		Bombuscaro	U5	08.11.2011	Beating
BT 3324	Eumolpinae sp 042	S455		Bombuscaro	U5	08.11.2011	Beating
BT 3325	Galerucinae sp 086	S455	KR425345	Bombuscaro	U5	08.11.2011	Beating
BT 3326	Eumolpinae sp 017	S455		Bombuscaro	U5	08.11.2011	Beating
BT 3327	Eumolpinae sp 050	S456	KR424842	Bombuscaro	U5	08.11.2011	Hand-Coll.(S)
BT 3328	Alticinae sp 210	S457	KR425035	Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3329	Alticinae sp 210	S457		Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3330	Alticinae sp 210	S457		Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3331	Alticinae sp 210	S457		Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3332	Alticinae sp 035	S457	KR425036	Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3333	Alticinae sp 173	S457	KR425037	Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3334	Alticinae sp 052	S457	KR425038	Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3335	Galerucinae sp 045	S457	KR425346	Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3336	Galerucinae sp 045	S457		Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3337	Eumolpinae sp 040	S457	KR424843	Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3338	Eumolpinae sp 050	S457	KR424844	Bombuscaro	U6	08.11.2011	Sweep Netting
BT 3339	Alticinae sp 127	S458	KR425039	Bombuscaro	U6	08.11.2011	Beating
BT 3340	Eumolpinae sp 053	S458		Bombuscaro	U6	08.11.2011	Beating
BT 3341	Alticinae sp 181	S458	KR425040	Bombuscaro	U6	08.11.2011	Beating
BT 3342	Alticinae sp 169	S458	KR425041	Bombuscaro	U6	08.11.2011	Beating
BT 3343	Alticinae sp 076	S458	KR425042	Bombuscaro	U6	08.11.2011	Beating
BT 3344	Alticinae sp 076	S458		Bombuscaro	U6	08.11.2011	Beating
BT 3345	Alticinae sp 076	S458	KR425043	Bombuscaro	U6	08.11.2011	Beating
BT 3346	Cassidinae sp 016	S459	KR424786	Bombuscaro	U6	08.11.2011	Hand-Coll.(S)
BT 3347	Galerucinae sp 108	S459	KR425347	Bombuscaro	U6	08.11.2011	Hand-Coll.(S)
BT 3348	Cassidinae sp 018	S459	KR424787	Bombuscaro	U6	08.11.2011	Hand-Coll.(S)
BT 3349	Eumolpinae sp 052	S461		Cajanuma	U4	10.11.2011	Beating
BT 3350	Hispiinae sp 014	S462	KR424797	Cajanuma	U4	10.11.2011	Hand-Coll.(S)
BT 3351	Alticinae sp 255	S464	KR425044	Cajanuma	U5	10.11.2011	Beating
BT 3352	Alticinae sp 236	S464		Cajanuma	U5	10.11.2011	Beating
BT 3353	Eumolpinae sp 054	S464		Cajanuma	U5	10.11.2011	Beating
BT 3354	Alticinae sp 172	S466	KR425045	Cajanuma	U6	10.11.2011	Sweep Netting
BT 3355	Eumolpinae sp 054	S466		Cajanuma	U6	10.11.2011	Sweep Netting
BT 3356	Eumolpinae sp 054	S466		Cajanuma	U6	10.11.2011	Sweep Netting
BT 3357	Alticinae sp 057	S469	KR425046	Bombuscaro	U1	15.11.2011	Sweep Netting
BT 3358	Alticinae sp 057	S469		Bombuscaro	U1	15.11.2011	Sweep Netting
BT 3359	Galerucinae sp 074	S469	KR425348	Bombuscaro	U1	15.11.2011	Sweep Netting
BT 3360	Galerucinae sp 089	S469	KR424786	Bombuscaro	U1	15.11.2011	Sweep Netting
BT 3361	Alticinae sp 009	S469	KR425047	Bombuscaro	U1	15.11.2011	Sweep Netting
BT 3362	Eumolpinae sp 046	S469	KR424845	Bombuscaro	U1	15.11.2011	Sweep Netting
BT 3363	Alticinae sp 106	S469	KR425048	Bombuscaro	U1	15.11.2011	Sweep Netting
BT 3364	Eumolpinae sp 024	S470	KR424846	Bombuscaro	U1	15.11.2011	Beating
BT 3365	Eumolpinae sp 024	S470		Bombuscaro	U1	15.11.2011	Beating
BT 3366	Alticinae sp 152	S470	KR425049	Bombuscaro	U1	15.11.2011	Beating
BT 3367	Alticinae sp 051	S470	KR425050	Bombuscaro	U1	15.11.2011	Beating
BT 3368	Alticinae sp 051	S470		Bombuscaro	U1	15.11.2011	Beating
BT 3369	Alticinae sp 051	S472	KR425051	Bombuscaro	U2	15.11.2011	Sweep Netting
BT 3370	Alticinae sp 051	S472		Bombuscaro	U2	15.11.2011	Sweep Netting
BT 3371	Alticinae sp 051	S472		Bombuscaro	U2	15.11.2011	Sweep Netting
BT 3372	Alticinae sp 051	S472		Bombuscaro	U2	15.11.2011	Sweep Netting
BT 3373	Eumolpinae sp 064	S472	KR424847	Bombuscaro	U2	15.11.2011	Sweep Netting
BT 3374	Eumolpinae sp 072	S472		Bombuscaro	U2	15.11.2011	Sweep Netting
BT 3375	Eumolpinae sp 024	S473	KR424848	Bombuscaro	U2	15.11.2011	Beating
BT 3376	Eumolpinae sp 024	S473		Bombuscaro	U2	15.11.2011	Beating
BT 3377	Eumolpinae sp 024	S473		Bombuscaro	U2	15.11.2011	Beating
BT 3378	Eumolpinae sp 024	S473		Bombuscaro	U2	15.11.2011	Beating
BT 3379	Eumolpinae sp 062	S475	KR424849	Bombuscaro	U3	15.11.2011	Sweep Netting
BT 3380	Galerucinae sp 074	S475	KR425350	Bombuscaro	U3	15.11.2011	Sweep Netting
BT 3381	Galerucinae sp 074	S475		Bombuscaro	U3	15.11.2011	Sweep Netting
BT 3382	Galerucinae sp 055	S475	KR425351	Bombuscaro	U3	15.11.2011	Sweep Netting
BT 3383	Eumolpinae sp 024	S475	KR424850	Bombuscaro	U3	15.11.2011	Sweep Netting
BT 3384	Alticinae sp 152	S475	KR425052	Bombuscaro	U3	15.11.2011	Sweep Netting
BT 3385	Alticinae sp 051	S475	KR425053	Bombuscaro	U3	15.11.2011	Sweep Netting
BT 3386	Hispiinae sp 011	S476	KR424798	Bombuscaro	U3	15.11.2011	Beating
BT 3387	Eumolpinae sp 024	S476	KR424851	Bombuscaro	U3	15.11.2011	Beating
BT 3388	Eumolpinae sp 024	S476	KR424852	Bombuscaro	U3	15.11.2011	Beating
BT 3389	Eumolpinae sp 064	S476	KR424853	Bombuscaro	U3	15.11.2011	Beating
BT 3390	Alticinae sp 104	S478	KR425054	ECSF	U4	17.11.2011	Sweep Netting
BT 3391	Alticinae sp 052	S478	KR425055	ECSF	U4	17.11.2011	Sweep Netting
BT 3392	Alticinae sp 052	S478		ECSF	U4	17.11.2011	Sweep Netting
BT 3393	Alticinae sp 118	S478	KR425056	ECSF	U4	17.11.2011	Sweep Netting
BT 3394	Alticinae sp 118	S478		ECSF	U4	17.11.2011	Sweep Netting
BT 3395	Alticinae sp 118	S478		ECSF	U4	17.11.2011	Sweep Netting
BT 3396	Alticinae sp 118	S478		ECSF	U4	17.11.2011	Sweep Netting
BT 3398	Eumolpinae sp 039	S479		ECSF	U4	17.11.2011	Beating
BT 3399	Eumolpinae sp 039	S479		ECSF	U4	17.11.2011	Beating
BT 3400	Eumolpinae sp 039	S479		ECSF	U4	17.11.2011	Beating
BT 3401	Eumolpinae sp 039	S479		ECSF	U4	17.11.2011	Beating
BT 3402	Eumolpinae sp 039	S479		ECSF	U4	17.11.2011	Beating
BT 3403	Eumolpinae sp 030	S479	KR424854	ECSF	U4	17.11.2011	Beating
BT 3405	Eumolpinae sp 039	S480	KR424855	ECSF	U4	17.11.2011	Hand-Coll.(S)
BT 3406	Eumolpinae sp 039	S480		ECSF	U4	17.11.2011	Hand-Coll.(S)

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SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method	
BT 3407	Alticinae sp_118	S480	KR425057	ECSF	U4	17.11.2011	Hand-Coll.(S)
BT 3408	Alticinae sp_104	S481	KR425058	ECSF	U5	17.11.2011	Sweep Netting
BT 3409	Alticinae sp_113	S481	KR425059	ECSF	U5	17.11.2011	Sweep Netting
BT 3410	Alticinae sp_131	S481		ECSF	U5	17.11.2011	Sweep Netting
BT 3411	Alticinae sp_118	S481		ECSF	U5	17.11.2011	Sweep Netting
BT 3412	Eumolpinae sp_039	S482		ECSF	U5	17.11.2011	Beating
BT 3413	Eumolpinae sp_039	S482		ECSF	U5	17.11.2011	Beating
BT 3414	Eumolpinae sp_039	S482		ECSF	U5	17.11.2011	Beating
BT 3415	Eumolpinae sp_039	S482		ECSF	U5	17.11.2011	Beating
BT 3416	Alticinae sp_118	S484		ECSF	U6	17.11.2011	Sweep Netting
BT 3417	Alticinae sp_118	S484		ECSF	U6	17.11.2011	Sweep Netting
BT 3418	Alticinae sp_118	S484		ECSF	U6	17.11.2011	Sweep Netting
BT 3419	Alticinae sp_118	S484		ECSF	U6	17.11.2011	Sweep Netting
BT 3420	Alticinae sp_118	S484		ECSF	U6	17.11.2011	Sweep Netting
BT 3421	Alticinae sp_118	S484		ECSF	U6	17.11.2011	Sweep Netting
BT 3422	Alticinae sp_118	S484		ECSF	U6	17.11.2011	Sweep Netting
BT 3423	Eumolpinae sp_029	S485	KR424856	ECSF	U6	17.11.2011	Beating
BT 3424	Alticinae sp_118	S485		ECSF	U6	17.11.2011	Beating
BT 3425	Alticinae sp_104	S486	KR425060	ECSF	U6	17.11.2011	Hand-Coll.(S)
BT 3426	Galerucinae sp_105	S488	KR425352	Cajanuma	L4	21.11.2011	Beating
BT 3427	Alticinae sp_172	S490	KR425061	Cajanuma	L5	21.11.2011	Sweep Netting
BT 3428	Hispinae sp_025	S490	KR424799	Cajanuma	L5	21.11.2011	Sweep Netting
BT 3429	Galerucinae sp_066	S490	KR425353	Cajanuma	L5	21.11.2011	Sweep Netting
BT 3430	Alticinae sp_172	S491	KR425062	Cajanuma	L5	21.11.2011	Beating
BT 3431	Alticinae sp_248	S493	KR425063	Cajanuma	L6	21.11.2011	Sweep Netting
BT 3432	Alticinae sp_209	S493	KR425064	Cajanuma	L6	21.11.2011	Sweep Netting
BT 3433	Eumolpinae sp_071	S494	KR424857	Cajanuma	L6	21.11.2011	Beating
BT 3434	Alticinae sp_187	S494	KR425065	Cajanuma	L6	21.11.2011	Beating
BT 3435	Eumolpinae sp_019	S496	KR424858	ECSF	L1	23.11.2011	Sweep Netting
BT 3436	Galerucinae sp_061	S496	KR425354	ECSF	L1	23.11.2011	Sweep Netting
BT 3437	Galerucinae sp_061	S496		ECSF	L1	23.11.2011	Sweep Netting
BT 3438	Galerucinae sp_031	S496	KR425355	ECSF	L1	23.11.2011	Sweep Netting
BT 3439	Alticinae sp_018	S496		ECSF	L1	23.11.2011	Sweep Netting
BT 3440	Alticinae sp_117	S496		ECSF	L1	23.11.2011	Sweep Netting
BT 3441	Alticinae sp_174	S496	KR425066	ECSF	L1	23.11.2011	Sweep Netting
BT 3442	Alticinae sp_207	S496		ECSF	L1	23.11.2011	Sweep Netting
BT 3443	Alticinae sp_207	S496		ECSF	L1	23.11.2011	Sweep Netting
BT 3444	Alticinae sp_064	S496		ECSF	L1	23.11.2011	Sweep Netting
BT 3445	Alticinae sp_092	S496	KR425067	ECSF	L1	23.11.2011	Sweep Netting
BT 3446	Alticinae sp_083	S496		ECSF	L1	23.11.2011	Sweep Netting
BT 3447	Alticinae sp_150	S497	KR425068	ECSF	L1	23.11.2011	Beating
BT 3448	Alticinae sp_117	S497	KR425069	ECSF	L1	23.11.2011	Beating
BT 3449	Alticinae sp_002	S497	KR425070	ECSF	L1	23.11.2011	Beating
BT 3450	Alticinae sp_087	S497	KR425071	ECSF	L1	23.11.2011	Beating
BT 3451	Alticinae sp_087	S497		ECSF	L1	23.11.2011	Beating
BT 3452	Alticinae sp_065	S497	KR425072	ECSF	L1	23.11.2011	Beating
BT 3453	Galerucinae sp_061	S497	KR425356	ECSF	L1	23.11.2011	Beating
BT 3454	Galerucinae sp_061	S497		ECSF	L1	23.11.2011	Beating
BT 3455	Galerucinae sp_031	S499	KR425357	ECSF	L3	23.11.2011	Sweep Netting
BT 3456	Alticinae sp_266	S499		ECSF	L3	23.11.2011	Sweep Netting
BT 3457	Hispinae sp_003	S499	KR424800	ECSF	L3	23.11.2011	Sweep Netting
BT 3458	Eumolpinae sp_039	S499	KR424859	ECSF	L3	23.11.2011	Sweep Netting
BT 3459	Alticinae sp_150	S499	KR425074	ECSF	L3	23.11.2011	Sweep Netting
BT 3460	Alticinae sp_096	S499	KR425075	ECSF	L3	23.11.2011	Sweep Netting
BT 3461	Alticinae sp_162	S499		ECSF	L3	23.11.2011	Sweep Netting
BT 3462	Chrysomelinae sp_001	S499	KR424781	ECSF	L3	23.11.2011	Sweep Netting
BT 3463	Alticinae sp_109	S499	KR425076	ECSF	L3	23.11.2011	Sweep Netting
BT 3464	Alticinae sp_071	S499		ECSF	L3	23.11.2011	Sweep Netting
BT 3465	Alticinae sp_071	S499	KR425077	ECSF	L3	23.11.2011	Sweep Netting
BT 3466	Alticinae sp_190	S499		ECSF	L3	23.11.2011	Sweep Netting
BT 3467	Alticinae sp_190	S499		ECSF	L3	23.11.2011	Sweep Netting
BT 3468	Alticinae sp_197	S499	KR425078	ECSF	L3	23.11.2011	Sweep Netting
BT 3471	Galerucinae sp_002	S500	KR425358	ECSF	L3	23.11.2011	Beating
BT 3472	Galerucinae sp_002	S500		ECSF	L3	23.11.2011	Beating
BT 3473	Alticinae sp_109	S500	KR425079	ECSF	L3	23.11.2011	Beating
BT 3474	Alticinae sp_197	S500	KR425080	ECSF	L3	23.11.2011	Beating
BT 3475	Galerucinae sp_104	S501	KR425359	ECSF	L3	23.11.2011	Hand-Coll.(S)
BT 3476	Eumolpinae sp_065	S501		ECSF	L3	23.11.2011	Hand-Coll.(S)
BT 3477	Alticinae sp_198	S501	KR425081	ECSF	L3	23.11.2011	Hand-Coll.(S)
BT 3478	Alticinae sp_104	S502	KR425082	ECSF	U3	23.11.2011	Sweep Netting
BT 3479	Alticinae sp_065	S502	KR425083	ECSF	U3	23.11.2011	Sweep Netting
BT 3480	Galerucinae sp_061	S502	KR425360	ECSF	U3	23.11.2011	Sweep Netting
BT 3481	Alticinae sp_064	S502	KR425084	ECSF	U3	23.11.2011	Sweep Netting
BT 3482	Galerucinae sp_054	S502		ECSF	U3	23.11.2011	Sweep Netting
BT 3483	Alticinae sp_123	S502		ECSF	U3	23.11.2011	Sweep Netting
BT 3484	Alticinae sp_123	S502		ECSF	U3	23.11.2011	Sweep Netting
BT 3485	Alticinae sp_249	S502	KR425085	ECSF	U3	23.11.2011	Sweep Netting
BT 3486	Alticinae sp_198	S502	KR425086	ECSF	U3	23.11.2011	Sweep Netting
BT 3487	Alticinae sp_111	S502		ECSF	U3	23.11.2011	Sweep Netting
BT 3488	Alticinae sp_176	S503	KR425087	ECSF	U3	23.11.2011	Beating
BT 3489	Galerucinae sp_066	S503	KR425361	ECSF	U3	23.11.2011	Beating
BT 3490	Galerucinae sp_066	S503	KR425362	ECSF	U3	23.11.2011	Beating
BT 3491	Galerucinae sp_066	S503		ECSF	U3	23.11.2011	Beating
BT 3492	Galerucinae sp_066	S503		ECSF	U3	23.11.2011	Beating
BT 3493	Alticinae sp_249	S503	KR425088	ECSF	U3	23.11.2011	Beating
BT 3494	Alticinae sp_249	S503		ECSF	U3	23.11.2011	Beating
BT 3495	Cassidinae sp_012	S504	KR424788	ECSF	U3	23.11.2011	Hand-Coll.(S)
BT 3496	Alticinae sp_104	S505	KR425089	ECSF	U2	28.11.2011	Sweep Netting
BT 3497	Alticinae sp_061	S505	KR425090	ECSF	U2	28.11.2011	Sweep Netting
BT 3498	Alticinae sp_260	S505		ECSF	U2	28.11.2011	Sweep Netting
BT 3499	Alticinae sp_005	S506	KR425091	ECSF	U2	28.11.2011	Beating
BT 3500	Galerucinae sp_066	S506	KR425363	ECSF	U2	28.11.2011	Beating
BT 3501	Galerucinae sp_094	S506	KR425364	ECSF	U2	28.11.2011	Beating
BT 3502	Eumolpinae sp_017	S506		ECSF	U2	28.11.2011	Beating
BT 3503	Eumolpinae sp_039	S506	KR424860	ECSF	U2	28.11.2011	Beating
BT 3504	Alticinae sp_019	S506	KR425092	ECSF	U2	28.11.2011	Beating
BT 3505	Alticinae sp_109	S506	KR425093	ECSF	U2	28.11.2011	Beating
BT 3506	Alticinae sp_008	S506	KR425094	ECSF	U2	28.11.2011	Beating
BT 3507	Alticinae sp_254	S507		ECSF	U2	28.11.2011	Hand-Coll.(S)
BT 3508	Alticinae sp_118	S507		ECSF	U2	28.11.2011	Hand-Coll.(S)
BT 3509	Alticinae sp_073	S508			L1	30.11.2011	Sweep Netting
BT 3510	Eumolpinae sp_024	S508	KR424861	Bombuscaro	L1	30.11.2011	Sweep Netting
BT 3511	Galerucinae sp_079	S508	KR425365	Bombuscaro	L1	30.11.2011	Sweep Netting
BT 3512	Galerucinae sp_079	S508		Bombuscaro	L1	30.11.2011	Sweep Netting
BT 3513	Alticinae sp_051	S509	KR425095	Bombuscaro	L1	30.11.2011	Beating
BT 3514	Alticinae sp_051	S509		Bombuscaro	L1	30.11.2011	Beating
BT 3515	Alticinae sp_152	S509	KR425096	Bombuscaro	L1	30.11.2011	Beating
BT 3516	Eumolpinae sp_024	S509	KR424862	Bombuscaro	L1	30.11.2011	Beating
BT 3517	Galerucinae sp_048	S509	KR425366	Bombuscaro	L1	30.11.2011	Beating
BT 3518	Galerucinae sp_058	S510	KR425367	Bombuscaro	L1	30.11.2011	Hand-Coll.(S)
BT 3519	Alticinae sp_201	S511		Bombuscaro	L4	06.12.2011	Sweep Netting
BT 3520	Alticinae sp_181	S511	KR425097	Bombuscaro	L4	06.12.2011	Sweep Netting
BT 3521	Alticinae sp_051	S512	KR425098	Bombuscaro	L4	06.12.2011	Beating
BT 3522	Alticinae sp_181	S513	KR425099	Bombuscaro	L4	06.12.2011	Hand-Coll.(S)
BT 3528	Galerucinae sp_093	S517	KR425368	Bombuscaro	L6	06.12.2011	Sweep Netting

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SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method	
BT 3529	Galerucinae sp 091	S517	KR425369	Bombuscaro	L6	06.12.2011	Sweep Netting
BT 3530	Alticinae sp 153	S517	KR425100	Bombuscaro	L6	06.12.2011	Sweep Netting
BT 3531	Alticinae sp 166	S518	KR425101	Bombuscaro	L6	06.12.2011	Beating
BT 3532	Alticinae sp 149	S518	KR425102	Bombuscaro	L6	06.12.2011	Beating
BT 3533	Alticinae sp 247	S520	KR425103	Cajanuma	L1	08.12.2011	Sweep Netting
BT 3534	Alticinae sp 247	S520		Cajanuma	L1	08.12.2011	Sweep Netting
BT 3535	Alticinae sp 172	S520	KR425104	Cajanuma	L1	08.12.2011	Sweep Netting
BT 3536	Alticinae sp 172	S520	KR425105	Cajanuma	L1	08.12.2011	Sweep Netting
BT 3537	Alticinae sp 172	S520	KR425106	Cajanuma	L1	08.12.2011	Sweep Netting
BT 3538	Alticinae sp 244	S522	KR425107	Cajanuma	L1	08.12.2011	Hand-Coll.(S)
BT 3539	Alticinae sp 241	S523	KR425108	Cajanuma	L2	08.12.2011	Sweep Netting
BT 3540	Eumolpinae sp 058	S523	KR424863	Cajanuma	L2	08.12.2011	Sweep Netting
BT 3542	Eumolpinae sp 049	S524	KR424864	Cajanuma	L2	08.12.2011	Beating
BT 3543	Eumolpinae sp 069	S524	KR424865	Cajanuma	L2	08.12.2011	Beating
BT 3544	Chrysomelinae sp 004	S526	KR424782	Cajanuma	L3	08.12.2011	Sweep Netting
BT 3545	Alticinae sp 184	S535	KR425109	Cajanuma	U3	13.12.2011	Sweep Netting
BT 3546	Eumolpinae sp 054	S535		Cajanuma	U3	13.12.2011	Sweep Netting
BT 3547	Alticinae sp 214	S538	KR425110	Bombuscaro	L1	15.12.2011	Sweep Netting
BT 3548	Eumolpinae sp 063	S539	KR424866	Bombuscaro	L1	15.12.2011	Beating
BT 3549	Hispiniae sp 012	S540	KR424801	Bombuscaro	L1	15.12.2011	Hand-Coll.(S)
BT 3550	Galerucinae sp 093	S540	KR425370	Bombuscaro	L1	15.12.2011	Hand-Coll.(S)
BT 3551	Galerucinae sp 093	S541	KR425371	Bombuscaro	L2	15.12.2011	Sweep Netting
BT 3552	Alticinae sp 026	S541	KR425111	Bombuscaro	L2	15.12.2011	Sweep Netting
BT 3553	Alticinae sp 166	S541		Bombuscaro	L2	15.12.2011	Sweep Netting
BT 3554	Alticinae sp 073	S543		Bombuscaro	L2	15.12.2011	Hand-Coll.(S)
BT 3555	Alticinae sp 181	S544	KR425112	Bombuscaro	L3	15.12.2011	Sweep Netting
BT 3556	Eumolpinae sp 050	S544	KR424867	Bombuscaro	L3	15.12.2011	Sweep Netting
BT 3557	Eumolpinae sp 070	S545	KR424868	Bombuscaro	L3	15.12.2011	Beating
BT 3558	Hispiniae sp 021	S546	KR424802	Bombuscaro	L3	15.12.2011	Hand-Coll.(S)
BT 3559	Alticinae sp 233	S547	KR425113	Cajanuma	U4	19.12.2011	Sweep Netting
BT 3560	Alticinae sp 188	S547	KR425114	Cajanuma	U5	19.12.2011	Beating
BT 3561	Alticinae sp 056	S553		Cajanuma	U6	19.12.2011	Sweep Netting
BT 3562	Eumolpinae sp 057	S554	KR424869	Cajanuma	U6	19.12.2011	Beating
BT 3563	Hispiniae sp 014	S554	KR424803	Cajanuma	U6	19.12.2011	Beating
BT 3564	Alticinae sp 127	S556	KR425115	Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3565	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3566	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3567	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3568	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3569	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3570	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3571	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3572	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3573	Alticinae sp 127	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3574	Alticinae sp 227	S556	KR425116	Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3575	Alticinae sp 227	S556	KR425117	Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3576	Alticinae sp 196	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3577	Alticinae sp 196	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3578	Alticinae sp 196	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3579	Alticinae sp 196	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3580	Alticinae sp 196	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3581	Alticinae sp 179	S556	KR425118	Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3582	Galerucinae sp 106	S556		Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3583	Alticinae sp 135	S556	KR425119	Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3584	Alticinae sp 167	S556	KR425120	Bombuscaro	U4	21.12.2011	Sweep Netting
BT 3585	Alticinae sp 127	S557	KR425121	Bombuscaro	U4	21.12.2011	Beating
BT 3586	Alticinae sp 127	S557		Bombuscaro	U4	21.12.2011	Beating
BT 3587	Alticinae sp 127	S557		Bombuscaro	U4	21.12.2011	Beating
BT 3588	Alticinae sp 127	S557		Bombuscaro	U4	21.12.2011	Beating
BT 3589	Alticinae sp 127	S557		Bombuscaro	U4	21.12.2011	Beating
BT 3590	Alticinae sp 127	S557		Bombuscaro	U4	21.12.2011	Beating
BT 3591	Alticinae sp 196	S557	KR425122	Bombuscaro	U4	21.12.2011	Beating
BT 3592	Alticinae sp 196	S557		Bombuscaro	U4	21.12.2011	Beating
BT 3593	Alticinae sp 076	S557	KR425123	Bombuscaro	U4	21.12.2011	Beating
BT 3594	Alticinae sp 181	S557	KR425124	Bombuscaro	U4	21.12.2011	Beating
BT 3596	Alticinae sp 111	S558	KR425125	Bombuscaro	U4	21.12.2011	Hand-Coll.(S)
BT 3597	Alticinae sp 196	S558	KR425126	Bombuscaro	U4	21.12.2011	Hand-Coll.(S)
BT 3598	Alticinae sp 196	S559		Bombuscaro	U5	21.12.2011	Sweep Netting
BT 3599	Alticinae sp 196	S559		Bombuscaro	U5	21.12.2011	Sweep Netting
BT 3600	Alticinae sp 253	S559		Bombuscaro	U5	21.12.2011	Sweep Netting
BT 3601	Alticinae sp 127	S559	KR425127	Bombuscaro	U5	21.12.2011	Sweep Netting
BT 3602	Alticinae sp 138	S559		Bombuscaro	U5	21.12.2011	Sweep Netting
BT 3603	Alticinae sp 182	S560		Bombuscaro	U5	21.12.2011	Beating
BT 3604	Alticinae sp 076	S560	KR425128	Bombuscaro	U5	21.12.2011	Beating
BT 3605	Alticinae sp 076	S560		Bombuscaro	U5	21.12.2011	Beating
BT 3606	Alticinae sp 127	S560		Bombuscaro	U5	21.12.2011	Beating
BT 3607	Alticinae sp 138	S560		Bombuscaro	U5	21.12.2011	Beating
BT 3608	Alticinae sp 253	S560		Bombuscaro	U5	21.12.2011	Beating
BT 3609	Eumolpinae sp 047	S560	KR424870	Bombuscaro	U5	21.12.2011	Beating
BT 3610	Alticinae sp 076	S561	KR425129	Bombuscaro	U5	21.12.2011	Hand-Coll.(S)
BT 3611	Alticinae sp 181	S562	KR425130	Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3612	Eumolpinae sp 059	S562	KR424871	Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3613	Criocerinae sp 007	S562	KR425415	Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3614	Eumolpinae sp 059	S562	KR424872	Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3615	Galerucinae sp 029	S562	KR425372	Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3616	Galerucinae sp 029	S562	KR425373	Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3617	Alticinae sp 127	S562	KR425131	Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3618	Alticinae sp 127	S562		Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3619	Alticinae sp 127	S562		Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3620	Alticinae sp 127	S562		Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3621	Alticinae sp 127	S562		Bombuscaro	U6	21.12.2011	Sweep Netting
BT 3622	Alticinae sp 254	S563	KR425132	Bombuscaro	U6	21.12.2011	Beating
BT 3623	Alticinae sp 253	S563	KR425133	Bombuscaro	U6	21.12.2011	Beating
BT 3624	Alticinae sp 127	S563	KR425134	Bombuscaro	U6	21.12.2011	Beating
BT 3625	Alticinae sp 051	S563	KR425135	Bombuscaro	U6	21.12.2011	Beating
BT 3626	Criocerinae sp 010	S563	KR425416	Bombuscaro	U6	21.12.2011	Beating
BT 3627	Eumolpinae sp 039	S565		ECSF	U1	26.12.2011	Sweep Netting
BT 3628	Alticinae sp 104	S565		ECSF	U1	26.12.2011	Sweep Netting
BT 3630	Alticinae sp 117	S566		ECSF	U1	26.12.2011	Beating
BT 3633	Alticinae sp 109	S566		ECSF	U1	26.12.2011	Beating
BT 3651	Galerucinae sp 041	S572		ECSF	L4	03.01.2012	Beating
BT 3652	Alticinae sp 086	S572		ECSF	L4	03.01.2012	Beating
BT 3653	Alticinae sp 065	S572		ECSF	L4	03.01.2012	Beating
BT 3656	Alticinae sp 115	S572		ECSF	L4	03.01.2012	Beating
BT 3658	Alticinae sp 198	S573		ECSF	L4	03.01.2012	Hand-Coll.(S)
BT 3659	Alticinae sp 237	S573		ECSF	L4	03.01.2012	Hand-Coll.(S)
BT 3660	Alticinae sp 092	S573		ECSF	L4	03.01.2012	Hand-Coll.(S)
BT 3685	Alticinae sp 018	S577	KR425136	ECSF	L6	03.01.2012	Sweep Netting
BT 3686	Alticinae sp 177	S577	KR425137	ECSF	L6	03.01.2012	Sweep Netting
BT 3687	Alticinae sp 115	S577	KR425138	ECSF	L6	03.01.2012	Sweep Netting
BT 3688	Alticinae sp 087	S578	KR425139	ECSF	L6	03.01.2012	Beating
BT 3689	Eumolpinae sp 040	S578	KR424873	ECSF	L6	03.01.2012	Beating
BT 3690	Alticinae sp 115	S578	KR425140	ECSF	L6	03.01.2012	Beating
BT 3691	Alticinae sp 147	S578	KR425141	ECSF	L6	03.01.2012	Beating
BT 3692	Alticinae sp 147	S578		ECSF	L6	03.01.2012	Beating

Continued on next page(s)



Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method
BT 3693	Alticinae sp 123	S578	ECSSF	L6	03.01.2012	Beating
BT 3694	Alticinae sp 197	S578	ECSSF	L6	03.01.2012	Beating
BT 3698	Galerucinae sp 089	S586	Bombuscaro	U3	05.01.2012	Sweep Netting
BT 3699	Eumolpinae sp 063	S586	Bombuscaro	U3	05.01.2012	Sweep Netting
BT 3700	Eumolpinae sp 024	S587	Bombuscaro	U3	05.01.2012	Beating
BT 3701	Alticinae sp 244	S590	Cajanuma	L4	09.01.2012	Beating
BT 3702	Alticinae sp 206	S591	Cajanuma	L4	09.01.2012	Hand-Coll.(S)
BT 3703	Alticinae sp 206	S591	Cajanuma	L4	09.01.2012	Hand-Coll.(S)
BT 3704	Alticinae sp 172	S597	Cajanuma	L6	09.01.2012	Hand-Coll.(S)
BT 3705	Alticinae sp 085	S598	Bombuscaro	L4	11.01.2012	Sweep Netting
BT 3706	Alticinae sp 182	S598	Bombuscaro	L4	11.01.2012	Sweep Netting
BT 3707	Alticinae sp 182	S598	Bombuscaro	L4	11.01.2012	Sweep Netting
BT 3708	Eumolpinae sp 059	S598	Bombuscaro	L4	11.01.2012	Sweep Netting
BT 3710	Alticinae sp 196	S598	Bombuscaro	L4	11.01.2012	Sweep Netting
BT 3711	Alticinae sp 076	S598	Bombuscaro	L4	11.01.2012	Sweep Netting
BT 3716	Alticinae sp 127	S599	Bombuscaro	L4	11.01.2012	Beating
BT 3719	Alticinae sp 093	S599	Bombuscaro	L4	11.01.2012	Beating
BT 3721	Alticinae sp 076	S599	Bombuscaro	L4	11.01.2012	Beating
BT 3723	Eumolpinae sp 059	S599	Bombuscaro	L4	11.01.2012	Beating
BT 3724	Alticinae sp 127	S600	Bombuscaro	L4	11.01.2012	Hand-Coll.(S)
BT 3725	Alticinae sp 138	S600	Bombuscaro	L4	11.01.2012	Hand-Coll.(S)
BT 3726	Alticinae sp 227	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3727	Alticinae sp 253	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3728	Alticinae sp 099	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3729	Alticinae sp 104	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3730	Alticinae sp 095	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3731	Alticinae sp 127	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3732	Alticinae sp 127	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3733	Alticinae sp 127	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3734	Alticinae sp 127	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3735	Alticinae sp 196	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3736	Alticinae sp 196	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3737	Alticinae sp 196	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3738	Alticinae sp 196	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3739	Galerucinae sp 085	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3740	Eumolpinae sp 047	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3741	Alticinae sp 138	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3750	Alticinae sp 076	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3751	Alticinae sp 076	S601	Bombuscaro	L5	11.01.2012	Sweep Netting
BT 3752	Eumolpinae sp 038	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3753	Galerucinae sp 093	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3754	Alticinae sp 076	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3755	Alticinae sp 076	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3756	Alticinae sp 127	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3757	Alticinae sp 127	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3758	Alticinae sp 127	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3759	Alticinae sp 127	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3760	Alticinae sp 127	S602	Bombuscaro	L5	11.01.2012	Beating
BT 3761	Alticinae sp 196	S603	Bombuscaro	L5	11.01.2012	Hand-Coll.(S)
BT 3762	Alticinae sp 127	S603	Bombuscaro	L5	11.01.2012	Hand-Coll.(S)
BT 3763	Alticinae sp 127	S603	Bombuscaro	L5	11.01.2012	Hand-Coll.(S)
BT 3765	Alticinae sp 138	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3767	Alticinae sp 226	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3768	Criocerinae sp 008	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3771	Alticinae sp 169	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3773	Alticinae sp 127	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3790	Alticinae sp 196	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3791	Alticinae sp 252	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3792	Alticinae sp 240	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3793	Alticinae sp 204	S604	Bombuscaro	L6	11.01.2012	Sweep Netting
BT 3795	Hispinae sp 019	S605	Bombuscaro	L6	11.01.2012	Beating
BT 3796	Galerucinae sp 007	S605	Bombuscaro	L6	11.01.2012	Beating
BT 3807	Alticinae sp 185	S607	Cajanuma	L1	16.01.2012	Sweep Netting
BT 3808	Alticinae sp 230	S607	Cajanuma	L1	16.01.2012	Sweep Netting
BT 3809	Alticinae sp 234	S607	Cajanuma	L1	16.01.2012	Sweep Netting
BT 3810	Alticinae sp 217	S608	Cajanuma	L1	16.01.2012	Beating
BT 3811	Hispinae sp 017	S610	Cajanuma	L2	16.01.2012	Sweep Netting
BT 3812	Alticinae sp 118	S610	Cajanuma	L2	16.01.2012	Sweep Netting
BT 3813	Alticinae sp 109	S611	Cajanuma	L2	16.01.2012	Beating
BT 3814	Alticinae sp 118	S611	Cajanuma	L2	16.01.2012	Beating
BT 3815	Alticinae sp 118	S611	Cajanuma	L2	16.01.2012	Beating
BT 3816	Alticinae sp 118	S611	Cajanuma	L2	16.01.2012	Beating
BT 3817	Alticinae sp 118	S611	Cajanuma	L2	16.01.2012	Beating
BT 3818	Alticinae sp 118	S611	Cajanuma	L2	16.01.2012	Beating
BT 3819	Alticinae sp 118	S611	Cajanuma	L2	16.01.2012	Beating
BT 3820	Alticinae sp 216	S613	Cajanuma	L3	16.01.2012	Sweep Netting
BT 3824	Galerucinae sp 013	S619	Bombuscaro	L2	18.01.2012	Sweep Netting
BT 3825	Alticinae sp 050	S619	Bombuscaro	L2	18.01.2012	Sweep Netting
BT 3826	Alticinae sp 201	S619	Bombuscaro	L2	18.01.2012	Sweep Netting
BT 3827	Galerucinae sp 090	S620	Bombuscaro	L2	18.01.2012	Beating
BT 3828	Alticinae sp 211	S620	Bombuscaro	L2	18.01.2012	Beating
BT 3829	Alticinae sp 181	S620	Bombuscaro	L2	18.01.2012	Beating
BT 3830	Alticinae sp 180	S623	Bombuscaro	L3	18.01.2012	Beating
BT 3832	Eumolpinae sp 050	S624	Bombuscaro	L3	18.01.2012	Hand-Coll.(S)
BT 3833	Alticinae sp 251	S631	Cajanuma	U3	23.01.2012	Sweep Netting
BT 3834	Alticinae sp 244	S631	Cajanuma	U3	23.01.2012	Sweep Netting
BT 3845	Galerucinae sp 007	S636	ECSSF	U4	25.01.2012	Hand-Coll.(S)
BT 3846	Alticinae sp 118	S637	ECSSF	U5	25.01.2012	Sweep Netting
BT 3847	Alticinae sp 118	S637	ECSSF	U5	25.01.2012	Sweep Netting
BT 3848	Eumolpinae sp 039	S637	ECSSF	U5	25.01.2012	Sweep Netting
BT 3849	Alticinae sp 175	S637	ECSSF	U5	25.01.2012	Sweep Netting
BT 3850	Alticinae sp 140	S637	ECSSF	U5	25.01.2012	Sweep Netting
BT 3851	Alticinae sp 118	S638	ECSSF	U5	25.01.2012	Beating
BT 3852	Alticinae sp 118	S638	ECSSF	U5	25.01.2012	Beating
BT 3853	Eumolpinae sp 039	S638	ECSSF	U5	25.01.2012	Beating
BT 3854	Eumolpinae sp 039	S638	ECSSF	U5	25.01.2012	Beating
BT 3855	Eumolpinae sp 039	S638	ECSSF	U5	25.01.2012	Beating
BT 3856	Alticinae sp 118	S639	ECSSF	U5	25.01.2012	Hand-Coll.(S)
BT 3857	Alticinae sp 118	S639	ECSSF	U5	25.01.2012	Hand-Coll.(S)
BT 3858	Alticinae sp 118	S639	ECSSF	U5	25.01.2012	Hand-Coll.(S)
BT 3859	Alticinae sp 118	S639	ECSSF	U5	25.01.2012	Hand-Coll.(S)
BT 3860	Eumolpinae sp 030	S639	ECSSF	U5	25.01.2012	Hand-Coll.(S)
BT 3861	Cassidinae sp 015	S639	ECSSF	U5	25.01.2012	Hand-Coll.(S)
BT 3869	Cassidinae sp 012	S646	ECSSF	L5	27.01.2012	Sweep Netting
BT 3870	Eumolpinae sp 002	S646	ECSSF	L5	27.01.2012	Sweep Netting
BT 3871	Alticinae sp 083	S646	ECSSF	L5	27.01.2012	Sweep Netting
BT 3872	Eumolpinae sp 040	S647	ECSSF	L5	27.01.2012	Beating
BT 3873	Cassidinae sp 020	S648	ECSSF	L5	27.01.2012	Hand-Coll.(S)
BT 3874	Eumolpinae sp 042	S648	ECSSF	L5	27.01.2012	Hand-Coll.(S)
BT 3885	Alticinae sp 007	S653	ECSSF	L6	27.01.2012	Beating
BT 3891	Galerucinae sp 090	S658	Bombuscaro	U1	29.01.2012	Sweep Netting
BT 3892	Galerucinae sp 090	S658	Bombuscaro	U1	29.01.2012	Sweep Netting
BT 3893	Alticinae sp 164	S658	Bombuscaro	U1	29.01.2012	Sweep Netting
BT 3894	Alticinae sp 200	S661	Bombuscaro	L1	29.01.2012	Sweep Netting
BT 3895	Galerucinae sp 079	S662	Bombuscaro	L1	29.01.2012	Beating

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SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method
BT 3896	S663	KR424805	Bombuscaro	L1	29.01.2012	Hand-Coll.(S)
BT 3915	S670	KR425380	ECSF	U4	31.01.2012	Sweep Netting
BT 3916	S670		ECSF	U4	31.01.2012	Sweep Netting
BT 3917	S671	KR425177	ECSF	U4	31.01.2012	Beating
BT 3918	S671	KR425178	ECSF	U4	31.01.2012	Beating
BT 3919	S671		ECSF	U4	31.01.2012	Beating
BT 3920	S671	KR425179	ECSF	U4	31.01.2012	Beating
BT 3921	S671	KR424883	ECSF	U4	31.01.2012	Beating
BT 3922	S671	KR425180	ECSF	U4	31.01.2012	Beating
BT 3923	S672		ECSF	U4	31.01.2012	Hand-Coll.(S)
BT 3947	S679	KR424884	ECSF	L2	31.01.2012	Sweep Netting
BT 3948	S679	KR425181	ECSF	L2	31.01.2012	Sweep Netting
BT 3949	S679	KR425182	ECSF	L2	31.01.2012	Sweep Netting
BT 3950	S679	KR425381	ECSF	L2	31.01.2012	Sweep Netting
BT 3951	S679	KR425183	ECSF	L2	31.01.2012	Sweep Netting
BT 3952	S679	KR425184	ECSF	L2	31.01.2012	Sweep Netting
BT 3953	S679	KR425185	ECSF	L2	31.01.2012	Sweep Netting
BT 3954	S679		ECSF	L2	31.01.2012	Sweep Netting
BT 3955	S679		ECSF	L2	31.01.2012	Sweep Netting
BT 3957	S680	KR425382	ECSF	L2	31.01.2012	Beating
BT 3958	S681	KR424885	ECSF	L2	31.01.2012	Hand-Coll.(S)
BT 3959	S681	KR425383	ECSF	L2	31.01.2012	Hand-Coll.(S)
BT 3960	S681	KR425186	ECSF	L2	31.01.2012	Hand-Coll.(S)
BT 3961	S681	KR424792	ECSF	L2	31.01.2012	Hand-Coll.(S)
BT 3962	S682		ECSF	U6	31.01.2012	Sweep Netting
BT 3963	S682		ECSF	U6	31.01.2012	Sweep Netting
BT 3964	S683	KR425187	ECSF	U6	31.01.2012	Beating
BT 3965	S683	KR425188	ECSF	U6	31.01.2012	Beating
BT 3966	S683		ECSF	U6	31.01.2012	Beating
BT 3967	S683		ECSF	U6	31.01.2012	Beating
BT 3968	S683		ECSF	U6	31.01.2012	Beating
BT 3969	S683	KR425189	ECSF	U6	31.01.2012	Beating
BT 3970	S683	KR425190	ECSF	U6	31.01.2012	Beating
BT 3971	S684	KR425384	ECSF	U6	31.01.2012	Hand-Coll.(S)
BT 3972	S686		Cajanuma	L6	03.02.2012	Beating
BT 3973	S694	KR425191	Cajanuma	U2	03.02.2012	Sweep Netting
BT 3974	S695	KR425192	Cajanuma	U2	03.02.2012	Beating
BT 3975	S697		Cajanuma	L4	03.02.2012	Sweep Netting
BT 3976	S700	KR424886	Bombuscaro	L4	06.02.2012	Sweep Netting
BT 3977	S701	KR424806	Bombuscaro	L4	06.02.2012	Beating
BT 3986	S705		Bombuscaro	U4	06.02.2012	Hand-Coll.(S)
BT 3988	S706	KR424887	Bombuscaro	L5	06.02.2012	Sweep Netting
BT 3989	S706	KR425194	Bombuscaro	L5	06.02.2012	Sweep Netting
BT 3990	S706	KR425195	Bombuscaro	L5	06.02.2012	Sweep Netting
BT 3994	S713		Bombuscaro	U6	06.02.2012	Beating
BT 3995	S715	KR425385	Bombuscaro	U6	06.02.2012	Sweep Netting
BT 3996	S715	KR425196	Bombuscaro	U6	06.02.2012	Sweep Netting
BT 3997	S716		Bombuscaro	U6	06.02.2012	Beating
BT 3998	S716	KR425386	Bombuscaro	U6	06.02.2012	Beating
BT 3999	S717	KR425197	Bombuscaro	U6	06.02.2012	Hand-Coll.(S)
BT 4000	S718	KR425198	ECSF	U2	09.02.2012	Sweep Netting
BT 4001	S718		ECSF	U2	09.02.2012	Sweep Netting
BT 4002	S719	KR425387	ECSF	U2	09.02.2012	Beating
BT 4003	S720	KR425388	ECSF	U2	09.02.2012	Hand-Coll.(S)
BT 4004	S720		ECSF	U2	09.02.2012	Hand-Coll.(S)
BT 4005	S720	KR425389	ECSF	U2	09.02.2012	Hand-Coll.(S)
BT 4006	S722	KR425199	ECSF	L4	09.02.2012	Beating
BT 4007	S722	KR425200	ECSF	L4	09.02.2012	Beating
BT 4008	S722	KR425390	ECSF	L4	09.02.2012	Beating
BT 4009	S722	KR425391	ECSF	L4	09.02.2012	Beating
BT 4010	S722	KR424888	ECSF	L4	09.02.2012	Beating
BT 4011	S722		ECSF	L4	09.02.2012	Beating
BT 4012	S722		ECSF	L4	09.02.2012	Beating
BT 4013	S723	KR425391	ECSF	L4	09.02.2012	Hand-Coll.(S)
BT 4014	S723	KR424793	ECSF	L4	09.02.2012	Hand-Coll.(S)
BT 4015	S723	KR425392	ECSF	L4	09.02.2012	Hand-Coll.(S)
BT 4016	S723	KR425393	ECSF	L4	09.02.2012	Hand-Coll.(S)
BT 4017	S724	KR425202	ECSF	U1	09.02.2012	Sweep Netting
BT 4018	S724		ECSF	U1	09.02.2012	Sweep Netting
BT 4019	S724	KR424889	ECSF	U1	09.02.2012	Sweep Netting
BT 4020	S724		ECSF	U1	09.02.2012	Sweep Netting
BT 4021	S725	KR425203	ECSF	U1	09.02.2012	Beating
BT 4022	S726	KR424794	ECSF	U1	09.02.2012	Hand-Coll.(S)
BT 4026	S728		ECSF	L5	09.02.2012	Beating
BT 4031	S730	KR425204	ECSF	U1	09.02.2012	Sweep Netting
BT 4032	S731	KR424807	ECSF	L6	09.02.2012	Beating
BT 4033	S731		ECSF	L6	09.02.2012	Beating
BT 4035	S732	KR424808	ECSF	L6	09.02.2012	Hand-Coll.(S)
BT 4036	S741	KR425394	Bombuscaro	U3	11.02.2012	Hand-Coll.(S)
BT 4047	S748	KR425205	Bombuscaro	L3	11.02.2012	Sweep Netting
BT 4048	S751	KR425417	ECSF	L1	12.02.2012	Sweep Netting
BT 4050	S752	KR425206	ECSF	L1	12.02.2012	Sweep Netting
BT 4051	S752	KR424890	ECSF	L1	12.02.2012	Beating
BT 4052	S752	KR424891	ECSF	L1	12.02.2012	Beating
BT 4052	S752	KR425207	ECSF	L1	12.02.2012	Beating
BT 4053	S752		ECSF	L1	12.02.2012	Beating
BT 4054	S752	KR425208	ECSF	L1	12.02.2012	Beating
BT 4055	S752		ECSF	L1	12.02.2012	Beating
BT 4056	S752	KR425209	ECSF	L1	12.02.2012	Beating
BT 4057	S752		ECSF	L1	12.02.2012	Beating
BT 4058	S752		ECSF	L1	12.02.2012	Beating
BT 4059	S753	KR425395	ECSF	L1	12.02.2012	Hand-Coll.(S)
BT 4060	S753		ECSF	L1	12.02.2012	Hand-Coll.(S)
BT 4061	S754	KR425210	ECSF	L2	12.02.2012	Sweep Netting
BT 4062	S755	KR424892	ECSF	L2	12.02.2012	Beating
BT 4063	S755	KR425211	ECSF	L2	12.02.2012	Beating
BT 4064	S755		ECSF	L2	12.02.2012	Beating
BT 4065	S755	KR425212	ECSF	L2	12.02.2012	Beating
BT 4066	S756	KR425213	ECSF	L2	12.02.2012	Hand-Coll.(S)
BT 4067	S756		ECSF	L2	12.02.2012	Hand-Coll.(S)
BT 4182	N149		ECSF		12.09.2011	Hand-Coll.(N)
BT 4186	N151		ECSF		19.09.2011	Hand-Coll.(N)
BT 4194	N155		Bombuscaro		04.10.2011	Hand-Coll.(N)
BT 4195	N155		Bombuscaro		04.10.2011	Hand-Coll.(N)
BT 4198	N155		Bombuscaro		04.10.2011	Hand-Coll.(N)
BT 4207	N155		Bombuscaro		04.10.2011	Hand-Coll.(N)
BT 4208	N155		Bombuscaro		04.10.2011	Hand-Coll.(N)
BT 4209	N155		Bombuscaro		04.10.2011	Hand-Coll.(N)
BT 4215	N157		Cajanuma		11.10.2011	Hand-Coll.(N)
BT 4216	N157		Cajanuma		11.10.2011	Hand-Coll.(N)
BT 4217	N157		Cajanuma		11.10.2011	Hand-Coll.(N)
BT 4241	N162		ECSF		13.10.2011	Hand-Coll.(N)
BT 4252	N163		ECSF		18.10.2011	Hand-Coll.(N)
BT 4253	N163		ECSF		18.10.2011	Hand-Coll.(N)

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Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/ Plot	Sampling Date	Sampling Method	
BT 4294	Hispinae_sp_020	N172	Bombuscaro		08.11.2011	Hand-Coll.(N)	
BT 4295	Galerucinae_sp_029	N172	Bombuscaro		08.11.2011	Hand-Coll.(N)	
BT 4306	Cassidinae_sp_019	N172	Bombuscaro		08.11.2011	Hand-Coll.(N)	
BT 4321	Galerucinae_sp_017	N175	Bombuscaro		17.02.2011	Hand-Coll.(N)	
BT 4344	Galerucinae_sp_011	N179	ECSF	L6	28.11.2011	Malaise Trap	
BT 4350	Alticinae_sp_218	N179	ECSF	L6	28.11.2011	Malaise Trap	
BT 4351	Galerucinae_sp_017	N179	ECSF	L6	28.11.2011	Malaise Trap	
BT 4354	Alticinae_sp_163	N179	ECSF	L6	28.11.2011	Malaise Trap	
BT 4402	Alticinae_sp_222	N184	Bombuscaro	L2	30.11.2011	Flight-Intercept.	
BT 4421	Alticinae_sp_223	N193	Bombuscaro		21.12.2011	Hand-Coll.(N)	
BT 4434	Alticinae_sp_163	N194	ECSF	L6	26.12.2011	Malaise Trap	
BT 4468	Galerucinae_sp_007	N195	ECSF	U1	26.12.2011	Malaise Trap	
BT 4477	Alticinae_sp_155	N201	Bombuscaro	U2	28.12.2011	Malaise Trap	
BT 4500	Galerucinae_sp_097	N207	Bombuscaro		11.01.2012	Hand-Coll.(N)	
BT 4509	Cassidinae_sp_018	N207	Bombuscaro		11.01.2012	Hand-Coll.(N)	
BT 4510	Hispinae_sp_026	N208	Cajanuma		16.01.2012	Hand-Coll.(N)	
BT 4511	Alticinae_sp_229	N208	Cajanuma		16.01.2012	Hand-Coll.(N)	
BT 4550	Alticinae_sp_220	N216	ECSF	L6	27.01.2012	Malaise Trap	
BT 4572	Alticinae_sp_219	N223	ECSF	U5	27.02.2012	Light Trap	
BT 4581	Galerucinae_sp_005	N223	ECSF	U5	30.01.2012	Light Trap	
BT 4605	Galerucinae_sp_110	N225	ECSF	U6	31.01.2012	Hand-Coll.(N)	
BT 4684	Cassidinae_sp_001	N234	ECSF	U6	09.02.2012	Hand-Coll.(N)	
BT 4687	Galerucinae_sp_036	N234	ECSF	U6	09.02.2012	Hand-Coll.(N)	
BT 4732	Alticinae_sp_095	S760	KR425214	Cajanuma	U4	16.02.2012	Sweep Netting
BT 4733	Eumolpinae_sp_068	S760	KR424893	Cajanuma	U4	16.02.2012	Sweep Netting
BT 4734	Alticinae_sp_172	S763	KR425215	Cajanuma	U5	16.02.2012	Sweep Netting
BT 4735	Alticinae_sp_073	S763	KR425216	Cajanuma	U5	16.02.2012	Sweep Netting
BT 4736	Alticinae_sp_172	S766	KR425217	Cajanuma	U6	16.02.2012	Sweep Netting
BT 4737	Alticinae_sp_172	S767	KR425218	Cajanuma	U6	16.02.2012	Beating
BT 4738	Alticinae_sp_172	S767		Cajanuma	U6	16.02.2012	Beating
BT 4739	Eumolpinae_sp_024	S769	KR424894	Bombuscaro	U4	27.02.2012	Sweep Netting
BT 4740	Eumolpinae_sp_024	S769	KR424895	Bombuscaro	U4	27.02.2012	Sweep Netting
BT 4741	Alticinae_sp_073	S772		Bombuscaro	U5	27.02.2012	Sweep Netting
BT 4742	Eumolpinae_sp_024	S772	KR424896	Bombuscaro	U4	27.02.2012	Sweep Netting
BT 4743	Alticinae_sp_181	S775	KR425219	Bombuscaro	U6	27.02.2012	Sweep Netting
BT 4744	Eumolpinae_sp_056	S781	KR424897	Bombuscaro	U2	29.02.2012	Sweep Netting
BT 4745	Galerucinae_sp_080	S781		Bombuscaro	U2	29.02.2012	Sweep Netting
BT 4746	Eumolpinae_sp_024	S781	KR424898	Bombuscaro	U2	29.02.2012	Sweep Netting
BT 4747	Galerucinae_sp_074	S781	KR425396	Bombuscaro	U2	29.02.2012	Sweep Netting
BT 4748	Eumolpinae_sp_024	S785	KR424899	Bombuscaro	U2	29.02.2012	Beating
BT 4749	Galerucinae_sp_074	S785	KR425397	Bombuscaro	U3	29.02.2012	Beating
BT 4750	Eumolpinae_sp_024	S785	KR424900	Bombuscaro	U3	29.02.2012	Beating
BT 4751	Alticinae_sp_253	S787	KR425220	Bombuscaro	L4	06.03.2012	Sweep Netting
BT 4753	Eumolpinae_sp_038	S788		Bombuscaro	L4	06.03.2012	Beating
BT 4754	Alticinae_sp_195	S788	KR425221	Bombuscaro	L4	06.03.2012	Beating
BT 4772	Alticinae_sp_009	S793	KR425222	Bombuscaro	L6	06.03.2012	Sweep Netting
BT 4773	Alticinae_sp_127	S793	KR425223	Bombuscaro	L6	06.03.2012	Sweep Netting
BT 4774	Alticinae_sp_050	S794	KR425224	Bombuscaro	L6	06.03.2012	Beating
BT 4775	Alticinae_sp_211	S795	KR425225	Bombuscaro	L6	06.03.2012	Hand-Coll.(S)
BT 4776	Galerucinae_sp_029	S798	KR425398	Bombuscaro	L1	08.03.2012	Hand-Coll.(S)
BT 4777	Alticinae_sp_183	S799		Bombuscaro	L2	08.03.2012	Sweep Netting
BT 4778	Alticinae_sp_181	S799	KR425226	Bombuscaro	L2	08.03.2012	Sweep Netting
BT 4779	Galerucinae_sp_106	S800	KR425399	Bombuscaro	L2	08.03.2012	Beating
BT 4780	Alticinae_sp_252	S800	KR425227	Bombuscaro	L2	08.03.2012	Beating
BT 4781	Alticinae_sp_141	S801		Bombuscaro	L2	08.03.2012	Hand-Coll.(S)
BT 4782	Galerucinae_sp_029	S801	KR425400	Bombuscaro	L2	08.03.2012	Hand-Coll.(S)
BT 4783	Hispinae_sp_016	S802	KR424809	Bombuscaro	L3	08.03.2012	Sweep Netting
BT 4784	Alticinae_sp_245	S805		Cajanuma	L1	13.03.2012	Sweep Netting
BT 4785	Hispinae_sp_027	S806		Cajanuma	L1	13.03.2012	Beating
BT 4786	Alticinae_sp_172	S814	KR425228	Cajanuma	L4	15.03.2012	Sweep Netting
BT 4787	Alticinae_sp_172	S814		Cajanuma	L4	15.03.2012	Sweep Netting
BT 4788	Eumolpinae_sp_058	S814	KR424901	Cajanuma	L4	15.03.2012	Sweep Netting
BT 4789	Alticinae_sp_172	S815	KR425229	Cajanuma	L4	15.03.2012	Beating
BT 4790	Alticinae_sp_189	S817	KR425230	Cajanuma	L5	15.03.2012	Sweep Netting
BT 4791	Alticinae_sp_172	S818	KR425231	Cajanuma	L5	15.03.2012	Beating
BT 4792	Alticinae_sp_187	S821	KR425232	Cajanuma	L6	15.03.2012	Beating
BT 4793	Alticinae_sp_263	S823	KR425233	ECSF	L4	20.03.2012	Sweep Netting
BT 4794	Alticinae_sp_198	S823	KR425234	ECSF	L4	20.03.2012	Sweep Netting
BT 4795	Alticinae_sp_249	S823	KR425235	ECSF	L4	20.03.2012	Sweep Netting
BT 4796	Alticinae_sp_261	S824		ECSF	L4	20.03.2012	Beating
BT 4797	Galerucinae_sp_034	S824	KR425401	ECSF	L4	20.03.2012	Beating
BT 4798	Alticinae_sp_115	S824	KR425236	ECSF	L4	20.03.2012	Beating
BT 4799	Eumolpinae_sp_038	S824		ECSF	L4	20.03.2012	Beating
BT 4800	Alticinae_sp_115	S824	KR425237	ECSF	L4	20.03.2012	Beating
BT 4801	Alticinae_sp_104	S824	KR425238	ECSF	L4	20.03.2012	Beating
BT 4802	Eumolpinae_sp_038	S826		ECSF	L5	20.03.2012	Sweep Netting
BT 4803	Alticinae_sp_101	S826	KR425239	ECSF	L5	20.03.2012	Sweep Netting
BT 4804	Alticinae_sp_115	S826	KR425240	ECSF	L5	20.03.2012	Sweep Netting
BT 4805	Alticinae_sp_115	S827	KR425241	ECSF	L5	20.03.2012	Beating
BT 4806	Alticinae_sp_115	S827		ECSF	L5	20.03.2012	Beating
BT 4807	Eumolpinae_sp_042	S827	KR424902	ECSF	L5	20.03.2012	Beating
BT 4808	Alticinae_sp_018	S827	KR425242	ECSF	L5	20.03.2012	Beating
BT 4809	Galerucinae_sp_034	S827	KR425402	ECSF	L5	20.03.2012	Beating
BT 4810	Alticinae_sp_115	S827	KR425243	ECSF	L5	20.03.2012	Beating
BT 4811	Alticinae_sp_115	S827	KR425244	ECSF	L5	20.03.2012	Beating
BT 4812	Alticinae_sp_101	S827	KR425245	ECSF	L5	20.03.2012	Beating
BT 4813	Alticinae_sp_198	S827	KR425246	ECSF	L5	20.03.2012	Beating
BT 4814	Alticinae_sp_199	S827	KR425247	ECSF	L5	20.03.2012	Beating
BT 4815	Alticinae_sp_086	S827	KR425248	ECSF	L5	20.03.2012	Beating
BT 4816	Alticinae_sp_086	S827		ECSF	L5	20.03.2012	Beating
BT 4817	Alticinae_sp_086	S827		ECSF	L5	20.03.2012	Beating
BT 4851	Eumolpinae_sp_024	S845	KR424903	Bombuscaro	U1	29.03.2012	Beating
BT 4856	Alticinae_sp_186	S853	KR425249	Cajanuma	U4	03.04.2012	Sweep Netting
BT 4857	Alticinae_sp_172	S853	KR425250	Cajanuma	U4	03.04.2012	Sweep Netting
BT 4858	Alticinae_sp_187	S854	KR425251	Cajanuma	U4	03.04.2012	Beating
BT 4859	Alticinae_sp_057	S862	KR425252	Bombuscaro	U4	05.04.2012	Sweep Netting
BT 4860	Eumolpinae_sp_046	S863	KR424904	Bombuscaro	U4	05.04.2012	Beating
BT 4861	Alticinae_sp_149	S863	KR425253	Bombuscaro	U4	05.04.2012	Beating
BT 4862	Alticinae_sp_149	S863	KR425254	Bombuscaro	U4	05.04.2012	Beating
BT 4863	Alticinae_sp_168	S863	KR425255	Bombuscaro	U4	05.04.2012	Beating
BT 4864	Alticinae_sp_149	S863	KR425256	Bombuscaro	U4	05.04.2012	Beating
BT 4867	Galerucinae_sp_003	S866		Bombuscaro	U5	05.04.2012	Beating
BT 4875	Criocerinae_sp_009	S868		Bombuscaro	U6	05.04.2012	Sweep Netting
BT 4885	Galerucinae_sp_056	S871	KR425403	Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4886	Alticinae_sp_121	S871	KR425257	Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4887	Alticinae_sp_102	S871	KR425258	Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4888	Eumolpinae_sp_059	S871	KR424905	Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4889	Alticinae_sp_051	S871	KR425259	Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4890	Alticinae_sp_051	S871		Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4891	Galerucinae_sp_087	S871	KR425404	Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4892	Alticinae_sp_149	S871	KR425260	Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4893	Alticinae_sp_149	S871		Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4894	Alticinae_sp_149	S871		Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4895	Alticinae_sp_149	S871		Bombuscaro	L4	10.04.2012	Sweep Netting
BT 4896	Alticinae_sp_149	S871		Bombuscaro	L4	10.04.2012	Sweep Netting

Continued on next page(s)

Table C.1 – continued from previous page(s)

SpecimenID	Sample ID	GenBank Acc. Nr.	Sampling Area	Site/Plot	Sampling Date	Sampling Method
BT_4897	Alticinae_sp_149	S871	Bombuscaro	L4	10.04.2012	Sweep Netting
BT_4898	Alticinae_sp_168	S871	Bombuscaro	L4	10.04.2012	Sweep Netting
BT_4899	Alticinae_sp_168	S871	Bombuscaro	L4	10.04.2012	Sweep Netting
BT_4900	Alticinae_sp_151	S871	Bombuscaro	L4	10.04.2012	Sweep Netting
BT_4901	Galerucinae_sp_057	S871	Bombuscaro	L4	10.04.2012	Sweep Netting
BT_4902	Galerucinae_sp_057	S871	Bombuscaro	L4	10.04.2012	Sweep Netting
BT_4903	Alticinae_sp_267	S872	Bombuscaro	L4	10.04.2012	Beating
BT_4904	Alticinae_sp_267	S872	Bombuscaro	L4	10.04.2012	Beating
BT_4905	Alticinae_sp_201	S872	Bombuscaro	L4	10.04.2012	Beating
BT_4906	Alticinae_sp_211	S873	Bombuscaro	L4	10.04.2012	Hand-Coll.(S)
BT_4907	Alticinae_sp_211	S873	Bombuscaro	L4	10.04.2012	Hand-Coll.(S)
BT_4908	Hispinae_sp_013	S873	Bombuscaro	L4	10.04.2012	Hand-Coll.(S)
BT_4909	Alticinae_sp_025	S873	Bombuscaro	L4	10.04.2012	Hand-Coll.(S)
BT_4910	Alticinae_sp_149	S873	Bombuscaro	L4	10.04.2012	Hand-Coll.(S)
BT_4911	Alticinae_sp_057	S873	Bombuscaro	L4	10.04.2012	Hand-Coll.(S)
BT_4912	Alticinae_sp_086	S873	Bombuscaro	L4	10.04.2012	Hand-Coll.(S)
BT_4914	Galerucinae_sp_003	S874	Bombuscaro	L5	10.04.2012	Sweep Netting
BT_4915	Alticinae_sp_050	S874	Bombuscaro	L5	10.04.2012	Sweep Netting
BT_4916	Eumolpinae_sp_063	S874	Bombuscaro	L5	10.04.2012	Sweep Netting
BT_4917	Alticinae_sp_166	S875	Bombuscaro	L5	10.04.2012	Beating
BT_4918	Alticinae_sp_050	S875	Bombuscaro	L5	10.04.2012	Beating
BT_4919	Eumolpinae_sp_056	S875	Bombuscaro	L5	10.04.2012	Beating
BT_4920	Alticinae_sp_057	S875	Bombuscaro	L5	10.04.2012	Beating
BT_4921	Alticinae_sp_057	S875	Bombuscaro	L5	10.04.2012	Beating
BT_4922	Alticinae_sp_009	S877	Bombuscaro	L6	10.04.2012	Sweep Netting
BT_4923	Galerucinae_sp_048	S877	Bombuscaro	L6	10.04.2012	Sweep Netting
BT_4924	Alticinae_sp_051	S877	Bombuscaro	L6	10.04.2012	Sweep Netting
BT_4925	Alticinae_sp_102	S878	Bombuscaro	L6	10.04.2012	Beating
BT_4926	Alticinae_sp_025	S879	Bombuscaro	L6	10.04.2012	Hand-Coll.(S)
BT_4927	Alticinae_sp_268	S880	Cajanuma	L1	12.04.2012	Sweep Netting
BT_4928	Alticinae_sp_172	S882	Cajanuma	L1	12.04.2012	Hand-Coll.(S)
BT_4929	Alticinae_sp_172	S884	Cajanuma	L2	12.04.2012	Beating
BT_4930	Alticinae_sp_172	S884	Cajanuma	L2	12.04.2012	Beating
BT_4931	Alticinae_sp_244	S888	Cajanuma	L3	12.04.2012	Hand-Coll.(S)
BT_4937	Alticinae_sp_127	S892	Bombuscaro	U3	17.04.2012	Sweep Netting
BT_4938	Alticinae_sp_149	S893	Bombuscaro	U3	17.04.2012	Beating
BT_4939	Alticinae_sp_149	S893	Bombuscaro	U3	17.04.2012	Beating
BT_4940	Alticinae_sp_059	S893	Bombuscaro	U3	17.04.2012	Beating
BT_4941	Alticinae_sp_127	S893	Bombuscaro	U3	17.04.2012	Beating
BT_4942	Alticinae_sp_102	S893	Bombuscaro	U3	17.04.2012	Beating
BT_4944	Alticinae_sp_150	S895	ECSF	L3	19.04.2012	Sweep Netting
BT_4945	Eumolpinae_sp_002	S895	ECSF	L3	19.04.2012	Sweep Netting
BT_4946	Eumolpinae_sp_034	S895	ECSF	L3	19.04.2012	Sweep Netting
BT_4947	Alticinae_sp_199	S895	ECSF	L3	19.04.2012	Sweep Netting
BT_4948	Alticinae_sp_150	S896	ECSF	L3	19.04.2012	Beating
BT_4949	Alticinae_sp_150	S896	ECSF	L3	19.04.2012	Beating
BT_4950	Galerucinae_sp_088	S896	ECSF	L3	19.04.2012	Beating
BT_4951	Alticinae_sp_061	S896	ECSF	L3	19.04.2012	Beating
BT_4952	Alticinae_sp_150	S896	ECSF	L3	19.04.2012	Beating
BT_4953	Alticinae_sp_199	S896	ECSF	L3	19.04.2012	Beating
BT_4954	Alticinae_sp_199	S896	ECSF	L3	19.04.2012	Beating
BT_4955	Alticinae_sp_199	S896	ECSF	L3	19.04.2012	Beating
BT_4956	Alticinae_sp_086	S896	ECSF	L3	19.04.2012	Beating
BT_4957	Alticinae_sp_150	S898	ECSF	U3	19.04.2012	Sweep Netting
BT_4958	Alticinae_sp_133	S898	ECSF	U3	19.04.2012	Sweep Netting
BT_4960	Alticinae_sp_086	S898	ECSF	U3	19.04.2012	Sweep Netting
BT_4961	Alticinae_sp_086	S898	ECSF	U3	19.04.2012	Sweep Netting
BT_4962	Alticinae_sp_086	S898	ECSF	U3	19.04.2012	Sweep Netting
BT_4963	Alticinae_sp_215	S899	ECSF	U3	19.04.2012	Beating
BT_4964	Alticinae_sp_150	S899	ECSF	U3	19.04.2012	Beating
BT_4965	Eumolpinae_sp_034	S899	ECSF	U3	19.04.2012	Beating
BT_4966	Alticinae_sp_008	S899	ECSF	U3	19.04.2012	Beating
BT_4967	Eumolpinae_sp_038	S899	ECSF	U3	19.04.2012	Beating
BT_4968	Eumolpinae_sp_038	S899	ECSF	U3	19.04.2012	Beating
BT_4969	Alticinae_sp_045	S900	ECSF	U3	19.04.2012	Hand-Coll.(S)
BT_4970	Galerucinae_sp_081	S901	Bombuscaro	U5	24.04.2012	Sweep Netting
BT_4971	Alticinae_sp_192	S901	Bombuscaro	U5	24.04.2012	Sweep Netting
BT_4972	Alticinae_sp_050	S902	Bombuscaro	U5	24.04.2012	Beating
BT_4986	Galerucinae_sp_101	S908	Bombuscaro	L4	24.04.2012	Beating
BT_4994	Alticinae_sp_172	S911	Cajanuma	L1	26.04.2012	Beating
BT_4995	Galerucinae_sp_103	S918	Cajanuma	U3	26.04.2012	Hand-Coll.(S)
BT_5029	Galerucinae_sp_099	N242	Bombuscaro		27.02.2012	Hand-Coll.(N)
BT_5122	Alticinae_sp_221	N266	Bombuscaro		24.04.2012	Hand-Coll.(N)

## Outgroup specimens:

Species	SequenceID/Accession Nr.	Source
<i>Anthonomus eugenii</i>	SequenceID ARBCP010-10	BOLD
<i>Dichromacalles dromedarius</i>	Accession number GU987917	GenBank
<i>Acalles camelus</i>	Accession number GU987989	GenBank

# Additional information: Data sets

Table D.1: Specimen list with data sets for which the specimen was used.

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT_0001 Eumolpinae sp_001	X	X	X	X	X				X
BT_0002 Alticinae sp_042	X	X	X	X	X				X
BT_0003 Alticinae sp_042				X					
BT_0003a Alticinae sp_042				X					
BT_0004 Eumolpinae sp_042	X	X	X	X	X				X
BT_0005 Galerucinae sp_040	X	X	X	X	X				X
BT_0006 Galerucinae sp_040				X					
BT_0006a Galerucinae sp_040				X					
BT_0007 Galerucinae sp_038	X	X	X	X	X				X
BT_0008 Alticinae sp_243	X	X	X	X	X				X
BT_0009 Eumolpinae sp_022	X		X	X					
BT_0010 Eumolpinae sp_022				X					
BT_0011 Eumolpinae sp_022				X					
BT_0012 Eumolpinae sp_021	X	X	X	X	X				X
BT_0013 Eumolpinae sp_021				X					
BT_0014 Eumolpinae sp_021				X					
BT_0015 Galerucinae sp_076	X	X	X	X	X				X
BT_0016 Galerucinae sp_076				X					
BT_0017 Alticinae sp_043	X	X	X	X	X				X
BT_0018 Alticinae sp_043				X					
BT_0019 Alticinae sp_043				X					
BT_0020 Alticinae sp_043				X					
BT_0021 Alticinae sp_007	X	X	X	X	X				X
BT_0022 Alticinae sp_219	X	X	X	X	X				X
BT_0024 Galerucinae sp_001	X	X	X	X	X				X
BT_0025 Galerucinae sp_001				X					
BT_0026 Galerucinae sp_001				X					
BT_0027 Galerucinae sp_001				X					
BT_0028 Galerucinae sp_001				X					
BT_0029 Galerucinae sp_001				X					
BT_0030 Galerucinae sp_001				X					
BT_0031 Galerucinae sp_001				X					
BT_0032 Galerucinae sp_001				X					
BT_0033 Galerucinae sp_037	X	X	X	X	X				X
BT_0034 Eumolpinae sp_014	X	X	X	X	X				X
BT_0035 Eumolpinae sp_006	X	X	X	X	X				X
BT_0036 Galerucinae sp_011	X	X	X	X	X				X
BT_0043 Galerucinae sp_005	X	X	X	X	X				X
BT_0044 Galerucinae sp_005				X					
BT_0045 Galerucinae sp_005				X					
BT_0045a Galerucinae sp_005				X					
BT_0046 Alticinae sp_243	X	X	X	X	X				X
BT_0047 Alticinae sp_042	X	X	X	X	X				X
BT_0048 Galerucinae sp_039	X	X	X	X	X				X
BT_0049 Galerucinae sp_041	X	X	X	X	X				X
BT_0050 Galerucinae sp_002				X					
BT_0051 Galerucinae sp_002				X					
BT_0052 Galerucinae sp_002				X					
BT_0053 Galerucinae sp_002				X					
BT_0054 Galerucinae sp_002				X					
BT_0055 Galerucinae sp_002				X					
BT_0056 Galerucinae sp_002				X					
BT_0057 Galerucinae sp_002				X					
BT_0058 Galerucinae sp_002				X					
BT_0059 Galerucinae sp_002				X					
BT_0060 Galerucinae sp_004				X					
BT_0061 Galerucinae sp_007				X					
BT_0062 Galerucinae sp_007				X					
BT_0063 Galerucinae sp_007				X					
BT_0064 Galerucinae sp_007				X					
BT_0065 Galerucinae sp_007				X					
BT_0066 Galerucinae sp_007				X					
BT_0067 Galerucinae sp_007				X					
BT_0068 Galerucinae sp_007				X					
BT_0069 Galerucinae sp_007				X					
BT_0070 Galerucinae sp_007				X					
BT_0071 Galerucinae sp_007				X					
BT_0072 Galerucinae sp_007				X					
BT_0085 Eumolpinae sp_048	X		X	X					
BT_0088 Galerucinae sp_007	X	X	X	X	X				X
BT_0089 Eumolpinae sp_001	X	X	X	X	X				X
BT_0090 Galerucinae sp_076	X	X	X	X	X				X
BT_0091 Eumolpinae sp_001	X	X	X	X	X				X
BT_0092 Eumolpinae sp_001				X					
BT_0093 Eumolpinae sp_001				X					
BT_0094 Galerucinae sp_011	X	X	X	X	X				X
BT_0095 Cassidinae sp_001	X	X	X	X	X				X
BT_0096 Alticinae sp_010	X	X	X	X	X				X
BT_0097 Alticinae sp_010				X					
BT_0098 Galerucinae sp_002	X	X	X	X	X				X
BT_0099 Galerucinae sp_007	X	X	X	X	X				X
BT_0100 Galerucinae sp_007				X					
BT_0101 Galerucinae sp_007				X					
BT_0102 Alticinae sp_044	X	X	X	X	X				X
BT_0103 Eumolpinae sp_038	X	X	X	X	X				X
BT_0104 Eumolpinae sp_038				X					
BT_0105 Eumolpinae sp_038				X					
BT_0106 Eumolpinae sp_038				X					
BT_0107 Galerucinae sp_046	X	X	X	X	X				X
BT_0108 Galerucinae sp_046				X					
BT_0109 Alticinae sp_251	X	X	X	X	X				X

Continued on next page(s)

Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT_0110_Alticinae_sp_087	X	X	X	X	X				X
BT_0111_Alticinae_sp_087				X	X				X
BT_0112_Alticinae_sp_087				X	X				X
BT_0113_Alticinae_sp_087				X	X				X
BT_0114_Galerucinae_sp_062	X	X	X	X	X				X
BT_0115_Alticinae_sp_098	X	X	X	X	X				X
BT_0116_Alticinae_sp_098				X	X				X
BT_0117_Eumolpinae_sp_036	X		X	X	X				X
BT_0118_Eumolpinae_sp_019	X	X	X	X	X				X
BT_0119_Alticinae_sp_124	X	X	X	X	X				X
BT_0120_Alticinae_sp_064	X		X	X	X				X
BT_0121_Alticinae_sp_107	X	X	X	X	X				X
BT_0122_Alticinae_sp_107				X	X				X
BT_0123_Alticinae_sp_129	X	X	X	X	X				X
BT_0124_Alticinae_sp_129				X	X				X
BT_0125_Alticinae_sp_097	X	X	X	X	X				X
BT_0126_Alticinae_sp_123	X	X	X	X	X				X
BT_0127_Hispinae_sp_001	X		X	X	X				X
BT_0128_Eumolpinae_sp_009	X		X	X	X				X
BT_0129_Eumolpinae_sp_009				X	X				X
BT_0130_Galerucinae_sp_034	X	X	X	X	X				X
BT_0131_Galerucinae_sp_034				X	X				X
BT_0133_Cassidinae_sp_003	X		X	X	X				X
BT_0134_Galerucinae_sp_007	X	X	X	X	X				X
BT_0135_Eumolpinae_sp_019	X	X	X	X	X				X
BT_0136_Eumolpinae_sp_019				X	X				X
BT_0137_Cassidinae_sp_004	X	X	X	X	X				X
BT_0138_Cassidinae_sp_004				X	X				X
BT_0139_Alticinae_sp_010	X	X	X	X	X				X
BT_0140_Alticinae_sp_028	X	X	X	X	X				X
BT_0144_Eumolpinae_sp_038	X	X	X	X	X				X
BT_0145_Galerucinae_sp_061	X	X	X	X	X				X
BT_0146_Alticinae_sp_029	X	X	X	X	X				X
BT_0147_Alticinae_sp_062	X	X	X	X	X				X
BT_0148_Alticinae_sp_066	X	X	X	X	X				X
BT_0149_Alticinae_sp_249	X	X	X	X	X				X
BT_0150_Alticinae_sp_249				X	X				X
BT_0151_Alticinae_sp_249				X	X				X
BT_0152_Alticinae_sp_249				X	X				X
BT_0153_Alticinae_sp_109	X	X	X	X	X				X
BT_0154_Alticinae_sp_115	X	X	X	X	X				X
BT_0155_Alticinae_sp_193	X	X	X	X	X				X
BT_0156_Alticinae_sp_019	X	X	X	X	X				X
BT_0157_Alticinae_sp_097	X	X	X	X	X				X
BT_0158_Eumolpinae_sp_002	X	X	X	X	X				X
BT_0159_Galerucinae_sp_096	X	X	X	X	X				X
BT_0160_Galerucinae_sp_096				X	X				X
BT_0161_Galerucinae_sp_096				X	X				X
BT_0162_Galerucinae_sp_096				X	X				X
BT_0163_Galerucinae_sp_096				X	X				X
BT_0164_Galerucinae_sp_096				X	X				X
BT_0165_Galerucinae_sp_096				X	X				X
BT_0166_Galerucinae_sp_096				X	X				X
BT_0167_Galerucinae_sp_096				X	X				X
BT_0168_Galerucinae_sp_096				X	X				X
BT_0169_Galerucinae_sp_096				X	X				X
BT_0170_Galerucinae_sp_096				X	X				X
BT_0171_Galerucinae_sp_096				X	X				X
BT_0172_Galerucinae_sp_096				X	X				X
BT_0173_Galerucinae_sp_096				X	X				X
BT_0174_Galerucinae_sp_046	X	X	X	X	X				X
BT_0175_Galerucinae_sp_046				X	X				X
BT_0176_Galerucinae_sp_046	X	X	X	X	X				X
BT_0177_Galerucinae_sp_046				X	X				X
BT_0178_Galerucinae_sp_046				X	X				X
BT_0179_Galerucinae_sp_046				X	X				X
BT_0180_Galerucinae_sp_046				X	X				X
BT_0181_Galerucinae_sp_046				X	X				X
BT_0182_Galerucinae_sp_046				X	X				X
BT_0183_Galerucinae_sp_034	X	X	X	X	X				X
BT_0184_Galerucinae_sp_034				X	X				X
BT_0188_Galerucinae_sp_011	X	X	X	X	X				X
BT_0189_Alticinae_sp_161	X	X	X	X	X				X
BT_0190_Eumolpinae_sp_001	X	X	X	X	X				X
BT_0191_Eumolpinae_sp_001				X	X				X
BT_0195_Galerucinae_sp_045	X	X	X	X	X				X
BT_0196_Galerucinae_sp_010	X	X	X	X	X				X
BT_0197_Galerucinae_sp_033	X	X	X	X	X				X
BT_0198_Alticinae_sp_137	X	X	X	X	X				X
BT_0199_Alticinae_sp_118	X	X	X	X	X				X
BT_0201_Chrysomelinae_sp_002	X	X	X	X	X				X
BT_0202_Galerucinae_sp_032	X	X	X	X	X				X
BT_0203_Galerucinae_sp_032				X	X				X
BT_0204_Hispinae_sp_002	X	X	X	X	X				X
BT_0206_Alticinae_sp_085	X	X	X	X	X				X
BT_0207_Galerucinae_sp_069	X	X	X	X	X				X
BT_0208_Eumolpinae_sp_019	X	X	X	X	X				X
BT_0209_Cassidinae_sp_005	X	X	X	X	X				X
BT_0210_Cassidinae_sp_006	X	X	X	X	X				X
BT_0211_Alticinae_sp_087	X	X	X	X	X				X
BT_0212_Galerucinae_sp_066	X	X	X	X	X				X
BT_0213_Galerucinae_sp_024	X	X	X	X	X				X
BT_0214_Alticinae_sp_028	X	X	X	X	X				X
BT_0215_Alticinae_sp_028				X	X				X
BT_0216_Eumolpinae_sp_038	X		X	X	X				X
BT_0217_Eumolpinae_sp_038				X	X				X
BT_0218_Galerucinae_sp_031	X	X	X	X	X				X
BT_0219_Eumolpinae_sp_073	X	X	X	X	X				X
BT_0220_Alticinae_sp_115	X	X	X	X	X				X
BT_0221_Eumolpinae_sp_010	X	X	X	X	X				X
BT_0223_Galerucinae_sp_034	X	X	X	X	X				X
BT_0224_Galerucinae_sp_034				X	X				X
BT_0227_Cassidinae_sp_014	X	X	X	X	X				X
BT_0228_Cassidinae_sp_007	X	X	X	X	X				X
BT_0229_Cassidinae_sp_007				X	X				X
BT_0230_Eumolpinae_sp_022	X	X	X	X	X				X
BT_0231_Hispinae_sp_003	X	X	X	X	X				X
BT_0232_Hispinae_sp_004	X	X	X	X	X				X
BT_0233_Alticinae_sp_061	X	X	X	X	X				X
BT_0234_Alticinae_sp_097	X	X	X	X	X				X
BT_0235_Alticinae_sp_010	X	X	X	X	X				X
BT_0236_Alticinae_sp_156	X	X	X	X	X				X
BT_0237_Alticinae_sp_156				X	X				X
BT_0239_Galerucinae_sp_002	X	X	X	X	X				X
BT_0240_Eumolpinae_sp_021	X	X	X	X	X				X
BT_0241_Eumolpinae_sp_038	X		X	X	X				X
BT_0242_Eumolpinae_sp_038				X	X				X

Continued on next page(s)

Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT_0243_Alticinae_sp_118	X	X	X	X	X				X
BT_0244_Alticinae_sp_097	X	X	X	X	X				X
BT_0245_Galerucinae_sp_034	X	X	X	X	X				X
BT_0246_Galerucinae_sp_030	X	X	X	X	X				X
BT_0247_Criocerinae_sp_001	X	X	X	X	X				X
BT_0249_Criocerinae_sp_001	X	X	X	X	X				X
BT_0250_Alticinae_sp_115	X	X	X	X	X				X
BT_0251_Alticinae_sp_115									
BT_0252_Galerucinae_sp_034	X	X	X	X	X				X
BT_0253_Galerucinae_sp_034									
BT_0254_Eumolpinae_sp_023	X	X	X	X	X				X
BT_0255_Eumolpinae_sp_023									
BT_0256_Alticinae_sp_031	X	X	X	X	X				X
BT_0257_Galerucinae_sp_001	X	X	X	X	X				X
BT_0258_Galerucinae_sp_005	X	X	X	X	X				X
BT_0259_Alticinae_sp_096	X	X	X	X	X				X
BT_0260_Eumolpinae_sp_038	X	X	X	X	X				X
BT_0261_Eumolpinae_sp_038									
BT_0266_Alticinae_sp_145	X	X	X	X	X				X
BT_0267_Alticinae_sp_029	X	X	X	X	X				X
BT_0268_Alticinae_sp_158	X	X	X	X	X				X
BT_0269_Alticinae_sp_086	X	X	X	X	X				X
BT_0270_Alticinae_sp_086									
BT_0271_Alticinae_sp_064	X	X	X	X	X				X
BT_0272_Alticinae_sp_064									
BT_0273_Alticinae_sp_141	X	X	X	X	X				X
BT_0274_Alticinae_sp_141									
BT_0275_Alticinae_sp_141									
BT_0276_Alticinae_sp_122									
BT_0277_Alticinae_sp_122									
BT_0278_Alticinae_sp_124	X	X	X	X	X				X
BT_0279_Alticinae_sp_115	X	X	X	X	X				X
BT_0280_Eumolpinae_sp_022	X	X	X	X	X				X
BT_0281_Eumolpinae_sp_038	X	X	X	X	X				X
BT_0282_Eumolpinae_sp_038									
BT_0283_Eumolpinae_sp_020	X	X	X	X	X				X
BT_0284_Alticinae_sp_087	X	X	X	X	X				X
BT_0285_Galerucinae_sp_022	X	X	X	X	X				X
BT_0286_Alticinae_sp_012	X	X	X	X	X				X
BT_0287_Eumolpinae_sp_038	X	X	X	X	X				X
BT_0288_Eumolpinae_sp_017	X	X	X	X	X				X
BT_0289_Eumolpinae_sp_001	X	X	X	X	X				X
BT_0290_Eumolpinae_sp_001									
BT_0291_Eumolpinae_sp_001									
BT_0292_Galerucinae_sp_076	X	X	X	X	X				X
BT_0293_Galerucinae_sp_076									
BT_0294_Galerucinae_sp_076									
BT_0295_Galerucinae_sp_031	X	X	X	X	X				X
BT_0296_Alticinae_sp_005	X	X	X	X	X				X
BT_0297_Alticinae_sp_142	X	X	X	X	X				X
BT_0298_Alticinae_sp_063	X	X	X	X	X				X
BT_0299_Alticinae_sp_064	X	X	X	X	X				X
BT_0300_Alticinae_sp_064									
BT_0301_Alticinae_sp_083	X	X	X	X	X				X
BT_0302_Alticinae_sp_086	X	X	X	X	X				X
BT_0303_Alticinae_sp_086									
BT_0304_Alticinae_sp_086									
BT_0305_Alticinae_sp_081	X	X	X	X	X				X
BT_0306_Alticinae_sp_019	X	X	X	X	X				X
BT_0307_Alticinae_sp_115	X	X	X	X	X				X
BT_0308_Alticinae_sp_018	X	X	X	X	X				X
BT_0309_Alticinae_sp_256	X	X	X	X	X				X
BT_0310_Eumolpinae_sp_017	X	X	X	X	X				X
BT_0311_Eumolpinae_sp_042	X	X	X	X	X				X
BT_0312_Eumolpinae_sp_021	X	X	X	X	X				X
BT_0313_Alticinae_sp_122	X	X	X	X	X				X
BT_0314_Eumolpinae_sp_043	X	X	X	X	X				X
BT_0318_Alticinae_sp_124	X	X	X	X	X				X
BT_0319_Galerucinae_sp_026	X	X	X	X	X				X
BT_0320_Galerucinae_sp_061	X	X	X	X	X				X
BT_0322_Alticinae_sp_143	X	X	X	X	X				X
BT_0323_Eumolpinae_sp_011	X	X	X	X	X				X
BT_0336_Galerucinae_sp_056	X	X	X	X	X				X
BT_0337_Galerucinae_sp_056	X	X	X	X	X				X
BT_0338_Galerucinae_sp_055	X	X	X	X	X				X
BT_0339_Alticinae_sp_071	X	X	X	X	X				X
BT_0340_Alticinae_sp_064	X	X	X	X	X				X
BT_0341_Alticinae_sp_073	X	X	X	X	X				X
BT_0342_Galerucinae_sp_051	X	X	X	X	X				X
BT_0343_Galerucinae_sp_052	X	X	X	X	X				X
BT_0344_Galerucinae_sp_052									
BT_0345_Alticinae_sp_051	X	X	X	X	X				X
BT_0346_Alticinae_sp_051	X	X	X	X	X				X
BT_0347_Alticinae_sp_051									
BT_0348_Alticinae_sp_051									
BT_0349_Eumolpinae_sp_020	X	X	X	X	X				X
BT_0350_Eumolpinae_sp_045	X	X	X	X	X				X
BT_0351_Eumolpinae_sp_045									
BT_0352_Alticinae_sp_130	X	X	X	X	X				X
BT_0353_Alticinae_sp_132	X	X	X	X	X				X
BT_0354_Alticinae_sp_125	X	X	X	X	X				X
BT_0355_Alticinae_sp_125									
BT_0356_Alticinae_sp_128	X	X	X	X	X				X
BT_0357_Alticinae_sp_128									
BT_0358_Alticinae_sp_128									
BT_0359_Alticinae_sp_128									
BT_0361_Alticinae_sp_050	X	X	X	X	X				X
BT_0362_Alticinae_sp_114	X	X	X	X	X				X
BT_0363_Alticinae_sp_051	X	X	X	X	X				X
BT_0364_Eumolpinae_sp_018	X	X	X	X	X				X
BT_0365_Galerucinae_sp_049	X	X	X	X	X				X
BT_0366_Galerucinae_sp_049									
BT_0367_Galerucinae_sp_049									
BT_0368_Galerucinae_sp_049									
BT_0369_Galerucinae_sp_049									
BT_0370_Galerucinae_sp_049									
BT_0371_Eumolpinae_sp_006	X	X	X	X	X				X
BT_0372_Alticinae_sp_009	X	X	X	X	X				X
BT_0373_Alticinae_sp_009									
BT_0374_Alticinae_sp_024	X	X	X	X	X				X
BT_0375_Eumolpinae_sp_032	X	X	X	X	X				X
BT_0376_Alticinae_sp_051	X	X	X	X	X				X
BT_0377_Eumolpinae_sp_024	X	X	X	X	X				X
BT_0378_Eumolpinae_sp_024									
BT_0380_Eumolpinae_sp_024	X	X	X	X	X				X
BT_0381_Galerucinae_sp_089	X	X	X	X	X				X
BT_0382_Galerucinae_sp_013	X	X	X	X	X				X

Continued on next page(s)

Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT_0383_Alticinae_sp_087	X	X	X	X	X	X	X	X	X
BT_0384_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0385_Alticinae_sp_064	X	X	X	X	X	X	X	X	X
BT_0386_Alticinae_sp_064	X	X	X	X	X	X	X	X	X
BT_0387_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_0388_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_0389_Alticinae_sp_105	X	X	X	X	X	X	X	X	X
BT_0390_Alticinae_sp_090	X	X	X	X	X	X	X	X	X
BT_0391_Alticinae_sp_090	X	X	X	X	X	X	X	X	X
BT_0392_Eumolpinae_sp_074	X	X	X	X	X	X	X	X	X
BT_0395_Eumolpinae_sp_074	X	X	X	X	X	X	X	X	X
BT_0396_Cassidinae_sp_004	X	X	X	X	X	X	X	X	X
BT_0397_Alticinae_sp_131	X	X	X	X	X	X	X	X	X
BT_0398_Cassidinae_sp_003	X	X	X	X	X	X	X	X	X
BT_0399_Galerucinae_sp_028	X	X	X	X	X	X	X	X	X
BT_0400_Galerucinae_sp_028	X	X	X	X	X	X	X	X	X
BT_0401_Galerucinae_sp_054	X	X	X	X	X	X	X	X	X
BT_0402_Galerucinae_sp_044	X	X	X	X	X	X	X	X	X
BT_0403_Galerucinae_sp_044	X	X	X	X	X	X	X	X	X
BT_0404_Galerucinae_sp_044	X	X	X	X	X	X	X	X	X
BT_0405_Galerucinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0406_Galerucinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0407_Galerucinae_sp_049	X	X	X	X	X	X	X	X	X
BT_0408_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_0409_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_0410_Alticinae_sp_141	X	X	X	X	X	X	X	X	X
BT_0411_Alticinae_sp_058	X	X	X	X	X	X	X	X	X
BT_0412_Alticinae_sp_058	X	X	X	X	X	X	X	X	X
BT_0413_Alticinae_sp_058	X	X	X	X	X	X	X	X	X
BT_0414_Alticinae_sp_058	X	X	X	X	X	X	X	X	X
BT_0415_Alticinae_sp_242	X	X	X	X	X	X	X	X	X
BT_0416_Alticinae_sp_242	X	X	X	X	X	X	X	X	X
BT_0417_Galerucinae_sp_074	X	X	X	X	X	X	X	X	X
BT_0418_Eumolpinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0419_Alticinae_sp_141	X	X	X	X	X	X	X	X	X
BT_0420_Alticinae_sp_128	X	X	X	X	X	X	X	X	X
BT_0421_Alticinae_sp_073	X	X	X	X	X	X	X	X	X
BT_0422_Galerucinae_sp_028	X	X	X	X	X	X	X	X	X
BT_0423_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_0425_Eumolpinae_sp_007	X	X	X	X	X	X	X	X	X
BT_0426_Alticinae_sp_150	X	X	X	X	X	X	X	X	X
BT_0427_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_0428_Alticinae_sp_105	X	X	X	X	X	X	X	X	X
BT_0429_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0430_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0431_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0432_Eumolpinae_sp_030	X	X	X	X	X	X	X	X	X
BT_0433_Galerucinae_sp_069	X	X	X	X	X	X	X	X	X
BT_0434_Galerucinae_sp_072	X	X	X	X	X	X	X	X	X
BT_0436_Alticinae_sp_149	X	X	X	X	X	X	X	X	X
BT_0437_Alticinae_sp_149	X	X	X	X	X	X	X	X	X
BT_0438_Galerucinae_sp_015	X	X	X	X	X	X	X	X	X
BT_0439_Galerucinae_sp_015	X	X	X	X	X	X	X	X	X
BT_0440_Alticinae_sp_025	X	X	X	X	X	X	X	X	X
BT_0441_Galerucinae_sp_082	X	X	X	X	X	X	X	X	X
BT_0442_Galerucinae_sp_049	X	X	X	X	X	X	X	X	X
BT_0443_Alticinae_sp_051	X	X	X	X	X	X	X	X	X
BT_0444_Alticinae_sp_143	X	X	X	X	X	X	X	X	X
BT_0445_Alticinae_sp_153	X	X	X	X	X	X	X	X	X
BT_0446_Alticinae_sp_153	X	X	X	X	X	X	X	X	X
BT_0447_Galerucinae_sp_007	X	X	X	X	X	X	X	X	X
BT_0448_Galerucinae_sp_049	X	X	X	X	X	X	X	X	X
BT_0449_Galerucinae_sp_049	X	X	X	X	X	X	X	X	X
BT_0451_Galerucinae_sp_015	X	X	X	X	X	X	X	X	X
BT_0452_Alticinae_sp_009	X	X	X	X	X	X	X	X	X
BT_0453_Alticinae_sp_135	X	X	X	X	X	X	X	X	X
BT_0454_Alticinae_sp_026	X	X	X	X	X	X	X	X	X
BT_0455_Alticinae_sp_026	X	X	X	X	X	X	X	X	X
BT_0456_Alticinae_sp_026	X	X	X	X	X	X	X	X	X
BT_0457_Alticinae_sp_181	X	X	X	X	X	X	X	X	X
BT_0458_Alticinae_sp_181	X	X	X	X	X	X	X	X	X
BT_0459_Galerucinae_sp_050	X	X	X	X	X	X	X	X	X
BT_0460_Galerucinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0461_Galerucinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0462_Eumolpinae_sp_042	X	X	X	X	X	X	X	X	X
BT_0463_Galerucinae_sp_070	X	X	X	X	X	X	X	X	X
BT_0464_Eumolpinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0465_Alticinae_sp_050	X	X	X	X	X	X	X	X	X
BT_0467_Eumolpinae_sp_018	X	X	X	X	X	X	X	X	X
BT_0468_Alticinae_sp_072	X	X	X	X	X	X	X	X	X
BT_0469_Galerucinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0470_Galerucinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0471_Galerucinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0472_Galerucinae_sp_045	X	X	X	X	X	X	X	X	X
BT_0473_Galerucinae_sp_069	X	X	X	X	X	X	X	X	X
BT_0474_Alticinae_sp_080	X	X	X	X	X	X	X	X	X
BT_0475_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0476_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0477_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_0478_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_0480_Hispinae_sp_005	X	X	X	X	X	X	X	X	X
BT_0489_Galerucinae_sp_067	X	X	X	X	X	X	X	X	X
BT_0490_Hispinae_sp_005	X	X	X	X	X	X	X	X	X
BT_0491_Alticinae_sp_157	X	X	X	X	X	X	X	X	X
BT_0492_Galerucinae_sp_061	X	X	X	X	X	X	X	X	X
BT_0493_Eumolpinae_sp_037	X	X	X	X	X	X	X	X	X
BT_0494_Alticinae_sp_250	X	X	X	X	X	X	X	X	X
BT_0495_Alticinae_sp_148	X	X	X	X	X	X	X	X	X
BT_0495a_Alticinae_sp_148	X	X	X	X	X	X	X	X	X
BT_0496_Alticinae_sp_112	X	X	X	X	X	X	X	X	X
BT_0497_Alticinae_sp_111	X	X	X	X	X	X	X	X	X
BT_0498_Alticinae_sp_113	X	X	X	X	X	X	X	X	X
BT_0499_Alticinae_sp_044	X	X	X	X	X	X	X	X	X
BT_0500_Alticinae_sp_044	X	X	X	X	X	X	X	X	X
BT_0501_Galerucinae_sp_066	X	X	X	X	X	X	X	X	X
BT_0502_Eumolpinae_sp_042	X	X	X	X	X	X	X	X	X
BT_0503_Alticinae_sp_111	X	X	X	X	X	X	X	X	X
BT_0504_Alticinae_sp_111	X	X	X	X	X	X	X	X	X
BT_0505_Alticinae_sp_096	X	X	X	X	X	X	X	X	X
BT_0506_Alticinae_sp_092	X	X	X	X	X	X	X	X	X
BT_0507_Alticinae_sp_092	X	X	X	X	X	X	X	X	X
BT_0508_Galerucinae_sp_053	X	X	X	X	X	X	X	X	X
BT_0509_Cassidinae_sp_003	X	X	X	X	X	X	X	X	X
BT_0510_Cassidinae_sp_004	X	X	X	X	X	X	X	X	X
BT_0511_Cassidinae_sp_008	X	X	X	X	X	X	X	X	X
BT_0512_Hispinae_sp_006	X	X	X	X	X	X	X	X	X
BT_0513_Hispinae_sp_006	X	X	X	X	X	X	X	X	X

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 0514 Eumolpinae sp 074	X	X	X	X	X	X	X	X	X
BT 0515 Eumolpinae sp 074			X	X					
BT 0516 Alticinae sp 126	X		X	X		X	X		X
BT 0517 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 0518 Alticinae sp 085	X	X	X	X	X	X	X	X	X
BT 0519 Alticinae sp 068	X	X	X	X	X	X	X	X	X
BT 0520 Alticinae sp 159	X	X	X	X	X	X	X	X	X
BT 0524 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 0525 Hispinae sp 023	X	X	X	X	X	X	X	X	X
BT 0526 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 0527 Galerucinae sp 036	X	X	X	X	X	X	X	X	X
BT 0528 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 0529 Alticinae sp 113	X	X	X	X	X	X	X	X	X
BT 0530 Galerucinae sp 031	X	X	X	X	X	X	X	X	X
BT 0531 Galerucinae sp 062	X	X	X	X	X	X	X	X	X
BT 0532 Galerucinae sp 075	X	X	X	X	X	X	X	X	X
BT 0533 Alticinae sp 097	X	X	X	X	X	X	X	X	X
BT 0534 Alticinae sp 111	X	X	X	X	X	X	X	X	X
BT 0535 Alticinae sp 097	X	X	X	X	X	X	X	X	X
BT 0536 Cassidinae sp 003	X	X	X	X	X	X	X	X	X
BT 0537 Alticinae sp 142	X	X	X	X	X	X	X	X	X
BT 0538 Alticinae sp 238	X	X	X	X	X	X	X	X	X
BT 0538a Alticinae sp 238									
BT 0539 Alticinae sp 013	X	X	X	X	X	X	X	X	X
BT 0540 Galerucinae sp 064	X	X	X	X	X	X	X	X	X
BT 0541 Galerucinae sp 064									
BT 0543 Cassidinae sp 009	X	X	X	X	X	X	X	X	X
BT 0544 Cassidinae sp 012	X	X	X	X	X	X	X	X	X
BT 0546 Alticinae sp 083	X	X	X	X	X	X	X	X	X
BT 0547 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 0548 Alticinae sp 112	X	X	X	X	X	X	X	X	X
BT 0549 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 0550 Alticinae sp 265	X	X	X	X	X	X	X	X	X
BT 0551 Galerucinae sp 064	X	X	X	X	X	X	X	X	X
BT 0552 Alticinae sp 117	X	X	X	X	X	X	X	X	X
BT 0553 Eumolpinae sp 042	X	X	X	X	X	X	X	X	X
BT 0554 Galerucinae sp 071	X	X	X	X	X	X	X	X	X
BT 0555 Alticinae sp 086	X	X	X	X	X	X	X	X	X
BT 0556 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 0557 Alticinae sp 181	X	X	X	X	X	X	X	X	X
BT 0558 Alticinae sp 149	X	X	X	X	X	X	X	X	X
BT 0559 Eumolpinae sp 042	X	X	X	X	X	X	X	X	X
BT 0560 Galerucinae sp 064	X	X	X	X	X	X	X	X	X
BT 0561 Galerucinae sp 066	X	X	X	X	X	X	X	X	X
BT 0562 Cassidinae sp 009	X		X	X	X	X	X	X	X
BT 0563 Cassidinae sp 009									
BT 0564 Cassidinae sp 009									
BT 0565 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 0566 Hispinae sp 006	X	X	X	X	X	X	X	X	X
BT 0567 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 0568 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 0569 Alticinae sp 118									
BT 0570 Alticinae sp 118									
BT 0571 Alticinae sp 118									
BT 0572 Alticinae sp 118									
BT 0573 Alticinae sp 118									
BT 0574 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 0575 Alticinae sp 140	X	X	X	X	X	X	X	X	X
BT 0576 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 0577 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 0578 Eumolpinae sp 039									
BT 0579 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 0580 Alticinae sp 104									
BT 0581 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 0582 Alticinae sp 118									
BT 0583 Alticinae sp 118									
BT 0584 Alticinae sp 118									
BT 0585 Eumolpinae sp 044	X		X	X	X	X	X	X	X
BT 0586 Alticinae sp 091	X		X	X	X	X	X	X	X
BT 0587 Alticinae sp 006	X	X	X	X	X	X	X	X	X
BT 0588 Galerucinae sp 069	X	X	X	X	X	X	X	X	X
BT 0589 Eumolpinae sp 017	X	X	X	X	X	X	X	X	X
BT 0590 Alticinae sp 140	X	X	X	X	X	X	X	X	X
BT 0592 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 0593 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 0594 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 0595 Alticinae sp 118									
BT 0596 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 0597 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 0598 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 0599 Alticinae sp 118									
BT 0600 Alticinae sp 118									
BT 0601 Alticinae sp 118									
BT 0602 Alticinae sp 118									
BT 0603 Alticinae sp 118									
BT 0604 Hispinae sp 005	X	X	X	X	X	X	X	X	X
BT 0605 Alticinae sp 087	X	X	X	X	X	X	X	X	X
BT 0606 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 0607 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 0608 Alticinae sp 118									
BT 0609 Alticinae sp 118									
BT 0610 Alticinae sp 118									
BT 0611 Alticinae sp 118									
BT 0612 Alticinae sp 118									
BT 0613 Alticinae sp 118									
BT 0614 Alticinae sp 118									
BT 0615 Alticinae sp 118									
BT 0616 Alticinae sp 118									
BT 0617 Alticinae sp 118									
BT 0618 Alticinae sp 118									
BT 0619 Alticinae sp 118									
BT 0620 Alticinae sp 118									
BT 0621 Alticinae sp 118									
BT 0622 Alticinae sp 118									
BT 0623 Alticinae sp 118									
BT 0624 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 0625 Cassidinae sp 003	X		X	X	X	X	X	X	X
BT 0626 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 0627 Alticinae sp 118									
BT 0628 Cassidinae sp 012	X	X	X	X	X	X	X	X	X
BT 0629 Cassidinae sp 012									
BT 0630 Alticinae sp 036	X	X	X	X	X	X	X	X	X
BT 0631 Alticinae sp 064	X	X	X	X	X	X	X	X	X
BT 0632 Alticinae sp 086	X	X	X	X	X	X	X	X	X
BT 0633 Alticinae sp 085	X	X	X	X	X	X	X	X	X
BT 0634 Alticinae sp 066	X	X	X	X	X	X	X	X	X
BT 0635 Alticinae sp 096	X	X	X	X	X	X	X	X	X

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT_0636_Alticinae_sp_096				X			X		
BT_0637_Alticinae_sp_096				X			X		
BT_0638_Alticinae_sp_084	X		X	X		X	X		
BT_0639_Alticinae_sp_096	X		X	X		X	X		
BT_0640_Alticinae_sp_018	X	X	X	X	X	X	X	X	X
BT_0641_Alticinae_sp_013	X	X	X	X	X	X	X	X	X
BT_0642_Eumolpinae_sp_042	X	X	X	X	X	X	X	X	X
BT_0643_Eumolpinae_sp_042				X			X		
BT_0644_Galerucinae_sp_066	X	X	X	X	X	X	X	X	X
BT_0645_Galerucinae_sp_011	X	X	X	X	X	X	X	X	X
BT_0646_Alticinae_sp_071	X	X	X	X	X	X	X	X	X
BT_0647_Galerucinae_sp_035	X	X	X	X	X	X	X	X	X
BT_0648_Alticinae_sp_140	X	X	X	X	X	X	X	X	X
BT_0649_Alticinae_sp_140				X			X		
BT_0650_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0651_Eumolpinae_sp_039				X			X		
BT_0652_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_0653_Alticinae_sp_104				X			X		
BT_0654_Alticinae_sp_104				X			X		
BT_0655_Alticinae_sp_257	X	X	X	X	X	X	X	X	X
BT_0656_Alticinae_sp_051	X	X	X	X	X	X	X	X	X
BT_0656a_Alticinae_sp_051				X			X		
BT_0657_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0658_Eumolpinae_sp_039				X			X		
BT_0659_Eumolpinae_sp_039				X			X		
BT_0660_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_0661_Alticinae_sp_104				X			X		
BT_0662_Alticinae_sp_126	X	X	X	X	X	X	X	X	X
BT_0663_Alticinae_sp_150	X	X	X	X	X	X	X	X	X
BT_0664_Alticinae_sp_150	X	X	X	X	X	X	X	X	X
BT_0665_Alticinae_sp_150	X	X	X	X	X	X	X	X	X
BT_0666_Alticinae_sp_150				X			X		
BT_0667_Alticinae_sp_150				X			X		
BT_0668_Alticinae_sp_150				X			X		
BT_0669_Alticinae_sp_150				X			X		
BT_0670_Alticinae_sp_150				X			X		
BT_0671_Alticinae_sp_150				X			X		
BT_0672_Alticinae_sp_150				X			X		
BT_0673_Alticinae_sp_150				X			X		
BT_0674_Alticinae_sp_150				X			X		
BT_0675_Alticinae_sp_008	X	X	X	X	X	X	X	X	X
BT_0676_Alticinae_sp_004	X	X	X	X	X	X	X	X	X
BT_0677_Eumolpinae_sp_039	X	X	X	X	X	X	X	X	X
BT_0678_Eumolpinae_sp_044	X	X	X	X	X	X	X	X	X
BT_0679_Eumolpinae_sp_001	X	X	X	X	X	X	X	X	X
BT_0680_Galerucinae_sp_076	X	X	X	X	X				X
BT_0681_Galerucinae_sp_076				X					
BT_0682_Galerucinae_sp_076				X					
BT_0683_Alticinae_sp_115	X	X	X	X	X				X
BT_0684_Alticinae_sp_115				X					
BT_0685_Alticinae_sp_115				X					
BT_0686_Eumolpinae_sp_023	X	X	X	X	X				X
BT_0687_Alticinae_sp_087	X	X	X	X	X				X
BT_0688_Criocerinae_sp_001	X	X	X	X	X				X
BT_0689_Criocerinae_sp_001				X					
BT_0690_Alticinae_sp_124	X	X	X	X	X				X
BT_0691_Galerucinae_sp_031	X	X	X	X	X				X
BT_0692_Alticinae_sp_014	X	X	X	X	X				X
BT_0693_Alticinae_sp_014				X					
BT_0694_Alticinae_sp_014				X					
BT_0695_Alticinae_sp_014				X					
BT_0696_Alticinae_sp_014				X					
BT_0697_Alticinae_sp_014				X					
BT_0698_Alticinae_sp_013	X	X	X	X	X				X
BT_0699_Alticinae_sp_083	X	X	X	X	X				X
BT_0700_Alticinae_sp_083				X					
BT_0701_Alticinae_sp_083				X					
BT_0705_Galerucinae_sp_008	X	X	X	X	X				X
BT_0706_Eumolpinae_sp_038				X					
BT_0708_Alticinae_sp_135	X	X							
BT_0709_Galerucinae_sp_064	X	X	X	X	X				X
BT_0710_Alticinae_sp_013	X	X	X	X	X				X
BT_0711_Galerucinae_sp_073	X	X	X	X	X				X
BT_0712_Galerucinae_sp_073				X					
BT_0713_Galerucinae_sp_073	X	X	X	X	X				X
BT_0714_Galerucinae_sp_042	X	X	X	X	X				X
BT_0715_Galerucinae_sp_076	X	X	X	X	X				X
BT_0716_Galerucinae_sp_011	X	X	X	X	X				X
BT_0717_Galerucinae_sp_002	X	X	X	X	X				X
BT_0718_Galerucinae_sp_002				X					
BT_0719_Eumolpinae_sp_043	X	X	X	X	X				X
BT_0720_Eumolpinae_sp_043				X					
BT_0721_Eumolpinae_sp_020	X	X	X	X	X				X
BT_0722_Cassidinae_sp_013	X	X	X	X	X				X
BT_0725_Alticinae_sp_243	X	X	X	X	X				X
BT_0726_Alticinae_sp_243				X					
BT_0727_Galerucinae_sp_073	X	X	X	X	X				X
BT_0728_Galerucinae_sp_064	X	X	X	X	X				X
BT_0729_Galerucinae_sp_015	X	X	X	X	X				X
BT_0730_Alticinae_sp_136	X	X	X	X	X				X
BT_0731_Alticinae_sp_004	X	X	X	X	X				X
BT_0732_Galerucinae_sp_004	X	X	X	X	X				X
BT_0733_Galerucinae_sp_097	X	X	X	X	X				X
BT_0734_Galerucinae_sp_039	X	X	X	X	X				X
BT_0735_Galerucinae_sp_038	X	X	X	X	X				X
BT_0736_Galerucinae_sp_026	X	X	X	X	X				X
BT_0737_Galerucinae_sp_026				X					
BT_0738_Eumolpinae_sp_004	X	X	X	X	X				X
BT_0739_Galerucinae_sp_018	X	X	X	X	X				X
BT_0740_Galerucinae_sp_076	X	X	X	X	X				X
BT_0741_Galerucinae_sp_002	X	X	X	X	X				X
BT_0742_Galerucinae_sp_019	X	X	X	X	X				X
BT_0743_Galerucinae_sp_014	X	X	X	X	X				X
BT_0744_Galerucinae_sp_007	X	X	X	X	X				X
BT_0745_Galerucinae_sp_007				X					
BT_0746_Galerucinae_sp_007				X					
BT_0747_Galerucinae_sp_038	X	X	X	X	X				X
BT_0748_Galerucinae_sp_038				X					
BT_0749_Galerucinae_sp_039	X	X	X	X	X				X
BT_0750_Cassidinae_sp_005	X	X	X	X	X				X
BT_0751_Galerucinae_sp_062	X	X	X	X	X				X
BT_0752_Alticinae_sp_039	X	X	X	X	X				X
BT_0753_Alticinae_sp_010	X	X	X	X	X				X
BT_0754_Eumolpinae_sp_008	X	X	X	X	X				X
BT_0755_Galerucinae_sp_036	X	X	X	X	X				X
BT_0756_Eumolpinae_sp_022	X	X	X	X	X				X
BT_0757_Eumolpinae_sp_022				X					

Continued on next page(s)

Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 0758 Eumolpinae sp 022				X					
BT 0759 Eumolpinae sp 038	X		X	X					
BT 0760 Eumolpinae sp 038				X					
BT 0761 Eumolpinae sp 038				X					
BT 0762 Eumolpinae sp 038				X					
BT 0763 Eumolpinae sp 038				X					
BT 0764 Eumolpinae sp 038				X					
BT 0765 Eumolpinae sp 038				X					
BT 0766 Alticinae sp 096	X	X	X	X	X				X
BT 0767 Alticinae sp 096				X					
BT 0768 Alticinae sp 096				X					
BT 0770 Eumolpinae sp 016	X	X	X	X	X				X
BT 0771 Eumolpinae sp 016				X					
BT 0772 Eumolpinae sp 016				X					
BT 0773 Eumolpinae sp 016				X					
BT 0774 Eumolpinae sp 016				X					
BT 0775 Eumolpinae sp 016				X					
BT 0776 Alticinae sp 018	X		X	X					
BT 0777 Alticinae sp 018				X					
BT 0778 Alticinae sp 018				X					
BT 0779 Alticinae sp 115	X	X	X	X	X				X
BT 0780 Alticinae sp 115				X					
BT 0781 Eumolpinae sp 020	X	X	X	X	X	X			X
BT 0782 Galerucinae sp 034	X	X	X	X	X	X			X
BT 0788 Alticinae sp 006	X	X	X	X	X	X			X
BT 0789 Alticinae sp 014	X	X	X	X	X	X			X
BT 0790 Alticinae sp 014				X					
BT 0791 Eumolpinae sp 016	X	X	X	X	X	X			X
BT 0792 Alticinae sp 115	X	X	X	X	X	X			X
BT 0793 Alticinae sp 115				X					
BT 0794 Alticinae sp 129	X	X	X	X	X	X			X
BT 0795 Alticinae sp 096	X	X	X	X	X	X			X
BT 0796 Alticinae sp 097	X	X	X	X	X	X			X
BT 0799 Cassidinae sp 014	X	X	X	X	X	X			X
BT 0800 Galerucinae sp 030	X	X	X	X	X	X			X
BT 0801 Cassidinae sp 006	X	X	X	X	X	X			X
BT 0802 Cassidinae sp 006				X					
BT 0803 Galerucinae sp 076	X	X	X	X	X	X			X
BT 0804 Alticinae sp 028	X	X	X	X	X	X			X
BT 0805 Alticinae sp 014	X	X	X	X	X	X			X
BT 0806 Alticinae sp 013	X	X	X	X	X	X			X
BT 0807 Alticinae sp 054	X	X	X	X	X	X			X
BT 0808 Alticinae sp 064	X	X	X	X	X	X			X
BT 0809 Alticinae sp 012	X	X	X	X	X	X			X
BT 0810 Galerucinae sp 046	X	X	X	X	X	X			X
BT 0811 Criocerinae sp 006	X	X	X	X	X	X			X
BT 0812 Criocerinae sp 006				X					
BT 0813 Criocerinae sp 004	X	X	X	X	X	X			X
BT 0814 Alticinae sp 086	X	X	X	X	X	X			X
BT 0815 Alticinae sp 032	X	X	X	X	X	X			X
BT 0816 Galerucinae sp 019	X	X	X	X	X	X			X
BT 0817 Alticinae sp 087	X	X	X	X	X	X			X
BT 0818 Galerucinae sp 061	X	X	X	X	X	X			X
BT 0819 Eumolpinae sp 074	X	X	X	X	X	X			X
BT 0820 Eumolpinae sp 038	X	X	X	X	X	X			X
BT 0821 Eumolpinae sp 038				X					
BT 0822 Galerucinae sp 033	X	X	X	X					
BT 0823 Galerucinae sp 034	X	X	X	X	X				X
BT 0824 Galerucinae sp 034				X					
BT 0825 Galerucinae sp 034				X					
BT 0826 Galerucinae sp 034				X					
BT 0827 Eumolpinae sp 020	X	X	X	X	X	X			X
BT 0828 Alticinae sp 118	X	X	X	X	X	X			X
BT 0829 Alticinae sp 115	X	X	X	X	X	X			X
BT 0830 Alticinae sp 115				X					
BT 0831 Alticinae sp 147	X	X	X	X	X				X
BT 0832 Alticinae sp 147				X					
BT 0835 Galerucinae sp 029	X	X	X	X	X	X			X
BT 0836 Alticinae sp 003	X	X	X	X	X	X			X
BT 0837 Alticinae sp 009	X	X	X	X	X	X			X
BT 0838 Alticinae sp 059	X	X	X	X	X	X			X
BT 0839 Alticinae sp 040	X	X	X	X	X	X			X
BT 0840 Alticinae sp 140	X	X	X	X	X	X			X
BT 0841 Alticinae sp 127	X	X	X	X	X	X			X
BT 0842 Alticinae sp 127				X					
BT 0843 Alticinae sp 136	X	X	X	X	X	X			X
BT 0844 Alticinae sp 138	X	X	X	X	X	X			X
BT 0847 Eumolpinae sp 005	X	X	X	X	X	X			X
BT 0848 Alticinae sp 045	X	X	X	X	X	X			X
BT 0849 Alticinae sp 045				X					
BT 0850 Alticinae sp 049	X	X	X	X	X	X			X
BT 0851 Alticinae sp 065	X	X	X	X	X	X			X
BT 0852 Alticinae sp 133	X	X	X	X	X	X			X
BT 0853 Alticinae sp 133				X					
BT 0854 Alticinae sp 133				X					
BT 0855 Alticinae sp 018	X	X	X	X	X				X
BT 0856 Alticinae sp 018				X					
BT 0857 Alticinae sp 018				X					
BT 0858 Alticinae sp 018				X					
BT 0859 Alticinae sp 018				X					
BT 0860 Alticinae sp 018				X					
BT 0861 Eumolpinae sp 020	X	X	X	X	X	X			X
BT 0862 Alticinae sp 115	X	X	X	X	X	X			X
BT 0863 Alticinae sp 115				X					
BT 0864 Alticinae sp 253	X	X	X	X	X	X			X
BT 0865 Alticinae sp 096	X	X	X	X	X	X			X
BT 0866 Alticinae sp 096				X					
BT 0867 Alticinae sp 096				X					
BT 0868 Alticinae sp 122	X	X	X	X	X				
BT 0869 Alticinae sp 122				X					
BT 0870 Alticinae sp 122				X					
BT 0871 Alticinae sp 086	X	X	X	X	X				X
BT 0872 Alticinae sp 086				X					
BT 0873 Alticinae sp 085	X	X	X	X	X				X
BT 0877 Cassidinae sp 005	X	X							
BT 0879 Eumolpinae sp 043	X	X							
BT 0880 Criocerinae sp 001	X	X							
BT 0892 Galerucinae sp 035	X	X							
BT 0893 Alticinae sp 083	X	X							
BT 0895 Cassidinae sp 006	X	X							
BT 0899 Chrysomelinae sp 001	X	X							
BT 0900 Alticinae sp 033	X	X							
BT 0902 Alticinae sp 062	X	X							
BT 0903 Alticinae sp 087	X	X							
BT 0904 Alticinae sp 046	X	X							
BT 0905 Galerucinae sp 060	X	X							
BT 0906 Galerucinae sp 046	X	X							

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT_0907_Galerucinae_sp_046	X	X							
BT_0908_Criocerinae_sp_005	X	X							
BT_0909_Criocerinae_sp_001	X	X							
BT_0924_Cryptocephalinae_sp_001	X	X							
BT_0925_Eumolpinae_sp_029	X	X							
BT_0926_Lamprosomatinae_sp_002	X								
BT_0930_Galerucinae_sp_046	X	X							
BT_0932_Alticinae_sp_015	X	X							
BT_0934_Alticinae_sp_116	X	X							
BT_0935_Alticinae_sp_082	X	X							
BT_0936_Eumolpinae_sp_026	X	X							
BT_0937_Alticinae_sp_101	X								
BT_0943_Galerucinae_sp_033	X								
BT_0949_Cassidinae_sp_002	X	X	X	X	X				X
BT_0950_Cassidinae_sp_010	X	X	X	X	X	X			X
BT_0951_Cassidinae_sp_011	X	X	X	X	X	X			X
BT_0952_Alticinae_sp_269	X	X	X	X	X	X			X
BT_0953_Galerucinae_sp_028	X	X	X	X	X	X			X
BT_0954_Galerucinae_sp_029	X	X	X	X	X	X			X
BT_0955_Galerucinae_sp_020	X	X	X	X	X	X			X
BT_0956_Alticinae_sp_011	X	X	X	X	X	X			X
BT_0957_Galerucinae_sp_036	X	X	X	X	X	X			X
BT_0958_Alticinae_sp_154	X	X	X	X	X	X			X
BT_0959_Galerucinae_sp_052	X	X	X	X	X	X			X
BT_0960_Galerucinae_sp_047	X	X	X	X	X	X			X
BT_0961_Alticinae_sp_077	X	X	X	X	X	X			X
BT_0962_Alticinae_sp_076	X	X	X	X	X	X			X
BT_0963_Eumolpinae_sp_033	X	X	X	X	X				X
BT_0964_Alticinae_sp_030	X	X	X	X	X	X			X
BT_0965_Criocerinae_sp_002	X	X	X	X	X				X
BT_0966_Criocerinae_sp_003	X	X	X	X	X				X
BT_0967_Eumolpinae_sp_046	X	X	X	X	X	X			X
BT_0968_Alticinae_sp_099	X	X	X	X	X	X			X
BT_0969_Alticinae_sp_099	X								
BT_0970_Alticinae_sp_023	X		X	X					
BT_0971_Alticinae_sp_194	X	X	X	X	X				X
BT_0972_Alticinae_sp_127	X	X	X	X	X	X			X
BT_0973_Alticinae_sp_097	X	X	X	X	X				X
BT_0979_Eumolpinae_sp_042	X	X							
BT_0980_Alticinae_sp_005	X	X							
BT_0981_Eumolpinae_sp_006	X	X							
BT_0982_Alticinae_sp_022	X	X							
BT_0983_Alticinae_sp_052	X	X							
BT_1021_Galerucinae_sp_027	X	X							
BT_1025_Eumolpinae_sp_038	X		X	X					
BT_1026_Eumolpinae_sp_038									
BT_1027_Eumolpinae_sp_038									
BT_1028_Alticinae_sp_052	X	X	X	X	X				
BT_1029_Alticinae_sp_052						X			
BT_1030_Alticinae_sp_240	X	X	X	X	X	X			X
BT_1031_Alticinae_sp_002	X	X	X	X	X	X			X
BT_1032_Alticinae_sp_002	X								
BT_1033_Alticinae_sp_149	X	X	X	X	X	X			X
BT_1034_Alticinae_sp_048	X	X	X	X	X	X			X
BT_1035_Alticinae_sp_150	X	X	X	X	X	X			X
BT_1036_Alticinae_sp_062	X	X	X	X	X	X			X
BT_1037_Alticinae_sp_062	X								
BT_1038_Alticinae_sp_065	X	X	X	X	X	X			X
BT_1039_Alticinae_sp_065									
BT_1040_Alticinae_sp_065									
BT_1041_Alticinae_sp_065									
BT_1042_Alticinae_sp_065									
BT_1043_Alticinae_sp_066	X	X	X	X	X	X			X
BT_1044_Alticinae_sp_081	X	X	X	X	X	X			
BT_1045_Alticinae_sp_086	X	X	X	X	X	X			
BT_1046_Alticinae_sp_086									
BT_1047_Alticinae_sp_086									
BT_1048_Alticinae_sp_086									
BT_1049_Alticinae_sp_086									
BT_1050_Alticinae_sp_086									
BT_1051_Alticinae_sp_086									
BT_1052_Alticinae_sp_086									
BT_1053_Alticinae_sp_086									
BT_1054_Alticinae_sp_086									
BT_1055_Alticinae_sp_085	X	X	X	X	X	X			X
BT_1056_Alticinae_sp_085									
BT_1057_Alticinae_sp_085									
BT_1058_Alticinae_sp_085									
BT_1059_Alticinae_sp_081	X	X	X	X	X	X			
BT_1060_Alticinae_sp_085	X	X	X	X	X	X			X
BT_1061_Galerucinae_sp_096	X	X	X	X	X	X			X
BT_1062_Galerucinae_sp_096									
BT_1064_Alticinae_sp_142	X	X	X	X	X	X	X	X	X
BT_1065_Alticinae_sp_020	X	X	X	X	X	X			X
BT_1066_Alticinae_sp_018	X		X	X					
BT_1067_Alticinae_sp_018									
BT_1068_Alticinae_sp_018	X		X	X	X				
BT_1069_Alticinae_sp_123	X	X	X	X	X	X			
BT_1070_Alticinae_sp_097	X	X	X	X	X	X			X
BT_1071_Alticinae_sp_096	X	X	X	X	X	X			X
BT_1072_Alticinae_sp_096									
BT_1073_Alticinae_sp_096									
BT_1074_Alticinae_sp_096									
BT_1075_Alticinae_sp_096									
BT_1076_Alticinae_sp_096									
BT_1079_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_1080_Eumolpinae_sp_074	X	X	X	X	X	X	X	X	X
BT_1081_Alticinae_sp_027	X	X	X	X	X	X	X	X	X
BT_1082_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_1083_Alticinae_sp_096	X	X	X	X	X	X	X	X	X
BT_1084_Alticinae_sp_096									
BT_1085_Alticinae_sp_061	X	X	X	X	X	X	X	X	X
BT_1086_Alticinae_sp_083	X	X	X	X	X	X	X	X	X
BT_1087_Alticinae_sp_109	X	X	X	X	X	X	X	X	X
BT_1088_Galerucinae_sp_031	X	X	X	X	X	X	X	X	X
BT_1089_Galerucinae_sp_031									
BT_1090_Alticinae_sp_019	X	X	X	X	X	X	X	X	X
BT_1091_Cassidinae_sp_012	X	X	X	X	X	X	X	X	X
BT_1092_Cassidinae_sp_003	X	X	X	X	X	X	X	X	X
BT_1093_Hispinae_sp_007	X	X	X	X	X	X	X	X	X
BT_1094_Eumolpinae_sp_038	X	X	X	X	X	X	X	X	X
BT_1095_Alticinae_sp_096	X	X	X	X	X	X	X	X	X
BT_1096_Galerucinae_sp_036	X	X	X	X	X	X	X	X	X
BT_1098_Alticinae_sp_097	X	X	X	X	X	X	X	X	X
BT_1101_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_1104_Alticinae_sp_140	X	X	X	X	X	X	X	X	X
BT_1105_Alticinae_sp_074	X	X	X	X	X	X	X	X	X

Continued on next page(s)

Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 1106 Galerucinae sp 064	X	X	X	X	X	X	X	X	X
BT 1107 Alticinae sp 099	X	X	X	X	X	X	X	X	X
BT 1108 Alticinae sp 092	X	X	X	X	X	X	X	X	X
BT 1109 Alticinae sp 041	X	X	X	X	X	X	X	X	X
BT 1110 Galerucinae sp 046	X	X	X	X	X	X	X	X	X
BT 1111 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 1112 Alticinae sp 112	X	X	X	X	X	X	X	X	X
BT 1114 Alticinae sp 041	X	X	X	X	X	X	X	X	X
BT 1116 Cassidinae sp 006	X	X	X	X	X	X	X	X	X
BT 1117 Alticinae sp 049	X	X	X	X	X	X	X	X	X
BT 1118 Alticinae sp 123	X	X	X	X	X	X	X	X	X
BT 1119 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 1120 Alticinae sp 181	X	X	X	X	X	X	X	X	X
BT 1121 Alticinae sp 124	X	X	X	X	X	X	X	X	X
BT 1122 Alticinae sp 265	X	X	X	X	X	X	X	X	X
BT 1124 Cassidinae sp 007	X	X	X	X	X	X	X	X	X
BT 1125 Galerucinae sp 059	X	X	X	X	X	X	X	X	X
BT 1126 Hispinae sp 003	X	X	X	X	X	X	X	X	X
BT 1127 Alticinae sp 066	X	X	X	X	X	X	X	X	X
BT 1128 Alticinae sp 018	X	X	X	X	X	X	X	X	X
BT 1129 Alticinae sp 123	X	X	X	X	X	X	X	X	X
BT 1130 Alticinae sp 140	X	X	X	X	X	X	X	X	X
BT 1131 Alticinae sp 081	X	X	X	X	X	X	X	X	X
BT 1132 Alticinae sp 086	X	X	X	X	X	X	X	X	X
BT 1133 Alticinae sp 086									
BT 1134 Alticinae sp 086									
BT 1135 Alticinae sp 086									
BT 1136 Alticinae sp 086									
BT 1137 Alticinae sp 086									
BT 1138 Alticinae sp 086									
BT 1142 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 1143 Hispinae sp 007	X	X	X	X	X	X	X	X	X
BT 1145 Cassidinae sp 003	X	X	X	X	X	X	X	X	X
BT 1146 Eumolpinae sp 014	X	X	X	X	X	X	X	X	X
BT 1147 Alticinae sp 152	X	X	X	X	X	X	X	X	X
BT 1148 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 1151 Alticinae sp 057	X	X	X	X	X	X	X	X	X
BT 1152 Alticinae sp 141	X	X	X	X	X	X	X	X	X
BT 1154 Alticinae sp 121	X	X	X	X	X	X	X	X	X
BT 1155 Alticinae sp 121	X	X	X	X	X	X	X	X	X
BT 1157 Alticinae sp 149	X	X	X	X	X	X	X	X	X
BT 1158 Alticinae sp 146	X	X	X	X	X	X	X	X	X
BT 1159 Alticinae sp 146	X	X	X	X	X	X	X	X	X
BT 1160 Alticinae sp 055	X	X	X	X	X	X	X	X	X
BT 1161 Alticinae sp 143	X	X	X	X	X	X	X	X	X
BT 1164 Alticinae sp 087	X	X	X	X	X	X	X	X	X
BT 1165 Alticinae sp 061	X	X	X	X	X	X	X	X	X
BT 1166 Alticinae sp 140	X	X	X	X	X	X	X	X	X
BT 1167 Alticinae sp 140									
BT 1168 Alticinae sp 140									
BT 1169 Alticinae sp 100	X	X	X	X	X	X	X	X	X
BT 1170 Alticinae sp 105	X	X	X	X	X	X	X	X	X
BT 1171 Alticinae sp 085	X	X	X	X	X	X	X	X	X
BT 1175 Alticinae sp 087	X	X	X	X	X	X	X	X	X
BT 1176 Alticinae sp 052	X	X	X	X	X	X	X	X	X
BT 1178 Alticinae sp 087	X	X	X	X	X	X	X	X	X
BT 1179 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 1180 Alticinae sp 118									
BT 1181 Alticinae sp 118									
BT 1182 Alticinae sp 118									
BT 1183 Alticinae sp 118									
BT 1184 Alticinae sp 118									
BT 1185 Alticinae sp 118									
BT 1186 Alticinae sp 118									
BT 1187 Alticinae sp 118									
BT 1188 Alticinae sp 118									
BT 1189 Alticinae sp 118									
BT 1190 Alticinae sp 118									
BT 1191 Alticinae sp 118									
BT 1192 Alticinae sp 118									
BT 1194 Alticinae sp 087	X	X	X	X	X	X	X	X	X
BT 1195 Alticinae sp 087									
BT 1196 Alticinae sp 108	X	X	X	X	X	X	X	X	X
BT 1197 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 1198 Alticinae sp 131	X	X	X	X	X	X	X	X	X
BT 1199 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 1200 Alticinae sp 118									
BT 1201 Alticinae sp 118									
BT 1202 Alticinae sp 118									
BT 1203 Alticinae sp 118									
BT 1204 Alticinae sp 111	X	X	X	X	X	X	X	X	X
BT 1205 Alticinae sp 052	X	X	X	X	X	X	X	X	X
BT 1206 Alticinae sp 052									
BT 1207 Alticinae sp 052									
BT 1208 Galerucinae sp 046	X	X	X	X	X	X	X	X	X
BT 1209 Galerucinae sp 046									
BT 1210 Alticinae sp 064	X	X	X	X	X	X	X	X	X
BT 1211 Alticinae sp 018	X	X	X	X	X	X	X	X	X
BT 1212 Alticinae sp 017	X	X	X	X	X	X	X	X	X
BT 1213 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 1214 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 1215 Alticinae sp 144	X	X	X	X	X	X	X	X	X
BT 1216 Alticinae sp 049	X	X	X	X	X	X	X	X	X
BT 1217 Alticinae sp 145	X	X	X	X	X	X	X	X	X
BT 1219 Alticinae sp 086	X	X	X	X	X	X	X	X	X
BT 1220 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 1221 Galerucinae sp 034									
BT 1222 Alticinae sp 052	X	X	X	X	X	X	X	X	X
BT 1223 Alticinae sp 028	X	X	X	X	X	X	X	X	X
BT 1224 Galerucinae sp 075	X	X	X	X	X	X	X	X	X
BT 1225 Eumolpinae sp 040	X	X	X	X	X	X	X	X	X
BT 1226 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 1227 Galerucinae sp 034									
BT 1228 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 1229 Alticinae sp 096									
BT 1230 Alticinae sp 115	X	X	X	X	X	X	X	X	X
BT 1233 Alticinae sp 052	X	X	X	X	X	X	X	X	X
BT 1234 Alticinae sp 052									
BT 1235 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 1236 Alticinae sp 096									
BT 1237 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 1238 Galerucinae sp 034									
BT 1239 Galerucinae sp 034									
BT 1240 Eumolpinae sp 041	X	X	X	X	X	X	X	X	X
BT 1242 Lamprosomatinae sp 003	X	X	X	X	X	X	X	X	X
BT 1244 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 1245 Alticinae sp 085	X	X	X	X	X	X	X	X	X

Continued on next page(s)

Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT-1248-Alticinae_sp_118	X		X	X		X	X		
BT-1249-Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT-1250-Alticinae_sp_104									
BT-1251-Alticinae_sp_131	X	X	X	X	X	X	X	X	X
BT-1252-Alticinae_sp_086	X	X	X	X	X	X	X	X	X
BT-1258-Galerucinae_sp_028	X	X	X	X	X	X	X	X	X
BT-1259-Galerucinae_sp_049	X	X	X	X	X	X	X	X	X
BT-1260-Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT-1261-Alticinae_sp_103	X	X	X	X	X	X	X	X	X
BT-1262-Alticinae_sp_141	X	X	X	X	X	X	X	X	X
BT-1263-Alticinae_sp_051	X	X	X	X	X	X	X	X	X
BT-1264-Alticinae_sp_078	X	X	X	X	X	X	X	X	X
BT-1265-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1266-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1267-Alticinae_sp_128	X	X	X	X	X	X	X	X	X
BT-1269-Alticinae_sp_009	X	X	X	X	X	X	X	X	X
BT-1270-Galerucinae_sp_028	X	X	X	X	X	X	X	X	X
BT-1271-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1272-Alticinae_sp_128	X	X	X	X	X	X	X	X	X
BT-1273-Galerucinae_sp_028	X	X	X	X	X	X	X	X	X
BT-1274-Alticinae_sp_042	X	X	X	X	X	X	X	X	X
BT-1275-Alticinae_sp_051	X	X	X	X	X	X	X	X	X
BT-1276-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1277-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1278-Alticinae_sp_143	X	X	X	X	X	X	X	X	X
BT-1279-Alticinae_sp_125	X	X	X	X	X	X	X	X	X
BT-1280-Alticinae_sp_069	X	X	X	X	X	X	X	X	X
BT-1281-Galerucinae_sp_047	X	X	X	X	X	X	X	X	X
BT-1282-Galerucinae_sp_052	X	X	X	X	X	X	X	X	X
BT-1283-Galerucinae_sp_006	X	X	X	X	X	X	X	X	X
BT-1284-Galerucinae_sp_063	X	X	X	X	X	X	X	X	X
BT-1285-Galerucinae_sp_047	X	X	X	X	X	X	X	X	X
BT-1286-Alticinae_sp_021	X	X	X	X	X	X	X	X	X
BT-1287-Alticinae_sp_067	X	X	X	X	X	X	X	X	X
BT-1288-Alticinae_sp_103	X	X	X	X	X	X	X	X	X
BT-1289-Alticinae_sp_102	X	X	X	X	X	X	X	X	X
BT-1293-Alticinae_sp_050	X	X	X	X	X	X	X	X	X
BT-1294-Galerucinae_sp_072	X	X	X	X	X	X	X	X	X
BT-1295-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1296-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1297-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1298-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1299-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1300-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1301-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1302-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1303-Alticinae_sp_057	X	X	X	X	X	X	X	X	X
BT-1304-Galerucinae_sp_047	X	X	X	X	X	X	X	X	X
BT-1305-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1306-Alticinae_sp_125	X	X	X	X	X	X	X	X	X
BT-1307-Eumolpinae_sp_016	X	X	X	X	X	X	X	X	X
BT-1308-Alticinae_sp_134	X	X	X	X	X	X	X	X	X
BT-1309-Alticinae_sp_118	X	X	X	X	X	X	X	X	X
BT-1310-Alticinae_sp_035	X	X	X	X	X	X	X	X	X
BT-1311-Alticinae_sp_069	X	X	X	X	X	X	X	X	X
BT-1312-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1313-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1314-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1315-Alticinae_sp_093	X	X	X	X	X	X	X	X	X
BT-1318-Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT-1319-Alticinae_sp_051	X	X	X	X	X	X	X	X	X
BT-1321-Galerucinae_sp_061	X	X	X	X	X	X	X	X	X
BT-1322-Alticinae_sp_019	X	X	X	X	X	X	X	X	X
BT-1323-Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT-1324-Alticinae_sp_092	X	X	X	X	X	X	X	X	X
BT-1325-Alticinae_sp_019	X	X	X	X	X	X	X	X	X
BT-1326-Eumolpinae_sp_019	X	X	X	X	X	X	X	X	X
BT-1327-Alticinae_sp_097	X	X	X	X	X	X	X	X	X
BT-1328-Alticinae_sp_106	X	X	X	X	X	X	X	X	X
BT-1329-Alticinae_sp_106	X	X	X	X	X	X	X	X	X
BT-1330-Alticinae_sp_140	X	X	X	X	X	X	X	X	X
BT-1331-Cassidinae_sp_004	X	X	X	X	X	X	X	X	X
BT-1332-Cassidinae_sp_004	X	X	X	X	X	X	X	X	X
BT-1334-Alticinae_sp_245	X	X	X	X	X	X	X	X	X
BT-1335-Alticinae_sp_118	X	X	X	X	X	X	X	X	X
BT-1336-Alticinae_sp_244	X	X	X	X	X	X	X	X	X
BT-1337-Lamprosomatinae_sp_001	X	X	X	X	X	X	X	X	X
BT-1339-Alticinae_sp_244	X	X	X	X	X	X	X	X	X
BT-1340-Galerucinae_sp_066	X	X	X	X	X	X	X	X	X
BT-1347-Alticinae_sp_092	X	X	X	X	X	X	X	X	X
BT-1349-Alticinae_sp_051	X	X	X	X	X	X	X	X	X
BT-1350-Alticinae_sp_131	X	X	X	X	X	X	X	X	X
BT-1756-Alticinae_sp_014	X	X	X	X	X	X	X	X	X
BT-1758-Alticinae_sp_014	X	X	X	X	X	X	X	X	X
BT-1773-Galerucinae_sp_030	X	X	X	X	X	X	X	X	X
BT-1775-Alticinae_sp_033	X	X	X	X	X	X	X	X	X
BT-1776-Galerucinae_sp_062	X	X	X	X	X	X	X	X	X
BT-1784-Lamprosomatinae_sp_001	X	X	X	X	X	X	X	X	X
BT-1785-Chrysomelinae_sp_003	X	X	X	X	X	X	X	X	X
BT-1789-Galerucinae_sp_022	X	X	X	X	X	X	X	X	X
BT-1794-Alticinae_sp_001	X	X	X	X	X	X	X	X	X
BT-1795-Chrysomelinae_sp_001	X	X	X	X	X	X	X	X	X
BT-1809-Alticinae_sp_001	X	X	X	X	X	X	X	X	X
BT-1810-Eumolpinae_sp_012	X	X	X	X	X	X	X	X	X
BT-1811-Galerucinae_sp_054	X	X	X	X	X	X	X	X	X
BT-1812-Alticinae_sp_002	X	X	X	X	X	X	X	X	X
BT-1813-Alticinae_sp_032	X	X	X	X	X	X	X	X	X
BT-1820-Alticinae_sp_142	X	X	X	X	X	X	X	X	X
BT-1821-Alticinae_sp_088	X	X	X	X	X	X	X	X	X
BT-1928-Galerucinae_sp_019	X	X	X	X	X	X	X	X	X
BT-1932-Eumolpinae_sp_042	X	X	X	X	X	X	X	X	X
BT-1934-Eumolpinae_sp_014	X	X	X	X	X	X	X	X	X
BT-1935-Alticinae_sp_074	X	X	X	X	X	X	X	X	X
BT-1936-Alticinae_sp_049	X	X	X	X	X	X	X	X	X
BT-1954-Galerucinae_sp_041	X	X	X	X	X	X	X	X	X
BT-1963-Alticinae_sp_037	X	X	X	X	X	X	X	X	X
BT-1964-Galerucinae_sp_022	X	X	X	X	X	X	X	X	X
BT-1970-Eumolpinae_sp_007	X	X	X	X	X	X	X	X	X
BT-1971-Eumolpinae_sp_015	X	X	X	X	X	X	X	X	X
BT-1972-Eumolpinae_sp_015	X	X	X	X	X	X	X	X	X
BT-1973-Eumolpinae_sp_015	X	X	X	X	X	X	X	X	X
BT-1974-Alticinae_sp_015	X	X	X	X	X	X	X	X	X
BT-1975-Alticinae_sp_015	X	X	X	X	X	X	X	X	X
BT-1976-Galerucinae_sp_102	X	X	X	X	X	X	X	X	X
BT-1977-Alticinae_sp_007	X	X	X	X	X	X	X	X	X
BT-1978-Eumolpinae_sp_040	X	X	X	X	X	X	X	X	X
BT-1981-Galerucinae_sp_065	X	X	X	X	X	X	X	X	X

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 1982 Alticinae sp 062	X	X							
BT 2050 Alticinae sp 123	X	X							
BT 2054 Alticinae sp 139	X	X							
BT 2055 Alticinae sp 145	X	X							
BT 2056 Alticinae sp 141	X	X							
BT 2059 Alticinae sp 142	X	X							
BT 2065 Alticinae sp 142	X	X							
BT 2067 Alticinae sp 142	X	X							
BT 2068 Alticinae sp 142	X	X							
BT 2073 Alticinae sp 022	X	X							
BT 2074 Galerucinae sp 034	X	X	X	X	X				X
BT 2075 Galerucinae sp 034	X	X	X	X	X				X
BT 2076 Alticinae sp 002	X	X	X	X	X				X
BT 2077 Galerucinae sp 007	X	X	X	X	X				X
BT 2078 Alticinae sp 265	X	X	X	X	X				X
BT 2079 Eumolpinae sp 038	X	X	X	X	X				X
BT 2080 Galerucinae sp 065	X	X	X	X	X				X
BT 2081 Eumolpinae sp 042	X	X	X	X	X				X
BT 2082 Alticinae sp 064	X	X	X	X	X				X
BT 2083 Alticinae sp 064	X	X	X	X	X				X
BT 2084 Alticinae sp 032	X	X	X	X	X				X
BT 2085 Eumolpinae sp 042	X	X	X	X	X				X
BT 2086 Eumolpinae sp 042	X	X	X	X	X				X
BT 2087 Alticinae sp 049	X	X	X	X	X				X
BT 2088 Alticinae sp 049	X	X	X	X	X				X
BT 2089 Alticinae sp 049	X	X	X	X	X				X
BT 2090 Alticinae sp 081	X	X	X	X	X				X
BT 2091 Alticinae sp 081	X	X	X	X	X				X
BT 2092 Alticinae sp 081	X	X	X	X	X				X
BT 2093 Alticinae sp 096	X	X	X	X	X				X
BT 2094 Alticinae sp 096	X	X	X	X	X				X
BT 2095 Alticinae sp 096	X	X	X	X	X				X
BT 2096 Alticinae sp 096	X	X	X	X	X				X
BT 2097 Alticinae sp 122	X	X	X	X	X				X
BT 2098 Alticinae sp 122	X	X	X	X	X				X
BT 2099 Alticinae sp 122	X	X	X	X	X				X
BT 2100 Alticinae sp 122	X	X	X	X	X				X
BT 2101 Alticinae sp 086	X	X	X	X	X				X
BT 2102 Alticinae sp 086	X	X	X	X	X				X
BT 2103 Alticinae sp 086	X	X	X	X	X				X
BT 2104 Alticinae sp 086	X	X	X	X	X				X
BT 2105 Alticinae sp 086	X	X	X	X	X				X
BT 2106 Alticinae sp 086	X	X	X	X	X				X
BT 2107 Alticinae sp 085	X	X	X	X	X				X
BT 2108 Alticinae sp 085	X	X	X	X	X				X
BT 2109 Alticinae sp 133	X	X	X	X	X				X
BT 2110 Alticinae sp 133	X	X	X	X	X				X
BT 2111 Alticinae sp 133	X	X	X	X	X				X
BT 2112 Alticinae sp 133	X	X	X	X	X				X
BT 2113 Alticinae sp 142	X	X	X	X	X				X
BT 2114 Alticinae sp 142	X	X	X	X	X				X
BT 2115 Alticinae sp 142	X	X	X	X	X				X
BT 2116 Alticinae sp 142	X	X	X	X	X				X
BT 2117 Alticinae sp 142	X	X	X	X	X				X
BT 2118 Alticinae sp 142	X	X	X	X	X				X
BT 2119 Eumolpinae sp 016	X	X	X	X	X				X
BT 2120 Eumolpinae sp 016	X	X	X	X	X				X
BT 2121 Eumolpinae sp 016	X	X	X	X	X				X
BT 2122 Eumolpinae sp 016	X	X	X	X	X				X
BT 2123 Alticinae sp 018	X	X	X	X	X				X
BT 2124 Alticinae sp 018	X	X	X	X	X				X
BT 2125 Alticinae sp 018	X	X	X	X	X				X
BT 2126 Alticinae sp 018	X	X	X	X	X				X
BT 2127 Alticinae sp 018	X	X	X	X	X				X
BT 2128 Alticinae sp 018	X	X	X	X	X				X
BT 2129 Alticinae sp 018	X	X	X	X	X				X
BT 2130 Alticinae sp 018	X	X	X	X	X				X
BT 2131 Alticinae sp 018	X	X	X	X	X				X
BT 2132 Alticinae sp 018	X	X	X	X	X				X
BT 2133 Alticinae sp 018	X	X	X	X	X				X
BT 2134 Alticinae sp 018	X	X	X	X	X				X
BT 2135 Alticinae sp 122	X	X	X	X	X				X
BT 2136 Alticinae sp 110	X	X	X	X	X				X
BT 2137 Alticinae sp 110	X	X	X	X	X				X
BT 2138 Alticinae sp 115	X	X	X	X	X				X
BT 2143 Eumolpinae sp 042	X	X	X	X	X				X
BT 2152 Eumolpinae sp 038	X	X	X	X	X				X
BT 2153 Eumolpinae sp 038	X	X	X	X	X				X
BT 2154 Galerucinae sp 017	X	X	X	X	X				X
BT 2155 Alticinae sp 115	X	X	X	X	X				X
BT 2156 Cassidinae sp 012	X	X	X	X	X				X
BT 2157 Eumolpinae sp 024	X	X	X	X	X				X
BT 2158 Alticinae sp 083	X	X	X	X	X				X
BT 2161 Cassidinae sp 012	X	X	X	X	X				X
BT 2162 Cassidinae sp 012	X	X	X	X	X				X
BT 2163 Cassidinae sp 012	X	X	X	X	X				X
BT 2164 Cassidinae sp 012	X	X	X	X	X				X
BT 2165 Cassidinae sp 012	X	X	X	X	X				X
BT 2166 Cassidinae sp 012	X	X	X	X	X				X
BT 2167 Cassidinae sp 012	X	X	X	X	X				X
BT 2168 Galerucinae sp 036	X	X	X	X	X				X
BT 2169 Galerucinae sp 033	X	X	X	X	X				X
BT 2170 Galerucinae sp 034	X	X	X	X	X				X
BT 2171 Galerucinae sp 034	X	X	X	X	X				X
BT 2172 Galerucinae sp 062	X	X	X	X	X				X
BT 2173 Eumolpinae sp 034	X	X	X	X	X				X
BT 2176 Galerucinae sp 043	X	X	X	X	X				X
BT 2177 Galerucinae sp 043	X	X	X	X	X				X
BT 2178 Galerucinae sp 043	X	X	X	X	X				X
BT 2179 Alticinae sp 011	X	X	X	X	X				X
BT 2180 Alticinae sp 075	X	X	X	X	X				X
BT 2181 Alticinae sp 251	X	X	X	X	X				X
BT 2182 Galerucinae sp 029	X	X	X	X	X				X
BT 2183 Galerucinae sp 029	X	X	X	X	X				X
BT 2184 Galerucinae sp 029	X	X	X	X	X				X
BT 2185 Galerucinae sp 029	X	X	X	X	X				X
BT 2186 Galerucinae sp 029	X	X	X	X	X				X
BT 2187 Galerucinae sp 029	X	X	X	X	X				X
BT 2188 Galerucinae sp 029	X	X	X	X	X				X
BT 2189 Hispinae sp 008	X	X	X	X	X				X
BT 2190 Hispinae sp 022	X	X	X	X	X				X
BT 2191 Criocerinae sp 005	X	X	X	X	X				X
BT 2192 Galerucinae sp 028	X	X	X	X	X				X
BT 2193 Alticinae sp 076	X	X	X	X	X				X
BT 2194 Alticinae sp 127	X	X	X	X	X				X
BT 2195 Alticinae sp 127	X	X	X	X	X				X
BT 2196 Alticinae sp 127	X	X	X	X	X				X

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 2197 Eumolpinae sp 047	X	X	X	X	X				X
BT 2198 Eumolpinae sp 047	X		X	X	X				
BT 2199 Eumolpinae sp 035	X		X	X	X				
BT 2203 Alticinae sp 016	X	X							
BT 2208 Alticinae sp 112	X	X	X	X	X				X
BT 2218 Galerucinae sp 012	X	X							
BT 2219 Eumolpinae sp 042	X	X							
BT 2220 Galerucinae sp 007	X	X							
BT 2221 Galerucinae sp 022	X	X							
BT 2222 Galerucinae sp 076	X	X							
BT 2229 Alticinae sp 063	X	X							
BT 2231 Alticinae sp 087	X	X							
BT 2232 Alticinae sp 087	X	X							
BT 2233 Alticinae sp 087	X	X							
BT 2310 Cassidinae sp 004	X	X	X	X	X				
BT 2311 Alticinae sp 074	X	X							
BT 2312 Alticinae sp 042	X	X	X	X	X				X
BT 2313 Eumolpinae sp 025	X		X	X					
BT 2314 Eumolpinae sp 030	X		X	X					
BT 2317 Alticinae sp 007	X	X	X	X	X				
BT 2353 Hispinae sp 006	X	X							
BT 2383 Galerucinae sp 059	X	X							
BT 2399 Alticinae sp 038	X	X							
BT 2405 Alticinae sp 042	X	X							
BT 2491 Alticinae sp 085	X	X	X	X	X	X	X	X	X
BT 2492 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 2495 Alticinae sp 243	X	X	X	X	X	X	X	X	X
BT 2496 Alticinae sp 140	X	X	X	X	X	X	X	X	X
BT 2498 Alticinae sp 052	X	X	X	X	X	X	X	X	X
BT 2499 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 2502 Alticinae sp 160	X	X	X	X	X	X	X	X	X
BT 2503 Alticinae sp 079	X	X	X	X	X	X	X	X	X
BT 2504 Galerucinae sp 098	X	X	X	X	X	X	X	X	X
BT 2505 Galerucinae sp 046	X	X	X	X	X	X	X	X	X
BT 2506 Alticinae sp 047	X	X	X	X	X	X	X	X	X
BT 2507 Alticinae sp 203	X	X	X	X	X	X	X	X	X
BT 2508 Galerucinae sp 034	X		X	X	X	X	X	X	X
BT 2509 Galerucinae sp 034	X		X	X	X	X	X	X	X
BT 2510 Hispinae sp 010	X								
BT 2511 Hispinae sp 010	X	X							
BT 2516 Galerucinae sp 009	X	X	X	X	X	X	X	X	X
BT 2517 Alticinae sp 201	X	X	X	X	X	X	X	X	X
BT 2518 Alticinae sp 070	X	X	X	X	X	X	X	X	X
BT 2519 Alticinae sp 153	X	X	X	X	X	X	X	X	X
BT 2520 Alticinae sp 073	X	X	X	X	X	X	X	X	X
BT 2521 Galerucinae sp 082	X	X	X	X	X	X	X	X	X
BT 2522 Alticinae sp 051	X	X	X	X	X	X	X	X	X
BT 2523 Alticinae sp 153	X	X	X	X	X	X	X	X	X
BT 2524 Alticinae sp 153	X		X	X	X	X	X	X	X
BT 2525 Alticinae sp 153	X		X	X	X	X	X	X	X
BT 2526 Alticinae sp 153	X		X	X	X	X	X	X	X
BT 2528 Eumolpinae sp 045	X		X	X	X	X	X	X	X
BT 2529 Galerucinae sp 055	X	X	X	X	X	X	X	X	X
BT 2531 Alticinae sp 008	X	X	X	X	X	X	X	X	X
BT 2544 Alticinae sp 097	X	X	X	X	X	X	X	X	X
BT 2546 Cassidinae sp 004	X	X	X	X	X	X	X	X	X
BT 2547 Cassidinae sp 004	X		X	X	X	X	X	X	X
BT 2548 Cassidinae sp 012	X	X	X	X	X	X	X	X	X
BT 2549 Alticinae sp 094	X	X	X	X	X	X	X	X	X
BT 2550 Alticinae sp 056	X	X	X	X	X	X	X	X	X
BT 2551 Alticinae sp 246	X	X	X	X	X	X	X	X	X
BT 2552 Alticinae sp 246	X		X	X	X	X	X	X	X
BT 2557 Galerucinae sp 007	X	X							
BT 2572 Alticinae sp 018	X	X	X	X	X	X	X	X	X
BT 2573 Cassidinae sp 012	X	X	X	X	X	X	X	X	X
BT 2574 Cassidinae sp 012	X		X	X	X	X	X	X	X
BT 2575 Eumolpinae sp 031	X	X	X	X	X	X	X	X	X
BT 2576 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 2578 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 2579 Alticinae sp 115	X	X	X	X	X	X	X	X	X
BT 2594 Galerucinae sp 002	X	X							
BT 2600 Alticinae sp 008	X	X	X	X	X	X	X	X	X
BT 2601 Galerucinae sp 007	X	X	X	X	X	X	X	X	X
BT 2602 Alticinae sp 085	X	X	X	X	X	X	X	X	X
BT 2603 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 2604 Alticinae sp 106	X	X	X	X	X	X	X	X	X
BT 2605 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 2606 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2607 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2608 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2609 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2610 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2611 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2612 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2613 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2614 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2615 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2616 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2617 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2618 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2619 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2620 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2621 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2622 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2623 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2624 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2625 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2626 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2627 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2628 Alticinae sp 118	X		X	X	X	X	X	X	X
BT 2629 Eumolpinae sp 029	X	X	X	X	X	X	X	X	X
BT 2631 Alticinae sp 135	X	X	X	X	X	X	X	X	X
BT 2632 Alticinae sp 143	X	X	X	X	X	X	X	X	X
BT 2634 Alticinae sp 196	X	X	X	X	X	X	X	X	X
BT 2637 Eumolpinae sp 040	X	X	X	X	X	X	X	X	X
BT 2638 Galerucinae sp 061	X	X	X	X	X	X	X	X	X
BT 2639 Alticinae sp 061	X	X	X	X	X	X	X	X	X
BT 2640 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 2641 Alticinae sp 097	X	X	X	X	X	X	X	X	X
BT 2642 Eumolpinae sp 019	X	X	X	X	X	X	X	X	X
BT 2643 Alticinae sp 042	X	X	X	X	X	X	X	X	X
BT 2644 Cricocerinae sp 001	X	X	X	X	X	X	X	X	X
BT 2645 Eumolpinae sp 038	X	X	X	X	X	X	X	X	X
BT 2646 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 2657 Alticinae sp 001	X	X	X	X	X	X	X	X	X
BT 2658 Alticinae sp 041	X	X	X	X	X	X	X	X	X
BT 2659 Alticinae sp 086	X	X	X	X	X	X	X	X	X

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 2660 Alticinae sp 097	X	X	X	X	X	X	X	X	
BT 2661 Hispinae sp 009	X	X	X	X	X	X	X	X	X
BT 2662 Alticinae sp 087	X	X	X	X	X	X	X	X	X
BT 2663 Hispinae sp 005	X	X	X	X	X	X	X	X	X
BT 2664 Cassidinae sp 003	X	X	X	X	X	X	X	X	
BT 2665 Criocerinae sp 004	X	X	X	X	X	X	X	X	X
BT 2666 Alticinae sp 089	X	X	X	X	X	X	X	X	X
BT 2668 Eumolpinae sp 028	X	X	X	X	X	X	X	X	
BT 2670 Alticinae sp 051	X	X	X	X	X	X	X	X	X
BT 2671 Galerucinae sp 072	X	X	X	X	X	X	X	X	X
BT 2672 Alticinae sp 051	X	X	X	X	X	X	X	X	X
BT 2673 Galerucinae sp 049	X	X	X	X	X	X	X	X	X
BT 2675 Galerucinae sp 002	X	X	X	X	X	X	X	X	
BT 2676 Galerucinae sp 046	X	X	X	X	X	X	X	X	
BT 2677 Galerucinae sp 061	X	X	X	X	X	X	X	X	
BT 2678 Alticinae sp 111	X	X	X	X	X	X	X	X	
BT 2679 Alticinae sp 096	X	X	X	X	X	X	X	X	
BT 2681 Criocerinae sp 001	X	X	X	X	X	X	X	X	
BT 2684 Alticinae sp 049	X	X	X	X	X	X	X	X	
BT 2697 Alticinae sp 053	X	X	X	X	X	X	X	X	X
BT 2698 Eumolpinae sp 071	X	X	X	X	X	X	X	X	X
BT 2699 Alticinae sp 060	X	X	X	X	X	X	X	X	
BT 2701 Alticinae sp 056	X	X	X	X	X	X	X	X	
BT 2702 Alticinae sp 056	X	X	X	X	X	X	X	X	
BT 2703 Alticinae sp 056	X	X	X	X	X	X	X	X	
BT 2704 Alticinae sp 246	X	X	X	X	X	X	X	X	
BT 2705 Alticinae sp 094	X	X	X	X	X	X	X	X	X
BT 2706 Alticinae sp 120	X	X	X	X	X	X	X	X	
BT 2707 Alticinae sp 034	X	X	X	X	X	X	X	X	X
BT 2708 Alticinae sp 024	X	X	X	X	X	X	X	X	
BT 2709 Hispinae sp 024	X	X	X	X	X	X	X	X	
BT 2717 Alticinae sp 001	X	X	X	X	X	X	X	X	
BT 2718 Alticinae sp 096	X	X	X	X	X	X	X	X	
BT 2719 Criocerinae sp 001	X	X	X	X	X	X	X	X	
BT 2723 Alticinae sp 225	X	X	X	X	X	X	X	X	
BT 2771 Galerucinae sp 100	X	X	X	X	X	X	X	X	
BT 2787 Galerucinae sp 002	X	X	X	X	X	X	X	X	
BT 2795 Galerucinae sp 017	X	X	X	X	X	X	X	X	
BT 2805 Eumolpinae sp 006	X	X	X	X	X	X	X	X	
BT 2806 Alticinae sp 050	X	X	X	X	X	X	X	X	
BT 2807 Alticinae sp 103	X	X	X	X	X	X	X	X	
BT 2809 Alticinae sp 259	X	X	X	X	X	X	X	X	
BT 2810 Galerucinae sp 072	X	X	X	X	X	X	X	X	
BT 2812 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 2813 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 2814 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 2816 Alticinae sp 150	X	X	X	X	X	X	X	X	
BT 2817 Alticinae sp 150	X	X	X	X	X	X	X	X	
BT 2818 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 2819 Eumolpinae sp 030	X	X	X	X	X	X	X	X	
BT 2820 Eumolpinae sp 030	X	X	X	X	X	X	X	X	
BT 2821 Eumolpinae sp 074	X	X	X	X	X	X	X	X	
BT 2822 Galerucinae sp 048	X	X	X	X	X	X	X	X	
BT 2823 Alticinae sp 068	X	X	X	X	X	X	X	X	
BT 2824 Alticinae sp 129	X	X	X	X	X	X	X	X	
BT 2825 Alticinae sp 126	X	X	X	X	X	X	X	X	
BT 2826 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 2827 Eumolpinae sp 067	X	X	X	X	X	X	X	X	
BT 2828 Eumolpinae sp 029	X	X	X	X	X	X	X	X	
BT 2829 Eumolpinae sp 030	X	X	X	X	X	X	X	X	
BT 2830 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 2831 Cassidinae sp 015	X	X	X	X	X	X	X	X	
BT 2832 Alticinae sp 072	X	X	X	X	X	X	X	X	
BT 2833 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 2834 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2835 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2837 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2838 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2839 Eumolpinae sp 066	X	X	X	X	X	X	X	X	
BT 2840 Alticinae sp 052	X	X	X	X	X	X	X	X	
BT 2841 Galerucinae sp 096	X	X	X	X	X	X	X	X	
BT 2842 Galerucinae sp 098	X	X	X	X	X	X	X	X	
BT 2843 Hispinae sp 005	X	X	X	X	X	X	X	X	
BT 2844 Alticinae sp 126	X	X	X	X	X	X	X	X	
BT 2845 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2846 Alticinae sp 152	X	X	X	X	X	X	X	X	
BT 2847 Alticinae sp 152	X	X	X	X	X	X	X	X	
BT 2848 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2849 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2850 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2851 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2852 Alticinae sp 052	X	X	X	X	X	X	X	X	
BT 2853 Alticinae sp 052	X	X	X	X	X	X	X	X	
BT 2854 Eumolpinae sp 017	X	X	X	X	X	X	X	X	
BT 2855 Galerucinae sp 098	X	X	X	X	X	X	X	X	
BT 2856 Cryptocephalinae sp 002	X	X	X	X	X	X	X	X	
BT 2863 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 2864 Eumolpinae sp 039	X	X	X	X	X	X	X	X	
BT 2865 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2866 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2867 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2868 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2869 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2870 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 2871 Alticinae sp 213	X	X	X	X	X	X	X	X	
BT 2872 Alticinae sp 064	X	X	X	X	X	X	X	X	
BT 2873 Alticinae sp 053	X	X	X	X	X	X	X	X	
BT 2874 Alticinae sp 094	X	X	X	X	X	X	X	X	
BT 2876 Alticinae sp 087	X	X	X	X	X	X	X	X	
BT 2877 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 2878 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 2879 Alticinae sp 164	X	X	X	X	X	X	X	X	
BT 2880 Eumolpinae sp 060	X	X	X	X	X	X	X	X	
BT 2881 Alticinae sp 085	X	X	X	X	X	X	X	X	
BT 2900 Alticinae sp 245	X	X	X	X	X	X	X	X	
BT 2901 Alticinae sp 244	X	X	X	X	X	X	X	X	
BT 2902 Alticinae sp 208	X	X	X	X	X	X	X	X	
BT 2903 Eumolpinae sp 071	X	X	X	X	X	X	X	X	
BT 2905 Alticinae sp 245	X	X	X	X	X	X	X	X	
BT 2907 Alticinae sp 191	X	X	X	X	X	X	X	X	
BT 2908 Alticinae sp 258	X	X	X	X	X	X	X	X	
BT 2912 Galerucinae sp 014	X	X	X	X	X	X	X	X	
BT 2935 Galerucinae sp 106	X	X	X	X	X	X	X	X	
BT 2938 Alticinae sp 244	X	X	X	X	X	X	X	X	
BT 2939 Alticinae sp 235	X	X	X	X	X	X	X	X	
BT 2948 Galerucinae sp 014	X	X	X	X	X	X	X	X	
BT 2949 Hispinae sp 007	X	X	X	X	X	X	X	X	

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 2954 Hispinae sp 018	X								
BT 2969 Eumolpinae sp 013	X	X							
BT 2999 Eumolpinae sp 003	X	X							
BT 3026 Galerucinae sp 029	X	X	X	X	X	X	X	X	X
BT 3027 Alticinae sp 173	X	X	X	X	X	X	X	X	X
BT 3028 Alticinae sp 204	X	X	X	X	X	X	X	X	X
BT 3029 Eumolpinae sp 017	X	X	X	X	X	X	X	X	X
BT 3030 Alticinae sp 193	X	X	X	X	X	X	X	X	X
BT 3031 Eumolpinae sp 055	X	X	X	X	X	X	X	X	X
BT 3032 Galerucinae sp 106	X	X	X	X	X	X	X	X	X
BT 3033 Alticinae sp 254	X	X	X	X	X	X	X	X	X
BT 3034 Alticinae sp 181	X	X	X	X	X	X	X	X	X
BT 3035 Galerucinae sp 051	X	X	X	X	X	X	X	X	X
BT 3036 Alticinae sp 042	X	X	X	X	X	X	X	X	X
BT 3037 Alticinae sp 135	X	X	X	X	X	X	X	X	X
BT 3038 Galerucinae sp 084	X	X	X	X	X	X	X	X	X
BT 3039 Eumolpinae sp 040	X	X	X	X	X	X	X	X	X
BT 3040 Eumolpinae sp 040				X					
BT 3041 Cassidinae sp 017	X	X	X	X	X	X	X	X	X
BT 3060 Alticinae sp 115	X	X	X	X	X	X	X	X	X
BT 3061 Alticinae sp 115				X					
BT 3062 Alticinae sp 249	X	X	X	X	X	X	X	X	X
BT 3063 Alticinae sp 249				X					
BT 3064 Alticinae sp 253	X	X	X	X	X	X	X	X	X
BT 3065 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 3066 Alticinae sp 147	X	X	X	X	X	X	X	X	X
BT 3067 Alticinae sp 147				X					
BT 3068 Alticinae sp 074	X		X	X		X	X		
BT 3070 Eumolpinae sp 038	X	X	X	X	X	X	X		X
BT 3071 Eumolpinae sp 038				X					
BT 3072 Eumolpinae sp 038				X					
BT 3073 Eumolpinae sp 038				X					
BT 3074 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 3075 Alticinae sp 096				X					
BT 3076 Criocerinae sp 001	X	X	X	X	X	X	X	X	X
BT 3077 Alticinae sp 064	X	X	X	X	X	X	X	X	X
BT 3079 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 3080 Alticinae sp 112	X	X	X	X	X	X	X	X	X
BT 3081 Alticinae sp 205	X	X	X	X	X	X	X	X	X
BT 3082 Galerucinae sp 012	X	X	X	X	X	X	X	X	X
BT 3083 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 3084 Alticinae sp 062	X	X	X	X	X	X	X	X	X
BT 3085 Alticinae sp 064	X	X	X	X	X	X	X	X	X
BT 3086 Alticinae sp 115	X	X	X	X	X	X	X	X	X
BT 3087 Eumolpinae sp 038	X	X	X	X	X	X	X	X	X
BT 3088 Alticinae sp 061	X	X	X	X	X	X	X	X	X
BT 3089 Eumolpinae sp 034	X	X	X	X	X	X	X	X	X
BT 3090 Eumolpinae sp 065	X	X	X	X	X	X	X	X	X
BT 3091 Eumolpinae sp 065				X					
BT 3092 Eumolpinae sp 002	X	X	X	X	X	X	X	X	X
BT 3093 Alticinae sp 051	X	X	X	X	X	X	X	X	X
BT 3094 Alticinae sp 051				X					
BT 3095 Alticinae sp 254	X	X	X	X	X	X	X	X	X
BT 3097 Eumolpinae sp 038	X	X	X	X	X	X	X	X	X
BT 3098 Eumolpinae sp 038				X					
BT 3099 Eumolpinae sp 038				X					
BT 3100 Alticinae sp 172	X	X	X	X	X	X	X	X	X
BT 3101 Alticinae sp 245	X	X	X	X	X	X	X	X	X
BT 3102 Alticinae sp 244	X	X	X	X	X	X	X	X	X
BT 3103 Alticinae sp 121	X	X	X	X	X	X	X	X	X
BT 3104 Alticinae sp 244	X	X	X	X	X	X	X	X	X
BT 3105 Alticinae sp 244				X					
BT 3106 Alticinae sp 171	X	X	X	X	X	X	X	X	X
BT 3107 Galerucinae sp 066	X	X	X	X	X	X	X	X	X
BT 3108 Alticinae sp 001	X	X	X	X	X	X	X	X	X
BT 3109 Galerucinae sp 109	X	X	X	X	X	X	X	X	X
BT 3110 Galerucinae sp 048	X	X	X	X	X	X	X	X	X
BT 3111 Alticinae sp 065	X	X	X	X	X	X	X	X	X
BT 3112 Alticinae sp 197	X	X	X	X	X	X	X	X	X
BT 3113 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 3114 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 3115 Galerucinae sp 098	X	X	X	X	X	X	X	X	X
BT 3116 Galerucinae sp 098				X					
BT 3117 Alticinae sp 163	X	X	X	X	X	X	X	X	X
BT 3118 Alticinae sp 178	X	X	X	X	X	X	X	X	X
BT 3120 Galerucinae sp 098				X					
BT 3121 Hispinae sp 025	X	X	X	X	X	X	X	X	X
BT 3122 Alticinae sp 005	X		X	X		X	X		
BT 3123 Eumolpinae sp 065	X		X	X		X	X		
BT 3124 Alticinae sp 087	X	X	X	X	X	X	X	X	X
BT 3125 Alticinae sp 065	X		X	X		X	X		
BT 3126 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 3127 Eumolpinae sp 039				X					
BT 3128 Eumolpinae sp 039				X					
BT 3143 Galerucinae sp 017	X	X							
BT 3157 Galerucinae sp 024	X	X							
BT 3177 Galerucinae sp 017	X	X							
BT 3191 Galerucinae sp 017	X	X							
BT 3197 Alticinae sp 228	X		X	X		X	X		
BT 3198 Alticinae sp 119	X		X	X		X	X		
BT 3200 Galerucinae sp 073	X	X	X	X	X	X	X	X	X
BT 3201 Alticinae sp 255	X	X	X	X	X	X	X	X	X
BT 3202 Alticinae sp 184	X	X	X	X	X	X	X	X	X
BT 3203 Galerucinae sp 073	X	X	X	X	X	X	X	X	X
BT 3204 Alticinae sp 172	X	X	X	X	X	X	X	X	X
BT 3205 Eumolpinae sp 052	X	X	X	X	X	X	X	X	X
BT 3206 Alticinae sp 245	X	X	X	X	X	X	X	X	X
BT 3207 Alticinae sp 197	X		X	X		X	X		
BT 3208 Alticinae sp 197				X					
BT 3209 Alticinae sp 123	X	X	X	X	X	X	X	X	X
BT 3210 Galerucinae sp 066	X	X	X	X	X	X	X	X	X
BT 3211 Alticinae sp 052	X	X	X	X	X	X	X	X	X
BT 3212 Criocerinae sp 001	X	X	X	X	X	X	X	X	X
BT 3213 Eumolpinae sp 040	X	X	X	X	X	X	X	X	X
BT 3214 Eumolpinae sp 014	X		X	X		X	X		
BT 3215 Eumolpinae sp 014				X					
BT 3216 Galerucinae sp 104	X	X	X	X	X	X	X	X	X
BT 3217 Alticinae sp 087	X	X	X	X	X	X	X	X	X
BT 3218 Alticinae sp 087				X					
BT 3219 Alticinae sp 087				X					
BT 3220 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 3221 Galerucinae sp 034				X					
BT 3222 Galerucinae sp 034				X					
BT 3223 Galerucinae sp 034				X					
BT 3224 Galerucinae sp 034				X					
BT 3225 Galerucinae sp 025	X	X	X	X	X	X	X	X	X
BT 3226 Galerucinae sp 025	X	X	X	X	X	X	X	X	X

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 3227 Galerucinae sp 025				X			X		
BT 3228 Galerucinae sp 025				X			X		
BT 3229 Galerucinae sp 025				X			X		
BT 3230 Alticinae sp 052	X	X	X	X	X	X	X	X	
BT 3231 Alticinae sp 052				X			X		
BT 3232 Alticinae sp 064	X	X	X	X	X	X	X	X	
BT 3233 Alticinae sp 097	X	X	X	X	X	X	X	X	
BT 3234 Eumolpinae sp 038	X		X	X		X	X		
BT 3235 Alticinae sp 018	X	X	X	X	X	X	X	X	
BT 3236 Galerucinae sp 034	X	X	X	X	X	X	X	X	
BT 3237 Galerucinae sp 034				X			X		
BT 3238 Galerucinae sp 034				X			X		
BT 3239 Galerucinae sp 034				X			X		
BT 3241 Alticinae sp 097	X		X	X		X	X		
BT 3242 Galerucinae sp 025	X	X	X	X	X	X	X	X	
BT 3243 Alticinae sp 240	X	X	X	X	X	X	X	X	
BT 3244 Alticinae sp 052	X	X	X	X	X	X	X	X	
BT 3248 Alticinae sp 116	X	X	X	X	X	X	X	X	
BT 3249 Galerucinae sp 034	X	X	X	X	X	X	X	X	
BT 3250 Galerucinae sp 034				X			X		
BT 3251 Galerucinae sp 034				X			X		
BT 3252 Galerucinae sp 034				X			X		
BT 3253 Alticinae sp 115	X	X	X	X	X	X	X	X	
BT 3254 Alticinae sp 115				X		X	X		
BT 3255 Galerucinae sp 061	X	X	X	X	X	X	X	X	
BT 3256 Galerucinae sp 092	X	X	X	X	X	X	X	X	
BT 3257 Alticinae sp 212	X	X	X	X	X	X	X	X	
BT 3258 Galerucinae sp 066	X	X	X	X	X	X	X	X	
BT 3259 Galerucinae sp 066				X			X		
BT 3260 Alticinae sp 109	X	X	X	X	X	X	X	X	
BT 3261 Alticinae sp 029	X	X	X	X	X	X	X	X	
BT 3262 Alticinae sp 042	X	X	X	X	X	X	X	X	
BT 3263 Eumolpinae sp 040	X	X	X	X	X	X	X	X	
BT 3264 Galerucinae sp 026	X	X	X	X	X	X	X	X	
BT 3265 Eumolpinae sp 042	X	X	X	X	X	X	X	X	
BT 3266 Galerucinae sp 021	X	X	X	X	X	X	X	X	
BT 3266a Alticinae sp 112	X	X	X	X	X	X	X	X	
BT 3267 Galerucinae sp 098	X	X	X	X	X	X	X	X	
BT 3268 Alticinae sp 170	X	X	X	X	X	X	X	X	
BT 3269 Alticinae sp 111	X	X	X	X	X	X	X	X	
BT 3270 Alticinae sp 092	X	X	X	X	X	X	X	X	
BT 3271 Alticinae sp 202	X	X	X	X	X	X	X	X	
BT 3272 Alticinae sp 096	X	X	X	X	X	X	X	X	
BT 3273 Galerucinae sp 034	X	X	X	X	X	X	X	X	
BT 3274 Alticinae sp 087	X	X	X	X	X	X	X	X	
BT 3275 Alticinae sp 087				X			X		
BT 3276 Alticinae sp 087				X			X		
BT 3277 Eumolpinae sp 039	X	X	X	X	X	X	X	X	
BT 3278 Galerucinae sp 061	X	X	X	X	X	X	X	X	
BT 3279 Alticinae sp 061	X	X	X	X	X	X	X	X	
BT 3280 Alticinae sp 051	X		X	X		X	X		
BT 3281 Alticinae sp 051				X			X		
BT 3282 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3283 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3284 Alticinae sp 260	X	X	X	X	X	X	X	X	
BT 3299 Eumolpinae sp 039	X	X	X	X	X	X	X	X	
BT 3300 Eumolpinae sp 039				X			X		
BT 3301 Alticinae sp 260	X	X	X	X		X	X		
BT 3302 Alticinae sp 260	X	X	X	X		X	X		
BT 3303 Hispinae sp 005	X	X	X	X	X	X	X	X	
BT 3304 Alticinae sp 260	X	X	X	X	X	X	X	X	
BT 3305 Galerucinae sp 028	X	X	X	X	X	X	X	X	
BT 3306 Alticinae sp 254	X	X	X	X	X	X	X	X	
BT 3307 Alticinae sp 181	X	X	X	X	X	X	X	X	
BT 3308 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 3309 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 3310 Alticinae sp 063	X	X	X	X	X	X	X	X	
BT 3311 Galerucinae sp 015	X	X	X	X	X	X	X	X	
BT 3312 Eumolpinae sp 042	X	X	X	X	X	X	X	X	
BT 3313 Eumolpinae sp 055	X	X	X	X	X	X	X	X	
BT 3314 Eumolpinae sp 042	X	X	X	X	X	X	X	X	
BT 3316 Alticinae sp 117	X	X	X	X	X	X	X	X	
BT 3317 Eumolpinae sp 051	X	X	X	X	X	X	X	X	
BT 3318 Alticinae sp 210	X	X	X	X	X	X	X	X	
BT 3319 Alticinae sp 173	X	X	X	X	X	X	X	X	
BT 3320 Galerucinae sp 078	X	X	X	X	X	X	X	X	
BT 3321 Alticinae sp 035	X	X	X	X	X	X	X	X	
BT 3322 Eumolpinae sp 042	X	X	X	X	X	X	X	X	
BT 3323 Eumolpinae sp 042				X			X		
BT 3324 Eumolpinae sp 042				X			X		
BT 3325 Galerucinae sp 086	X	X	X	X	X	X	X	X	
BT 3326 Eumolpinae sp 017	X		X	X		X	X		
BT 3327 Eumolpinae sp 050	X	X	X	X	X	X	X	X	
BT 3328 Alticinae sp 210	X	X	X	X	X	X	X	X	
BT 3329 Alticinae sp 210				X			X		
BT 3330 Alticinae sp 210				X			X		
BT 3331 Alticinae sp 210				X			X		
BT 3332 Alticinae sp 035	X	X	X	X	X	X	X	X	
BT 3333 Alticinae sp 253	X	X	X	X	X	X	X	X	
BT 3334 Alticinae sp 052	X	X	X	X	X	X	X	X	
BT 3335 Galerucinae sp 045	X	X	X	X	X	X	X	X	
BT 3336 Galerucinae sp 045				X			X		
BT 3337 Eumolpinae sp 040	X	X	X	X	X	X	X	X	
BT 3338 Eumolpinae sp 050	X	X	X	X	X	X	X	X	
BT 3339 Alticinae sp 127	X	X	X	X	X	X	X	X	
BT 3340 Eumolpinae sp 053	X			X			X		
BT 3341 Alticinae sp 181	X	X	X	X	X	X	X	X	
BT 3342 Alticinae sp 169	X	X	X	X	X	X	X	X	
BT 3343 Alticinae sp 076	X	X	X	X	X	X	X	X	
BT 3344 Alticinae sp 076				X			X		
BT 3345 Alticinae sp 076	X	X	X	X	X	X	X	X	
BT 3346 Cassidinae sp 016	X	X	X	X	X	X	X	X	
BT 3347 Galerucinae sp 108	X	X	X	X	X	X	X	X	
BT 3348 Cassidinae sp 018	X	X	X	X	X	X	X	X	
BT 3349 Eumolpinae sp 052	X			X			X		
BT 3350 Hispinae sp 014	X	X	X	X	X	X	X	X	
BT 3351 Alticinae sp 255	X	X	X	X	X	X	X	X	
BT 3352 Alticinae sp 236	X			X			X		
BT 3353 Eumolpinae sp 054	X			X			X		
BT 3354 Alticinae sp 172	X	X	X	X	X	X	X	X	
BT 3355 Eumolpinae sp 054	X			X			X		
BT 3356 Eumolpinae sp 054	X			X			X		
BT 3357 Alticinae sp 057	X	X	X	X	X	X	X	X	
BT 3358 Alticinae sp 057				X			X		
BT 3359 Galerucinae sp 074	X	X	X	X	X	X	X	X	
BT 3360 Galerucinae sp 089	X	X	X	X	X	X	X	X	
BT 3361 Alticinae sp 009	X	X	X	X	X	X	X	X	

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 3362 Eumolpinae sp 046	X	X	X	X	X	X	X	X	
BT 3363 Alticinae sp 106	X	X	X	X	X	X	X	X	
BT 3364 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3365 Eumolpinae sp 024				X	X	X	X	X	
BT 3366 Alticinae sp 152	X	X	X	X	X	X	X	X	
BT 3367 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 3368 Alticinae sp 051				X	X	X	X	X	
BT 3369 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 3370 Alticinae sp 051				X	X	X	X	X	
BT 3371 Alticinae sp 051				X	X	X	X	X	
BT 3372 Alticinae sp 051				X	X	X	X	X	
BT 3373 Eumolpinae sp 064	X	X	X	X	X	X	X	X	
BT 3374 Eumolpinae sp 072	X		X	X	X	X	X	X	
BT 3375 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3376 Eumolpinae sp 024				X	X	X	X	X	
BT 3377 Eumolpinae sp 024				X	X	X	X	X	
BT 3378 Eumolpinae sp 024				X	X	X	X	X	
BT 3379 Eumolpinae sp 062	X	X	X	X	X	X	X	X	
BT 3380 Galerucinae sp 074	X	X	X	X	X	X	X	X	
BT 3381 Galerucinae sp 074				X	X	X	X	X	
BT 3382 Galerucinae sp 055	X	X	X	X	X	X	X	X	
BT 3383 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3384 Alticinae sp 152	X	X	X	X	X	X	X	X	
BT 3385 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 3386 Hispinae sp 011	X	X	X	X	X	X	X	X	
BT 3387 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3388 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3389 Eumolpinae sp 064	X	X	X	X	X	X	X	X	
BT 3390 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 3391 Alticinae sp 052	X	X	X	X	X	X	X	X	
BT 3392 Alticinae sp 052				X	X	X	X	X	
BT 3393 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 3394 Alticinae sp 118				X	X	X	X	X	
BT 3395 Alticinae sp 118				X	X	X	X	X	
BT 3396 Alticinae sp 118				X	X	X	X	X	
BT 3398 Eumolpinae sp 039	X		X	X	X	X	X	X	
BT 3399 Eumolpinae sp 039				X	X	X	X	X	
BT 3400 Eumolpinae sp 039				X	X	X	X	X	
BT 3401 Eumolpinae sp 039				X	X	X	X	X	
BT 3402 Eumolpinae sp 039				X	X	X	X	X	
BT 3403 Eumolpinae sp 030	X	X	X	X	X	X	X	X	
BT 3405 Eumolpinae sp 039	X	X	X	X	X	X	X	X	
BT 3406 Eumolpinae sp 039				X	X	X	X	X	
BT 3407 Alticinae sp 118	X	X	X	X	X	X	X	X	
BT 3408 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 3409 Alticinae sp 113	X	X	X	X	X	X	X	X	
BT 3410 Alticinae sp 131	X	X	X	X	X	X	X	X	
BT 3411 Alticinae sp 118	X		X	X	X	X	X	X	
BT 3412 Eumolpinae sp 039	X		X	X	X	X	X	X	
BT 3413 Eumolpinae sp 039				X	X	X	X	X	
BT 3414 Eumolpinae sp 039				X	X	X	X	X	
BT 3415 Eumolpinae sp 039				X	X	X	X	X	
BT 3416 Alticinae sp 118	X		X	X	X	X	X	X	
BT 3417 Alticinae sp 118				X	X	X	X	X	
BT 3418 Alticinae sp 118	X		X	X	X	X	X	X	
BT 3419 Alticinae sp 118				X	X	X	X	X	
BT 3420 Alticinae sp 118				X	X	X	X	X	
BT 3421 Alticinae sp 118				X	X	X	X	X	
BT 3422 Alticinae sp 118				X	X	X	X	X	
BT 3423 Eumolpinae sp 029	X	X	X	X	X	X	X	X	
BT 3424 Alticinae sp 118	X		X	X	X	X	X	X	
BT 3425 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 3426 Galerucinae sp 105	X	X	X	X	X	X	X	X	
BT 3427 Alticinae sp 172	X	X	X	X	X	X	X	X	
BT 3428 Hispinae sp 025	X	X	X	X	X	X	X	X	
BT 3429 Galerucinae sp 066	X	X	X	X	X	X	X	X	
BT 3430 Alticinae sp 172	X	X	X	X	X	X	X	X	
BT 3431 Alticinae sp 248	X	X	X	X	X	X	X	X	
BT 3432 Alticinae sp 209	X	X	X	X	X	X	X	X	
BT 3433 Eumolpinae sp 071	X	X	X	X	X	X	X	X	
BT 3434 Alticinae sp 187	X	X	X	X	X	X	X	X	
BT 3435 Eumolpinae sp 019	X	X	X	X	X	X	X	X	
BT 3436 Galerucinae sp 061	X	X	X	X	X	X	X	X	
BT 3437 Galerucinae sp 061				X	X	X	X	X	
BT 3438 Galerucinae sp 031	X	X	X	X	X	X	X	X	
BT 3439 Alticinae sp 018	X		X	X	X	X	X	X	
BT 3440 Alticinae sp 117	X		X	X	X	X	X	X	
BT 3441 Alticinae sp 174	X	X	X	X	X	X	X	X	
BT 3442 Alticinae sp 207	X		X	X	X	X	X	X	
BT 3443 Alticinae sp 207				X	X	X	X	X	
BT 3444 Alticinae sp 064	X		X	X	X	X	X	X	
BT 3445 Alticinae sp 092	X	X	X	X	X	X	X	X	
BT 3446 Alticinae sp 083	X		X	X	X	X	X	X	
BT 3447 Alticinae sp 150	X	X	X	X	X	X	X	X	
BT 3448 Alticinae sp 117	X	X	X	X	X	X	X	X	
BT 3449 Alticinae sp 002	X	X	X	X	X	X	X	X	
BT 3450 Alticinae sp 087	X	X	X	X	X	X	X	X	
BT 3451 Alticinae sp 087				X	X	X	X	X	
BT 3452 Alticinae sp 065	X	X	X	X	X	X	X	X	
BT 3453 Galerucinae sp 061	X	X	X	X	X	X	X	X	
BT 3454 Galerucinae sp 061				X	X	X	X	X	
BT 3455 Galerucinae sp 031	X	X	X	X	X	X	X	X	
BT 3456 Alticinae sp 266	X	X	X	X	X	X	X	X	
BT 3457 Hispinae sp 003	X	X	X	X	X	X	X	X	
BT 3458 Eumolpinae sp 039	X	X	X	X	X	X	X	X	
BT 3459 Alticinae sp 150	X	X	X	X	X	X	X	X	
BT 3460 Alticinae sp 096	X	X	X	X	X	X	X	X	
BT 3461 Alticinae sp 162	X		X	X	X	X	X	X	
BT 3462 Chrysomelinae sp 001	X	X	X	X	X	X	X	X	
BT 3463 Alticinae sp 109	X	X	X	X	X	X	X	X	
BT 3464 Alticinae sp 071	X		X	X	X	X	X	X	
BT 3465 Alticinae sp 071	X	X	X	X	X	X	X	X	
BT 3466 Alticinae sp 190	X		X	X	X	X	X	X	
BT 3467 Alticinae sp 190				X	X	X	X	X	
BT 3468 Alticinae sp 197	X	X	X	X	X	X	X	X	
BT 3471 Galerucinae sp 002	X	X	X	X	X	X	X	X	
BT 3472 Galerucinae sp 002				X	X	X	X	X	
BT 3473 Alticinae sp 109	X	X	X	X	X	X	X	X	
BT 3474 Alticinae sp 197	X	X	X	X	X	X	X	X	
BT 3475 Galerucinae sp 104	X	X	X	X	X	X	X	X	
BT 3476 Eumolpinae sp 065	X		X	X	X	X	X	X	
BT 3477 Alticinae sp 198	X	X	X	X	X	X	X	X	
BT 3478 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 3479 Alticinae sp 065	X	X	X	X	X	X	X	X	
BT 3480 Galerucinae sp 061	X	X	X	X	X	X	X	X	
BT 3481 Alticinae sp 064	X	X	X	X	X	X	X	X	
BT 3482 Galerucinae sp 054	X		X	X	X	X	X	X	

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 3483 Alticinae sp 123	X		X	X		X	X		
BT 3484 Alticinae sp 123				X			X		
BT 3485 Alticinae sp 249	X	X	X	X	X	X	X	X	
BT 3486 Alticinae sp 198	X	X	X	X	X	X	X	X	
BT 3487 Alticinae sp 111	X		X	X		X	X		
BT 3488 Alticinae sp 176	X	X	X	X	X	X	X	X	
BT 3489 Galerucinae sp 066	X	X	X	X	X	X	X	X	
BT 3490 Galerucinae sp 066	X	X	X	X	X	X	X	X	
BT 3491 Galerucinae sp 066				X			X		
BT 3492 Galerucinae sp 066				X			X		
BT 3493 Alticinae sp 249	X	X	X	X	X	X	X	X	
BT 3494 Alticinae sp 249				X			X		
BT 3495 Cassidinae sp 012	X	X	X	X	X	X	X	X	
BT 3496 Alticinae sp 104	X	X	X	X	X	X	X	X	
BT 3497 Alticinae sp 061	X	X	X	X	X	X	X	X	
BT 3498 Alticinae sp 260	X			X			X		
BT 3499 Alticinae sp 005	X	X	X	X	X	X	X	X	
BT 3500 Galerucinae sp 066	X	X	X	X	X	X	X	X	
BT 3501 Galerucinae sp 094	X	X	X	X	X	X	X	X	
BT 3502 Eumolpinae sp 017	X		X	X		X	X		
BT 3503 Eumolpinae sp 039	X	X	X	X	X	X	X	X	
BT 3504 Alticinae sp 019	X	X	X	X	X	X	X	X	
BT 3505 Alticinae sp 109	X	X	X	X	X	X	X	X	
BT 3506 Alticinae sp 008	X	X	X	X	X	X	X	X	
BT 3507 Alticinae sp 254	X		X	X		X	X		
BT 3508 Alticinae sp 118	X		X	X		X	X		
BT 3509 Alticinae sp 073	X		X	X		X	X		
BT 3510 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3511 Galerucinae sp 079	X	X	X	X	X	X	X	X	
BT 3512 Galerucinae sp 079				X			X		
BT 3513 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 3514 Alticinae sp 051				X			X		
BT 3515 Alticinae sp 152	X	X	X	X	X	X	X	X	
BT 3516 Eumolpinae sp 024	X	X	X	X	X	X	X	X	
BT 3517 Galerucinae sp 048	X	X	X	X	X	X	X	X	
BT 3518 Galerucinae sp 058	X	X	X	X	X	X	X	X	
BT 3519 Alticinae sp 201	X		X	X		X	X		
BT 3520 Alticinae sp 181	X	X	X	X	X	X	X	X	
BT 3521 Alticinae sp 051	X	X	X	X	X	X	X	X	
BT 3522 Alticinae sp 181	X	X	X	X	X	X	X	X	
BT 3528 Galerucinae sp 093	X	X	X	X	X	X	X	X	
BT 3529 Galerucinae sp 091	X	X	X	X	X	X	X	X	
BT 3530 Alticinae sp 153	X	X	X	X	X	X	X	X	
BT 3531 Alticinae sp 166	X	X	X	X	X	X	X	X	
BT 3532 Alticinae sp 149	X	X	X	X	X	X	X	X	
BT 3533 Alticinae sp 247	X	X	X	X	X	X	X	X	
BT 3534 Alticinae sp 247				X			X		
BT 3535 Alticinae sp 172	X	X	X	X	X	X	X	X	
BT 3536 Alticinae sp 172	X	X	X	X	X	X	X	X	
BT 3537 Alticinae sp 172	X	X	X	X	X	X	X	X	
BT 3538 Alticinae sp 244	X	X	X	X	X	X	X	X	
BT 3539 Alticinae sp 241	X	X	X	X	X	X	X	X	
BT 3540 Eumolpinae sp 058	X	X	X	X	X	X	X	X	
BT 3542 Eumolpinae sp 049	X	X	X	X	X	X	X	X	
BT 3543 Eumolpinae sp 069	X	X	X	X	X	X	X	X	
BT 3544 Chrysomelinae sp 004	X	X	X	X	X	X	X	X	
BT 3545 Alticinae sp 184	X	X	X	X	X	X	X	X	
BT 3546 Eumolpinae sp 054	X		X	X		X	X		
BT 3547 Alticinae sp 214	X	X	X	X	X	X	X	X	
BT 3548 Eumolpinae sp 063	X	X	X	X	X	X	X	X	
BT 3549 Hispinae sp 012	X	X	X	X	X	X	X	X	
BT 3550 Galerucinae sp 093	X	X	X	X	X	X	X	X	
BT 3551 Galerucinae sp 093	X	X	X	X	X	X	X	X	
BT 3552 Alticinae sp 025	X	X	X	X	X	X	X	X	
BT 3553 Alticinae sp 166	X		X	X		X	X		
BT 3554 Alticinae sp 073	X		X	X		X	X		
BT 3555 Alticinae sp 181	X	X	X	X	X	X	X	X	
BT 3556 Eumolpinae sp 050	X	X	X	X	X	X	X	X	
BT 3557 Eumolpinae sp 070	X	X	X	X	X	X	X	X	
BT 3558 Hispinae sp 021	X	X	X	X	X	X	X	X	
BT 3559 Alticinae sp 233	X	X	X	X	X	X	X	X	
BT 3560 Alticinae sp 188	X	X	X	X	X	X	X	X	
BT 3561 Alticinae sp 056	X		X	X		X	X		
BT 3562 Eumolpinae sp 057	X	X	X	X	X	X	X	X	
BT 3563 Hispinae sp 014	X	X	X	X	X	X	X	X	
BT 3564 Alticinae sp 127	X	X	X	X	X	X	X	X	
BT 3565 Alticinae sp 127				X			X		
BT 3566 Alticinae sp 127				X			X		
BT 3567 Alticinae sp 127				X			X		
BT 3568 Alticinae sp 127				X			X		
BT 3569 Alticinae sp 127				X			X		
BT 3570 Alticinae sp 127				X			X		
BT 3571 Alticinae sp 127				X			X		
BT 3572 Alticinae sp 127				X			X		
BT 3573 Alticinae sp 127				X			X		
BT 3574 Alticinae sp 227	X	X	X	X	X	X	X	X	
BT 3575 Alticinae sp 227	X	X	X	X	X	X	X	X	
BT 3576 Alticinae sp 196	X		X	X		X	X		
BT 3577 Alticinae sp 196				X			X		
BT 3578 Alticinae sp 196				X			X		
BT 3579 Alticinae sp 196				X			X		
BT 3580 Alticinae sp 196				X			X		
BT 3581 Alticinae sp 179	X	X	X	X	X	X	X	X	
BT 3582 Galerucinae sp 106	X		X	X		X	X		
BT 3583 Alticinae sp 135	X	X	X	X	X	X	X	X	
BT 3584 Alticinae sp 167	X	X	X	X	X	X	X	X	
BT 3585 Alticinae sp 127	X	X	X	X	X	X	X	X	
BT 3586 Alticinae sp 127				X			X		
BT 3587 Alticinae sp 127				X			X		
BT 3588 Alticinae sp 127				X			X		
BT 3589 Alticinae sp 127				X			X		
BT 3590 Alticinae sp 127				X			X		
BT 3591 Alticinae sp 196	X	X	X	X	X	X	X	X	
BT 3592 Alticinae sp 196				X			X		
BT 3593 Alticinae sp 076	X	X	X	X	X	X	X	X	
BT 3594 Alticinae sp 181	X	X	X	X	X	X	X	X	
BT 3596 Alticinae sp 111	X	X	X	X	X	X	X	X	
BT 3597 Alticinae sp 196	X	X	X	X	X	X	X	X	
BT 3598 Alticinae sp 196				X			X		
BT 3599 Alticinae sp 196				X			X		
BT 3600 Alticinae sp 253	X		X	X		X	X		
BT 3601 Alticinae sp 127	X	X	X	X	X	X	X	X	
BT 3602 Alticinae sp 138	X		X	X		X	X		
BT 3603 Alticinae sp 182	X		X	X		X	X		
BT 3604 Alticinae sp 076	X	X	X	X	X	X	X	X	
BT 3605 Alticinae sp 076				X			X		
BT 3606 Alticinae sp 127	X		X	X		X	X		

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 3607 - Alticinae sp 138	X		X	X		X	X		
BT 3608 - Alticinae sp 253	X		X	X		X	X		
BT 3609 - Eumolpinae sp 047	X	X	X	X	X	X	X		X
BT 3610 - Alticinae sp 076	X	X	X	X	X	X	X		X
BT 3611 - Alticinae sp 181	X	X	X	X	X	X	X		X
BT 3612 - Eumolpinae sp 059	X	X	X	X	X	X	X		X
BT 3613 - Criocerinae sp 007	X	X	X	X	X	X	X		X
BT 3614 - Eumolpinae sp 059	X	X	X	X	X	X	X		X
BT 3615 - Galerucinae sp 029	X	X	X	X	X	X	X		X
BT 3616 - Galerucinae sp 029	X	X	X	X	X	X	X		X
BT 3617 - Alticinae sp 127	X	X	X	X	X	X	X		X
BT 3618 - Alticinae sp 127				X			X		
BT 3619 - Alticinae sp 127				X			X		
BT 3620 - Alticinae sp 127				X			X		
BT 3621 - Alticinae sp 127				X			X		
BT 3622 - Alticinae sp 254	X	X	X		X	X	X		X
BT 3623 - Alticinae sp 253	X	X	X	X	X	X	X		X
BT 3624 - Alticinae sp 127	X	X	X	X	X	X	X		X
BT 3625 - Alticinae sp 051	X	X	X	X	X	X	X		X
BT 3626 - Criocerinae sp 010	X	X	X	X	X	X	X		X
BT 3627 - Eumolpinae sp 039	X								
BT 3628 - Alticinae sp 104	X								
BT 3630 - Alticinae sp 117	X								
BT 3633 - Alticinae sp 109	X								
BT 3651 - Galerucinae sp 041	X	X							
BT 3652 - Alticinae sp 086	X								
BT 3653 - Alticinae sp 065	X								
BT 3656 - Alticinae sp 115	X								
BT 3658 - Alticinae sp 198	X								
BT 3659 - Alticinae sp 237	X								
BT 3660 - Alticinae sp 092	X								
BT 3685 - Alticinae sp 018	X	X	X	X	X	X	X		X
BT 3686 - Alticinae sp 177	X	X	X	X	X	X	X		X
BT 3687 - Alticinae sp 115	X	X	X	X	X	X	X		X
BT 3688 - Alticinae sp 087	X	X	X	X	X	X	X		X
BT 3689 - Eumolpinae sp 040	X	X	X	X	X	X	X		X
BT 3690 - Alticinae sp 115	X	X	X	X	X	X	X		X
BT 3691 - Alticinae sp 147	X	X	X	X	X	X	X		X
BT 3692 - Alticinae sp 147				X			X		
BT 3693 - Alticinae sp 123	X	X	X	X	X	X	X		X
BT 3694 - Alticinae sp 197	X	X	X	X	X	X	X		X
BT 3698 - Galerucinae sp 089	X								
BT 3699 - Eumolpinae sp 063	X	X	X	X	X	X	X		X
BT 3700 - Eumolpinae sp 024	X	X	X	X	X	X	X		X
BT 3701 - Alticinae sp 244	X	X	X	X	X	X	X		X
BT 3702 - Alticinae sp 206	X		X	X		X	X		
BT 3703 - Alticinae sp 206	X		X	X		X	X		
BT 3704 - Alticinae sp 172	X	X	X	X	X	X	X		X
BT 3705 - Alticinae sp 085	X								
BT 3706 - Alticinae sp 182	X								
BT 3707 - Alticinae sp 182	X								
BT 3708 - Eumolpinae sp 059	X								
BT 3710 - Alticinae sp 196	X								
BT 3711 - Alticinae sp 076	X								
BT 3716 - Alticinae sp 127	X								
BT 3719 - Alticinae sp 093	X								
BT 3721 - Alticinae sp 076	X								
BT 3723 - Eumolpinae sp 059	X								
BT 3724 - Alticinae sp 127	X								
BT 3725 - Alticinae sp 138	X								
BT 3726 - Alticinae sp 227	X	X	X	X	X	X	X		X
BT 3727 - Alticinae sp 253	X	X	X	X	X	X	X		X
BT 3728 - Alticinae sp 099	X	X	X	X	X	X	X		X
BT 3729 - Alticinae sp 104	X	X	X	X	X	X	X		X
BT 3730 - Alticinae sp 095	X	X	X	X	X	X	X		X
BT 3731 - Alticinae sp 127	X	X	X	X	X	X	X		X
BT 3732 - Alticinae sp 127				X			X		
BT 3733 - Alticinae sp 127				X			X		
BT 3734 - Alticinae sp 127				X			X		
BT 3735 - Alticinae sp 196	X	X	X		X	X	X		X
BT 3736 - Alticinae sp 196				X			X		
BT 3737 - Alticinae sp 196				X			X		
BT 3738 - Alticinae sp 196				X			X		
BT 3739 - Galerucinae sp 085	X	X	X	X	X	X	X		X
BT 3740 - Eumolpinae sp 047	X								
BT 3741 - Alticinae sp 138	X								
BT 3750 - Alticinae sp 076	X	X	X	X	X	X	X		X
BT 3751 - Alticinae sp 076	X								
BT 3752 - Eumolpinae sp 038	X	X	X	X	X	X	X		X
BT 3753 - Galerucinae sp 093	X	X	X	X	X	X	X		X
BT 3754 - Alticinae sp 076	X	X	X	X	X	X	X		X
BT 3755 - Alticinae sp 076	X	X	X	X	X	X	X		X
BT 3756 - Alticinae sp 127	X	X	X	X	X	X	X		X
BT 3757 - Alticinae sp 127				X			X		
BT 3758 - Alticinae sp 127				X			X		
BT 3759 - Alticinae sp 127				X			X		
BT 3760 - Alticinae sp 127				X			X		
BT 3761 - Alticinae sp 196	X	X	X	X	X	X	X		X
BT 3762 - Alticinae sp 127	X	X	X	X	X	X	X		X
BT 3763 - Alticinae sp 127				X			X		
BT 3765 - Alticinae sp 138	X								
BT 3767 - Alticinae sp 226	X								
BT 3768 - Criocerinae sp 008	X								
BT 3771 - Alticinae sp 169	X								
BT 3773 - Alticinae sp 127	X								
BT 3790 - Alticinae sp 196	X								
BT 3791 - Alticinae sp 232	X								
BT 3792 - Alticinae sp 240	X								
BT 3793 - Alticinae sp 204	X								
BT 3795 - Hispinae sp 019	X								
BT 3796 - Galerucinae sp 007	X	X							
BT 3807 - Alticinae sp 185	X		X	X		X	X		X
BT 3808 - Alticinae sp 230	X		X	X		X	X		X
BT 3809 - Alticinae sp 234	X		X	X		X	X		X
BT 3810 - Alticinae sp 217	X	X	X	X	X	X	X		X
BT 3811 - Hispinae sp 017	X	X	X	X	X	X	X		X
BT 3812 - Alticinae sp 118	X	X	X	X	X	X	X		X
BT 3813 - Alticinae sp 109	X	X	X	X	X	X	X		X
BT 3814 - Alticinae sp 118	X	X	X	X	X	X	X		X
BT 3815 - Alticinae sp 118	X	X	X	X	X	X	X		X
BT 3816 - Alticinae sp 118				X			X		
BT 3817 - Alticinae sp 118				X			X		
BT 3818 - Alticinae sp 118				X			X		
BT 3819 - Alticinae sp 118				X			X		
BT 3820 - Alticinae sp 216	X	X	X	X	X	X	X		X
BT 3824 - Galerucinae sp 013	X	X	X	X	X	X	X		X
BT 3825 - Alticinae sp 050	X	X	X	X	X	X	X		X

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 3826 Alticinae sp 201	X	X	X	X	X	X	X	X	X
BT 3827 Galerucinae sp 090	X	X	X	X	X	X	X	X	X
BT 3828 Alticinae sp 211	X	X	X	X	X	X	X	X	X
BT 3829 Alticinae sp 181	X	X	X	X	X	X	X	X	X
BT 3830 Alticinae sp 180	X	X	X	X	X	X	X	X	X
BT 3832 Eumolpinae sp 050	X	X	X	X	X	X	X	X	X
BT 3833 Alticinae sp 231	X	X	X	X	X	X	X	X	X
BT 3834 Alticinae sp 244	X	X	X	X	X	X	X	X	X
BT 3845 Galerucinae sp 007	X	X	X	X	X	X	X	X	X
BT 3846 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 3847 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 3848 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 3849 Alticinae sp 175	X	X	X	X	X	X	X	X	X
BT 3850 Alticinae sp 140	X	X	X	X	X	X	X	X	X
BT 3851 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 3852 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 3853 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 3854 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 3855 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 3856 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 3857 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 3858 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 3859 Alticinae sp 118	X	X	X	X	X	X	X	X	X
BT 3860 Eumolpinae sp 030	X	X	X	X	X	X	X	X	X
BT 3861 Cassidinae sp 015	X	X	X	X	X	X	X	X	X
BT 3869 Cassidinae sp 012	X	X	X	X	X	X	X	X	X
BT 3870 Eumolpinae sp 002	X	X	X	X	X	X	X	X	X
BT 3871 Alticinae sp 083	X	X	X	X	X	X	X	X	X
BT 3872 Eumolpinae sp 040	X	X	X	X	X	X	X	X	X
BT 3873 Cassidinae sp 020	X	X	X	X	X	X	X	X	X
BT 3874 Eumolpinae sp 042	X	X	X	X	X	X	X	X	X
BT 3885 Alticinae sp 007	X	X	X	X	X	X	X	X	X
BT 3891 Galerucinae sp 090	X	X	X	X	X	X	X	X	X
BT 3892 Galerucinae sp 090	X	X	X	X	X	X	X	X	X
BT 3893 Alticinae sp 164	X	X	X	X	X	X	X	X	X
BT 3894 Alticinae sp 200	X	X	X	X	X	X	X	X	X
BT 3895 Galerucinae sp 079	X	X	X	X	X	X	X	X	X
BT 3896 Hispinae sp 012	X	X	X	X	X	X	X	X	X
BT 3915 Galerucinae sp 007	X	X	X	X	X	X	X	X	X
BT 3916 Galerucinae sp 007	X	X	X	X	X	X	X	X	X
BT 3917 Alticinae sp 065	X	X	X	X	X	X	X	X	X
BT 3918 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 3919 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 3920 Alticinae sp 165	X	X	X	X	X	X	X	X	X
BT 3921 Eumolpinae sp 042	X	X	X	X	X	X	X	X	X
BT 3922 Alticinae sp 264	X	X	X	X	X	X	X	X	X
BT 3923 Cassidinae sp 006	X	X	X	X	X	X	X	X	X
BT 3947 Eumolpinae sp 019	X	X	X	X	X	X	X	X	X
BT 3948 Alticinae sp 044	X	X	X	X	X	X	X	X	X
BT 3949 Alticinae sp 077	X	X	X	X	X	X	X	X	X
BT 3950 Galerucinae sp 061	X	X	X	X	X	X	X	X	X
BT 3951 Alticinae sp 238	X	X	X	X	X	X	X	X	X
BT 3952 Alticinae sp 092	X	X	X	X	X	X	X	X	X
BT 3953 Alticinae sp 112	X	X	X	X	X	X	X	X	X
BT 3954 Alticinae sp 112	X	X	X	X	X	X	X	X	X
BT 3955 Alticinae sp 112	X	X	X	X	X	X	X	X	X
BT 3957 Galerucinae sp 066	X	X	X	X	X	X	X	X	X
BT 3958 Eumolpinae sp 019	X	X	X	X	X	X	X	X	X
BT 3959 Galerucinae sp 069	X	X	X	X	X	X	X	X	X
BT 3960 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 3961 Cassidinae sp 004	X	X	X	X	X	X	X	X	X
BT 3962 Alticinae sp 117	X	X	X	X	X	X	X	X	X
BT 3963 Alticinae sp 085	X	X	X	X	X	X	X	X	X
BT 3964 Alticinae sp 045	X	X	X	X	X	X	X	X	X
BT 3965 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 3966 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 3967 Galerucinae sp 034	X	X	X	X	X	X	X	X	X
BT 3968 Eumolpinae sp 061	X	X	X	X	X	X	X	X	X
BT 3969 Alticinae sp 096	X	X	X	X	X	X	X	X	X
BT 3970 Alticinae sp 085	X	X	X	X	X	X	X	X	X
BT 3971 Galerucinae sp 026	X	X	X	X	X	X	X	X	X
BT 3972 Alticinae sp 231	X	X	X	X	X	X	X	X	X
BT 3973 Alticinae sp 248	X	X	X	X	X	X	X	X	X
BT 3974 Alticinae sp 094	X	X	X	X	X	X	X	X	X
BT 3975 Alticinae sp 172	X	X	X	X	X	X	X	X	X
BT 3976 Eumolpinae sp 050	X	X	X	X	X	X	X	X	X
BT 3977 Hispinae sp 016	X	X	X	X	X	X	X	X	X
BT 3986 Galerucinae sp 007	X	X	X	X	X	X	X	X	X
BT 3988 Eumolpinae sp 059	X	X	X	X	X	X	X	X	X
BT 3989 Alticinae sp 238	X	X	X	X	X	X	X	X	X
BT 3990 Alticinae sp 173	X	X	X	X	X	X	X	X	X
BT 3994 Hispinae sp 015	X	X	X	X	X	X	X	X	X
BT 3995 Galerucinae sp 074	X	X	X	X	X	X	X	X	X
BT 3996 Alticinae sp 042	X	X	X	X	X	X	X	X	X
BT 3997 Galerucinae sp 033	X	X	X	X	X	X	X	X	X
BT 3998 Galerucinae sp 007	X	X	X	X	X	X	X	X	X
BT 3999 Alticinae sp 009	X	X	X	X	X	X	X	X	X
BT 4000 Alticinae sp 175	X	X	X	X	X	X	X	X	X
BT 4001 Alticinae sp 175	X	X	X	X	X	X	X	X	X
BT 4002 Galerucinae sp 066	X	X	X	X	X	X	X	X	X
BT 4003 Galerucinae sp 083	X	X	X	X	X	X	X	X	X
BT 4004 Galerucinae sp 083	X	X	X	X	X	X	X	X	X
BT 4005 Galerucinae sp 031	X	X	X	X	X	X	X	X	X
BT 4006 Alticinae sp 251	X	X	X	X	X	X	X	X	X
BT 4007 Alticinae sp 145	X	X	X	X	X	X	X	X	X
BT 4008 Galerucinae sp 061	X	X	X	X	X	X	X	X	X
BT 4009 Alticinae sp 018	X	X	X	X	X	X	X	X	X
BT 4010 Eumolpinae sp 040	X	X	X	X	X	X	X	X	X
BT 4011 Eumolpinae sp 040	X	X	X	X	X	X	X	X	X
BT 4012 Eumolpinae sp 040	X	X	X	X	X	X	X	X	X
BT 4013 Galerucinae sp 046	X	X	X	X	X	X	X	X	X
BT 4014 Cassidinae sp 005	X	X	X	X	X	X	X	X	X
BT 4015 Galerucinae sp 064	X	X	X	X	X	X	X	X	X
BT 4016 Galerucinae sp 110	X	X	X	X	X	X	X	X	X
BT 4017 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 4018 Alticinae sp 104	X	X	X	X	X	X	X	X	X
BT 4019 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 4020 Eumolpinae sp 039	X	X	X	X	X	X	X	X	X
BT 4021 Alticinae sp 008	X	X	X	X	X	X	X	X	X
BT 4022 Cassidinae sp 004	X	X	X	X	X	X	X	X	X
BT 4026 Galerucinae sp 017	X	X	X	X	X	X	X	X	X
BT 4031 Alticinae sp 145	X	X	X	X	X	X	X	X	X
BT 4032 Hispinae sp 007	X	X	X	X	X	X	X	X	X
BT 4033 Eumolpinae sp 037	X	X	X	X	X	X	X	X	X
BT 4035 Hispinae sp 003	X	X	X	X	X	X	X	X	X
BT 4036 Galerucinae sp 078	X	X	X	X	X	X	X	X	X
BT 4047 Alticinae sp 181	X	X	X	X	X	X	X	X	X

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Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT_4048_Criocerinae_sp_001	X	X	X	X	X	X	X	X	X
BT_4049_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4050_Eumolpinae_sp_042	X	X	X	X	X	X	X	X	X
BT_4051_Eumolpinae_sp_042	X	X	X	X	X	X	X	X	X
BT_4052_Alticinae_sp_2T3	X	X	X	X	X	X	X	X	X
BT_4053_Eumolpinae_sp_065	X	X	X	X	X	X	X	X	X
BT_4054_Alticinae_sp_018	X	X	X	X	X	X	X	X	X
BT_4055_Alticinae_sp_018	X	X	X	X	X	X	X	X	X
BT_4056_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4057_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4058_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4059_Galerucinae_sp_107	X	X	X	X	X	X	X	X	X
BT_4060_Eumolpinae_sp_038	X	X	X	X	X	X	X	X	X
BT_4061_Alticinae_sp_106	X	X	X	X	X	X	X	X	X
BT_4062_Eumolpinae_sp_038	X	X	X	X	X	X	X	X	X
BT_4063_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4064_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4065_Alticinae_sp_018	X	X	X	X	X	X	X	X	X
BT_4066_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4067_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4182_Galerucinae_sp_092	X	X	X	X	X	X	X	X	X
BT_4186_Galerucinae_sp_020	X	X	X	X	X	X	X	X	X
BT_4194_Galerucinae_sp_095	X	X	X	X	X	X	X	X	X
BT_4195_Galerucinae_sp_029	X	X	X	X	X	X	X	X	X
BT_4198_Criocerinae_sp_002	X	X	X	X	X	X	X	X	X
BT_4207_Cassidinae_sp_021	X	X	X	X	X	X	X	X	X
BT_4208_Galerucinae_sp_028	X	X	X	X	X	X	X	X	X
BT_4209_Criocerinae_sp_007	X	X	X	X	X	X	X	X	X
BT_4215_Alticinae_sp_239	X	X	X	X	X	X	X	X	X
BT_4216_Alticinae_sp_187	X	X	X	X	X	X	X	X	X
BT_4217_Alticinae_sp_262	X	X	X	X	X	X	X	X	X
BT_4241_Cassidinae_sp_014	X	X	X	X	X	X	X	X	X
BT_4252_Galerucinae_sp_007	X	X	X	X	X	X	X	X	X
BT_4253_Galerucinae_sp_002	X	X	X	X	X	X	X	X	X
BT_4294_Hispinae_sp_020	X	X	X	X	X	X	X	X	X
BT_4295_Galerucinae_sp_029	X	X	X	X	X	X	X	X	X
BT_4306_Cassidinae_sp_019	X	X	X	X	X	X	X	X	X
BT_4321_Galerucinae_sp_017	X	X	X	X	X	X	X	X	X
BT_4344_Galerucinae_sp_011	X	X	X	X	X	X	X	X	X
BT_4350_Alticinae_sp_2T8	X	X	X	X	X	X	X	X	X
BT_4351_Galerucinae_sp_017	X	X	X	X	X	X	X	X	X
BT_4354_Alticinae_sp_163	X	X	X	X	X	X	X	X	X
BT_4402_Alticinae_sp_222	X	X	X	X	X	X	X	X	X
BT_4421_Alticinae_sp_223	X	X	X	X	X	X	X	X	X
BT_4434_Alticinae_sp_163	X	X	X	X	X	X	X	X	X
BT_4468_Galerucinae_sp_007	X	X	X	X	X	X	X	X	X
BT_4477_Alticinae_sp_155	X	X	X	X	X	X	X	X	X
BT_4500_Galerucinae_sp_097	X	X	X	X	X	X	X	X	X
BT_4509_Cassidinae_sp_018	X	X	X	X	X	X	X	X	X
BT_4510_Hispinae_sp_026	X	X	X	X	X	X	X	X	X
BT_4511_Alticinae_sp_229	X	X	X	X	X	X	X	X	X
BT_4550_Alticinae_sp_220	X	X	X	X	X	X	X	X	X
BT_4572_Alticinae_sp_219	X	X	X	X	X	X	X	X	X
BT_4581_Galerucinae_sp_005	X	X	X	X	X	X	X	X	X
BT_4605_Galerucinae_sp_110	X	X	X	X	X	X	X	X	X
BT_4684_Cassidinae_sp_001	X	X	X	X	X	X	X	X	X
BT_4687_Galerucinae_sp_036	X	X	X	X	X	X	X	X	X
BT_4732_Alticinae_sp_095	X	X	X	X	X	X	X	X	X
BT_4733_Eumolpinae_sp_068	X	X	X	X	X	X	X	X	X
BT_4734_Alticinae_sp_172	X	X	X	X	X	X	X	X	X
BT_4735_Alticinae_sp_235	X	X	X	X	X	X	X	X	X
BT_4736_Alticinae_sp_172	X	X	X	X	X	X	X	X	X
BT_4737_Alticinae_sp_172	X	X	X	X	X	X	X	X	X
BT_4738_Alticinae_sp_172	X	X	X	X	X	X	X	X	X
BT_4739_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_4740_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_4741_Alticinae_sp_073	X	X	X	X	X	X	X	X	X
BT_4742_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_4743_Alticinae_sp_181	X	X	X	X	X	X	X	X	X
BT_4744_Eumolpinae_sp_001	X	X	X	X	X	X	X	X	X
BT_4745_Galerucinae_sp_056	X	X	X	X	X	X	X	X	X
BT_4745_Galerucinae_sp_080	X	X	X	X	X	X	X	X	X
BT_4746_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_4747_Galerucinae_sp_074	X	X	X	X	X	X	X	X	X
BT_4748_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_4749_Galerucinae_sp_074	X	X	X	X	X	X	X	X	X
BT_4750_Eumolpinae_sp_024	X	X	X	X	X	X	X	X	X
BT_4751_Alticinae_sp_253	X	X	X	X	X	X	X	X	X
BT_4753_Eumolpinae_sp_038	X	X	X	X	X	X	X	X	X
BT_4754_Alticinae_sp_195	X	X	X	X	X	X	X	X	X
BT_4772_Alticinae_sp_009	X	X	X	X	X	X	X	X	X
BT_4773_Alticinae_sp_127	X	X	X	X	X	X	X	X	X
BT_4774_Alticinae_sp_050	X	X	X	X	X	X	X	X	X
BT_4775_Alticinae_sp_211	X	X	X	X	X	X	X	X	X
BT_4776_Galerucinae_sp_029	X	X	X	X	X	X	X	X	X
BT_4777_Alticinae_sp_183	X	X	X	X	X	X	X	X	X
BT_4778_Alticinae_sp_181	X	X	X	X	X	X	X	X	X
BT_4779_Galerucinae_sp_106	X	X	X	X	X	X	X	X	X
BT_4780_Alticinae_sp_252	X	X	X	X	X	X	X	X	X
BT_4781_Alticinae_sp_141	X	X	X	X	X	X	X	X	X
BT_4782_Galerucinae_sp_029	X	X	X	X	X	X	X	X	X
BT_4783_Hispinae_sp_016	X	X	X	X	X	X	X	X	X
BT_4784_Alticinae_sp_245	X	X	X	X	X	X	X	X	X
BT_4785_Hispinae_sp_027	X	X	X	X	X	X	X	X	X
BT_4786_Alticinae_sp_172	X	X	X	X	X	X	X	X	X
BT_4787_Alticinae_sp_172	X	X	X	X	X	X	X	X	X
BT_4788_Eumolpinae_sp_058	X	X	X	X	X	X	X	X	X
BT_4789_Alticinae_sp_172	X	X	X	X	X	X	X	X	X
BT_4790_Alticinae_sp_189	X	X	X	X	X	X	X	X	X
BT_4791_Alticinae_sp_172	X	X	X	X	X	X	X	X	X
BT_4792_Alticinae_sp_187	X	X	X	X	X	X	X	X	X
BT_4793_Alticinae_sp_263	X	X	X	X	X	X	X	X	X
BT_4794_Alticinae_sp_198	X	X	X	X	X	X	X	X	X
BT_4795_Alticinae_sp_249	X	X	X	X	X	X	X	X	X
BT_4796_Alticinae_sp_261	X	X	X	X	X	X	X	X	X
BT_4797_Galerucinae_sp_034	X	X	X	X	X	X	X	X	X
BT_4798_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4799_Eumolpinae_sp_038	X	X	X	X	X	X	X	X	X
BT_4800_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4801_Alticinae_sp_104	X	X	X	X	X	X	X	X	X
BT_4802_Eumolpinae_sp_038	X	X	X	X	X	X	X	X	X
BT_4803_Alticinae_sp_101	X	X	X	X	X	X	X	X	X
BT_4804_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4805_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4806_Alticinae_sp_115	X	X	X	X	X	X	X	X	X
BT_4807_Eumolpinae_sp_042	X	X	X	X	X	X	X	X	X
BT_4808_Alticinae_sp_018	X	X	X	X	X	X	X	X	X
BT_4809_Galerucinae_sp_034	X	X	X	X	X	X	X	X	X

Continued on next page(s)



Table S3 – continued from previous page(s)

Specimen	1	1b	2	2a	2b	3	3a	3b	4
BT 4810 Alticinae sp 115	X	X	X	X	X	X	X	X	X
BT 4811 Alticinae sp 115	X	X	X	X	X	X	X	X	X
BT 4812 Alticinae sp 101	X	X	X	X	X	X	X	X	X
BT 4813 Alticinae sp 198	X	X	X	X	X	X	X	X	X
BT 4814 Alticinae sp 199	X	X	X	X	X	X	X	X	X
BT 4815 Alticinae sp 086	X	X	X	X	X	X	X	X	X
BT 4816 Alticinae sp 086									
BT 4817 Alticinae sp 086									
BT 4851 Eumolpinae sp 024	X	X	X	X	X	X	X	X	X
BT 4856 Alticinae sp 186	X	X	X	X	X	X	X	X	X
BT 4857 Alticinae sp 172	X	X	X	X	X	X	X	X	X
BT 4858 Alticinae sp 187	X	X	X	X	X	X	X	X	X
BT 4859 Alticinae sp 057	X	X	X	X	X	X	X	X	X
BT 4860 Eumolpinae sp 046	X	X	X	X	X	X	X	X	X
BT 4861 Alticinae sp 149	X	X	X	X	X	X	X	X	X
BT 4862 Alticinae sp 149	X	X	X	X	X	X	X	X	X
BT 4863 Alticinae sp 168	X	X	X	X	X	X	X	X	X
BT 4864 Alticinae sp 149	X	X	X	X	X	X	X	X	X
BT 4867 Galerucinae sp 003	X	X							
BT 4875 Criocerinae sp 009	X	X							
BT 4885 Galerucinae sp 056	X	X	X	X	X	X	X	X	X
BT 4886 Alticinae sp 121	X	X	X	X	X	X	X	X	X
BT 4887 Alticinae sp 102	X	X	X	X	X	X	X	X	X
BT 4888 Eumolpinae sp 059	X	X	X	X	X	X	X	X	X
BT 4889 Alticinae sp 051	X	X	X	X	X	X	X	X	X
BT 4890 Alticinae sp 051									
BT 4891 Galerucinae sp 087	X	X	X	X	X	X	X	X	X
BT 4892 Alticinae sp 149	X	X	X	X	X	X	X	X	X
BT 4893 Alticinae sp 149									
BT 4894 Alticinae sp 149									
BT 4895 Alticinae sp 149									
BT 4896 Alticinae sp 149									
BT 4897 Alticinae sp 149									
BT 4898 Alticinae sp 168	X	X	X	X	X	X	X	X	X
BT 4899 Alticinae sp 168									
BT 4900 Alticinae sp 151	X	X	X	X	X	X	X	X	X
BT 4901 Galerucinae sp 057	X	X	X	X	X	X	X	X	X
BT 4902 Galerucinae sp 057									
BT 4903 Alticinae sp 267	X	X	X	X	X	X	X	X	X
BT 4904 Alticinae sp 267									
BT 4905 Alticinae sp 201	X	X	X	X	X	X	X	X	X
BT 4906 Alticinae sp 211	X	X	X	X	X	X	X	X	X
BT 4907 Alticinae sp 211									
BT 4908 Hispinae sp 013	X	X	X	X	X	X	X	X	X
BT 4909 Alticinae sp 025	X	X	X	X	X	X	X	X	X
BT 4910 Alticinae sp 149	X	X	X	X	X	X	X	X	X
BT 4911 Alticinae sp 057	X	X	X	X	X	X	X	X	X
BT 4912 Alticinae sp 086	X	X	X	X	X	X	X	X	X
BT 4914 Galerucinae sp 003	X	X	X	X	X	X	X	X	X
BT 4915 Alticinae sp 050	X	X	X	X	X	X	X	X	X
BT 4916 Eumolpinae sp 063	X	X	X	X	X	X	X	X	X
BT 4917 Alticinae sp 166	X	X	X	X	X	X	X	X	X
BT 4918 Alticinae sp 050	X	X	X	X	X	X	X	X	X
BT 4919 Eumolpinae sp 056	X	X	X	X	X	X	X	X	X
BT 4920 Alticinae sp 057									
BT 4921 Alticinae sp 057									
BT 4922 Alticinae sp 009	X	X	X	X	X	X	X	X	X
BT 4923 Galerucinae sp 048	X	X	X	X	X	X	X	X	X
BT 4924 Alticinae sp 051	X	X	X	X	X	X	X	X	X
BT 4925 Alticinae sp 102	X	X	X	X	X	X	X	X	X
BT 4926 Alticinae sp 025	X	X	X	X	X	X	X	X	X
BT 4927 Alticinae sp 268	X	X	X	X	X	X	X	X	X
BT 4928 Alticinae sp 172	X	X	X	X	X	X	X	X	X
BT 4929 Alticinae sp 172	X	X	X	X	X	X	X	X	X
BT 4930 Alticinae sp 172	X	X	X	X	X	X	X	X	X
BT 4931 Alticinae sp 244	X	X	X	X	X	X	X	X	X
BT 4937 Alticinae sp 127	X	X	X	X	X	X	X	X	X
BT 4938 Alticinae sp 149	X	X	X	X	X	X	X	X	X
BT 4939 Alticinae sp 149									
BT 4940 Alticinae sp 059									
BT 4941 Alticinae sp 127	X	X	X	X	X	X	X	X	X
BT 4942 Alticinae sp 102	X	X	X	X	X	X	X	X	X
BT 4944 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 4945 Eumolpinae sp 002	X	X	X	X	X	X	X	X	X
BT 4946 Eumolpinae sp 034	X	X	X	X	X	X	X	X	X
BT 4947 Alticinae sp 199	X	X	X	X	X	X	X	X	X
BT 4948 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 4949 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 4950 Galerucinae sp 088	X	X	X	X	X	X	X	X	X
BT 4951 Alticinae sp 061	X	X	X	X	X	X	X	X	X
BT 4952 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 4953 Alticinae sp 199	X	X	X	X	X	X	X	X	X
BT 4954 Alticinae sp 199									
BT 4955 Alticinae sp 199									
BT 4956 Alticinae sp 086	X	X	X	X	X	X	X	X	X
BT 4957 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 4958 Alticinae sp 133	X	X	X	X	X	X	X	X	X
BT 4960 Alticinae sp 086	X	X	X	X	X	X	X	X	X
BT 4961 Alticinae sp 086									
BT 4962 Alticinae sp 086									
BT 4963 Alticinae sp 215	X	X	X	X	X	X	X	X	X
BT 4964 Alticinae sp 150	X	X	X	X	X	X	X	X	X
BT 4965 Eumolpinae sp 034	X	X	X	X	X	X	X	X	X
BT 4966 Alticinae sp 008	X	X	X	X	X	X	X	X	X
BT 4967 Eumolpinae sp 038	X	X	X	X	X	X	X	X	X
BT 4968 Eumolpinae sp 038									
BT 4969 Alticinae sp 045	X	X	X	X	X	X	X	X	X
BT 4970 Galerucinae sp 081	X	X	X	X	X	X	X	X	X
BT 4971 Alticinae sp 192	X	X	X	X	X	X	X	X	X
BT 4972 Alticinae sp 050	X	X	X	X	X	X	X	X	X
BT 4986 Galerucinae sp 101	X	X	X	X	X	X	X	X	X
BT 4994 Alticinae sp 172	X	X	X	X	X	X	X	X	X
BT 4995 Galerucinae sp 103	X	X	X	X	X	X	X	X	X
BT 5029 Galerucinae sp 099	X	X							
BT 5122 Alticinae sp 221	X	X							



# Additional information: Species delimitation results

Table E.1: Results of species delimitation for each specimen. Only specimens from data set 4. D.-Cluster = Distance-Cluster

Specimen	Haplotype-Network	3% D.-Cluster	5% D.-Cluster	7.5% D.-Cluster	GMYC-Cluster	PTP-Cluster	Haplotype	
BT-0001	Eumolpinae sp. 1	Network282	3C1001	5C1001	75C1001	GMYC008	PTP023	H416
BT-0002	Alticinae sp. 42	Network52	3C1002	5C1002	75C1002	GMYC219	PTP114	H085
BT-0004	Eumolpinae sp. 42	Network257	3C1003	5C1003	75C1003	GMYC001	PTP003	H388
BT-0005	Galerucinae sp. 40	Network216	3C1004	5C1004	75C1004	GMYC100	PTP238	H329
BT-0007	Galerucinae sp. 38	Network98	3C1005	5C1005	75C1005	GMYC117	PTP272	H154
BT-0008	Alticinae sp. 243	Network54	3C1006	5C1006	75C1006	GMYC220	PTP115	H088
BT-0012	Eumolpinae sp. 21	Network270	3C1007	5C1007	75C1007	GMYC022	PTP007	H401
BT-0015	Galerucinae sp. 76	Network118	3C1008	5C1008	75C1008	GMYC113	PTP288	H185
BT-0017	Alticinae sp. 43	Network49	3C1009	5C1009	75C1009	GMYC227	PTP122	H081
BT-0021	Alticinae sp. 7	Network187	3C1010	5C1010	75C1010	GMYC087	PTP204	H285
BT-0022	Alticinae sp. 219	Network190	3C1115	5C1011	75C1011	GMYC089	PTP202	H288
BT-0024	Galerucinae sp. 1	Network112	3C1011	5C1012	75C1012	GMYC138	PTP264	H176
BT-0033	Galerucinae sp. 37	Network116	3C1012	5C1013	75C1013	GMYC115	PTP286	H181
BT-0034	Eumolpinae sp. 14	Network285	3C1013	5C1014	75C1014	GMYC009	PTP016	H419
BT-0035	Eumolpinae sp. 006	Network275	3C1014	5C1015	75C1015	GMYC017	PTP025	H407
BT-0036	Galerucinae sp. 11	Network103	3C1015	5C1016	75C1016	GMYC143	PTP258	H160
BT-0043	Galerucinae sp. 5	Network110	3C1016	5C1017	75C1017	GMYC137	PTP265	H172
BT-0046	Alticinae sp. 243	Network53	3C1006	5C1006	75C1006	GMYC221	PTP116	H087
BT-0047	Alticinae sp. 42	Network52	3C1002	5C1002	75C1002	GMYC219	PTP114	H086
BT-0048	Galerucinae sp. 39	Network97	3C1017	5C1018	75C1018	GMYC118	PTP271	H153
BT-0049	Galerucinae sp. 41	Network217	3C1018	5C1019	75C1019	GMYC101	PTP237	H330
BT-0088	Galerucinae sp. 7	Network109	3C1019	5C1020	75C1020	GMYC134	PTP267	H170
BT-0089	Eumolpinae sp. 1	Network282	3C1001	5C1001	75C1001	GMYC008	PTP023	H416
BT-0090	Galerucinae sp. 76	Network118	3C1008	5C1008	75C1008	GMYC113	PTP288	H184
BT-0091	Eumolpinae sp. 1	Network282	3C1001	5C1001	75C1001	GMYC008	PTP023	H416
BT-0094	Galerucinae sp. 11	Network103	3C1015	5C1016	75C1016	GMYC143	PTP258	H161
BT-0095	Cassidinae sp. 1	Network261	3C1020	5C1021	75C1021	GMYC280	PTP038	H392
BT-0096	Alticinae sp. 10	Network195	3C1021	5C1022	75C1022	GMYC097	PTP195	H303
BT-0098	Galerucinae sp. 002	Network111	3C1022	5C1023	75C1023	GMYC139	PTP263	H173
BT-0099	Galerucinae sp. 7	Network92	3C1023	5C1024	75C1024	GMYC126	PTP284	H147
BT-0102	Alticinae sp. 44	Network42	3C1024	5C1025	75C1025	GMYC214	PTP102	H060
BT-0103	Eumolpinae sp. 38	Network284	3C1025	5C1026	75C1026	GMYC010	PTP017	H418
BT-0107	Galerucinae sp. 46	Network155	3C1026	5C1027	75C1027	GMYC178	PTP249	H241
BT-0109	Alticinae sp. 251	Network81	3C1027	5C1028	75C1028	GMYC057	PTP156	H124
BT-0110	Alticinae sp. 87	Network88	3C1028	5C1029	75C1029	GMYC061	PTP164	H136
BT-0114	Galerucinae sp. 62	Network205	3C1029	5C1030	75C1030	GMYC111	PTP239	H317
BT-0115	Alticinae sp. 98	Network10	3C1030	5C1031	75C1031	GMYC253	PTP060	H010
BT-0118	Eumolpinae sp. 19	Network265	3C1031	5C1032	75C1032	GMYC028	PTP004	H396
BT-0119	Alticinae sp. 124	Network87	3C1032	5C1033	75C1033	GMYC048	PTP166	H135
BT-0121	Alticinae sp. 107	Network218	3C1033	5C1034	75C1034	GMYC062	PTP141	H331
BT-0123	Alticinae sp. 129	Network212	3C1034	5C1035	75C1035	GMYC065	PTP098	H325
BT-0125	Alticinae sp. 97	Network13	3C1035	5C1036	75C1036	GMYC258	PTP065	H020
BT-0126	Alticinae sp. 123	Network148	3C1036	5C1037	75C1037	GMYC070	PTP144	H228
BT-0130	Galerucinae sp. 34	Network183	3C1037	5C1038	75C1038	GMYC103	PTP232	H280
BT-0134	Galerucinae sp. 7	Network92	3C1023	5C1024	75C1024	GMYC126	PTP284	H148
BT-0135	Eumolpinae sp. 19	Network265	3C1031	5C1032	75C1032	GMYC028	PTP004	H396
BT-0137	Cassidinae sp. 4	Network253	3C1038	5C1039	75C1039	GMYC284	PTP053	H379
BT-0139	Alticinae sp. 10	Network195	3C1021	5C1022	75C1022	GMYC097	PTP195	H304
BT-0140	Alticinae sp. 28	Network29	3C1039	5C1040	75C1040	GMYC209	PTP129	H041
BT-0144	Eumolpinae sp. 38	Network283	3C1040	5C1041	75C1041	GMYC011	PTP018	H417
BT-0145	Galerucinae sp. 61	Network93	3C1041	5C1042	75C1042	GMYC131	PTP279	H149
BT-0146	Alticinae sp. 29	Network68	3C1042	5C1043	75C1043	GMYC049	PTP168	H107
BT-0147	Alticinae sp. 62	Network57	3C1043	5C1044	75C1044	GMYC224	PTP117	H092
BT-0148	Alticinae sp. 66	Network85	3C1044	5C1045	75C1045	GMYC051	PTP169	H131
BT-0149	Alticinae sp. 249	Network79	3C1045	5C1046	75C1046	GMYC037	PTP147	H121
BT-0153	Alticinae sp. 109	Network171	3C1046	5C1047	75C1047	GMYC254	PTP062	H264
BT-0154	Alticinae sp. 115	Network10	3C1030	5C1031	75C1031	GMYC253	PTP060	H010
BT-0155	Alticinae sp. 193	Network172	3C1047	5C1048	75C1048	GMYC243	PTP206	H266
BT-0157	Alticinae sp. 97	Network11	3C1048	5C1049	75C1049	GMYC256	PTP064	H015
BT-0158	Eumolpinae sp. 2	Network285	3C1013	5C1014	75C1014	GMYC009	PTP016	H419
BT-0159	Galerucinae sp. 096	Network180	3C1049	5C1050	75C1050	GMYC174	PTP252	H277
BT-0174	Galerucinae sp. 46	Network155	3C1026	5C1027	75C1027	GMYC178	PTP249	H241
BT-0176	Galerucinae sp. 46	Network156	3C1050	5C1051	75C1051	GMYC179	PTP250	H242
BT-0183	Galerucinae sp. 34	Network183	3C1037	5C1038	75C1038	GMYC103	PTP232	H280
BT-0188	Galerucinae sp. 11	Network103	3C1015	5C1016	75C1016	GMYC143	PTP258	H162
BT-0189	Alticinae sp.161	Network38	3C1051	5C1052	75C1052	GMYC151	PTP071	H053
BT-0190	Eumolpinae sp. 1	Network282	3C1001	5C1001	75C1001	GMYC008	PTP023	H416
BT-0196	Galerucinae sp. 10	Network232	3C1052	5C1053	75C1053	GMYC076	PTP228	H353
BT-0199	Alticinae sp. 118	Network170	3C1053	5C1054	75C1054	GMYC187	PTP108	H262
BT-0201	Chrysomelinae sp. 2	Network206	3C1054	5C1055	75C1055	GMYC102	PTP236	H318
BT-0202	Galerucinae sp. 32	Network2	3C1055	5C1056	75C1056	GMYC175	PTP247	H002
BT-0204	Hispinae sp. 2	Network255	3C1056	5C1057	75C1057	GMYC273	PTP051	H382
BT-0207	Galerucinae sp. 69	Network219	3C1057	5C1058	75C1058	GMYC086	PTP215	H333
BT-0208	Eumolpinae sp. 19	Network265	3C1031	5C1032	75C1032	GMYC028	PTP004	H396
BT-0209	Cassidinae sp. 5	Network242	3C1058	5C1059	75C1059	GMYC277	PTP043	H364
BT-0211	Alticinae sp. 87	Network88	3C1028	5C1029	75C1029	GMYC061	PTP164	H136
BT-0212	Galerucinae sp. 66	Network223	3C1059	5C1060	75C1060	GMYC079	PTP222	H341
BT-0213	Galerucinae sp. 24	Network197	3C1060	5C1061	75C1061	GMYC108	PTP230	H306
BT-0214	Alticinae sp. 28	Network29	3C1039	5C1040	75C1040	GMYC209	PTP129	H043
BT-0218	Galerucinae sp. 31	Network203	3C1061	5C1062	75C1062	GMYC084	PTP217	H313
BT-0219	Eumolpinae sp. 073	Network244	3C1062	5C1063	75C1063	GMYC031	PTP002	H368
BT-0220	Alticinae sp. 115	Network10	3C1030	5C1031	75C1031	GMYC253	PTP060	H010
BT-0221	Eumolpinae sp. 10	Network274	3C1063	5C1064	75C1064	GMYC021	PTP008	H406

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APPENDIX G

# Erklärung

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Ich versichere, dass ich diese Arbeit selbständig verfasst, keine anderen Quellen und Hilfsmittel als die angegebenen benutzt und die Stellen der Arbeit, die anderen Werken dem Wortlaut oder Sinn nach entnommen sind, kenntlich gemacht habe.

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