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A cladistic analysis of whiteflies, subfamily Aleyrodinae (Hemiptera: Sternorrhyncha: Aleyrodidae)

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Abstract

Relationships within the Aleyrodinae are investigated phylogenetically through a cladistic analysis of 94 (70 binary and 24 multistate) morphological characters derived from the pupal case. This is the first attempt to construct a phylogenetic classification at genus level on a world basis for the subfamily Aleyrodinae. More than 430 species, about one third of all described species, were examined. These belonged to 124 genera, of which 117 were aleyrodines (including two undescribed genera) and seven were out-groups from the subfamily Aleurodicinae (Hemiptera: Aleyrodidae). Maximum parsimony analysis yielded more than 30 000 most-parsimonious trees (length=2730, RI=0.672, CI=0.137). Based upon the strict consensus tree, only 33 of the currently accepted genera, for which we had multiple representatives, were found to be monophyletic (44 genera of the 117 examined genera were monobasic and/or only one species was available for study). Monophyly of some economically important genera, e.g. *Aleurolobus*, *Bemisia* and *Dialeurodes*, as well as that of most previously proposed tribes, was not supported. No parallel diversification of whiteflies with host plants was detected, mostly because of within-species host switching. Problems in determining generic level characters and applicability of tribal classification for this subfamily are discussed. One new genus, *Pseudozaphanera* Manzari **gen. n.**, and seven new combinations, *Aleuroclava lanceolata* (Takahashi) **comb. n.**, *Crescentaleyrodes fumipennis* (Hempel) **comb. n.**, *Pseudozaphanera niger* (Maskell) **comb. n.**, *Pseudozaphanera papyrocarpae* (Martin) **comb. n.**, *Pseudozaphanera rhachisreticulata* (Martin) **comb. n.**, *Pseudozaphanera splendida* (Martin) **comb. n.**, *Pseudozaphanera wariensis* (Martin) **comb. n.**, are proposed as well as three new generic synonyms; *Taiwanaleyrodes* Takahashi is placed in synonymy with *Aleuroclava* Singh **syn. n.**, *Hempelia* Sampson & Drews with *Aleurothrixus* Quaintance & Baker **syn. n.** and *Rositalleyrodes* Meganathan & David with *Aleurolobus* Quaintance & Baker **syn. n.**

Keywords: *Host–plant relationships, phylogeny, successive approximations character weighting, tribal classification*

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Introduction

The Aleyrodidae is composed of tiny insects (1–3 mm in body length), usually called whiteflies, though one of the most widespread pest species, *Aleurocanthus woglumi* Ashby, is called the citrus blackfly (Mound 1973). Mound and Halsey (1978) catalogued the whiteflies of the world and listed 1156 species, which has now increased to approximately 1450 (Martin et al. 2000). However, the real number is certainly much higher.

Many whiteflies are agricultural pests, especially in the tropics and subtropics (Calvert et al. 2001). Feeding activity damages their host plants and some species are known to transmit diseases, especially those caused by viruses (Gullan and Martin 2003). In spite of this, the family is poorly studied taxonomically, particularly in a phylogenetic context. Almost all publications dealing with whitefly taxonomy are restricted to alpha taxonomy. Major publications provide lists of species and descriptions of new species and/or the genera, mostly with restricted geographical compass (e.g. Singh 1931; Takahashi 1932; Corbett 1935; Russell 1948; Mound 1965; Cohic 1966; David and Subramaniam 1976; Bink-Moenen 1983; Jesudasan and David 1991; Martin 1999).

The taxonomy of the Aleyrodidae has long been problematic. Unusually amongst insects, whitefly generic classification is largely based on one of the immature stages, the so-called, pupal case (=puparium=fourth instar=last nymphal stage=last larval stage), rather than on the adults (Gill 1990). Quite often, third instar larvae have been used in species descriptions, having been mistaken for the so-called puparial stage (Martin 1999). Unfortunately, whitefly puparia are notorious for displaying variation induced by environmental and physical factors, such as temperature and humidity (Mohanty and Basu 1986), as well as by the type of leaf surface upon which they have developed (Russell 1948; Mound 1963; David and Ananthakrishnan 1976; Neal and Bentz 1999; Guershon and Gerling 2001). Their generic classification was largely laid out by Quaintance and Baker (1913, 1914) and is based mainly on pupal characters; the morphological traits of adults are currently poorly understood and so do not readily permit differentiation between genera or species (Frohlich et al. 1999).

Apart from a few genealogical diagrams proposed (e.g. Quaintance and Baker (1913) (Figure 1); Bondar 1923), the first attempt to construct a phylogenetic classification for whiteflies using morphological characters was made by Jensen (1999) for the species of *Dialeurodes* Cockerell. In Jensen's article, alternative generic names were not proposed despite the fact that his results showed the genus to be paraphyletic with respect to several others. Subsequently, Jensen (2001) proposed the use of three pre-existing generic names, *Dialeurodes*, *Massilieurodes* Goux and *Singhiella* Sampson, for many of the *Dialeurodes* of the world based on the placement of the type-species of these genera in the cladograms or based on the examination of original descriptions.

Phylogenetic analysis of 18S rDNA nucleotide sequences of hemipteran exemplars, shows the Sternorrhyncha appears to be monophyletic forming a sister group to all other hemipterans (the Euhemiptera, including Auchenorrhyncha and Heteroptera) (Campbell et al. 1994, 1995a, 1995b; von Dohlen and Moran 1995). However, relationships within the Sternorrhyncha (which includes psyllids, whiteflies, aphids and scales) are controversial with molecular and morphological data supporting different scenarios. Based upon the same molecular evidence, whiteflies form a sister group to aphids and scales, while psyllids form a sister group to all other Sternorrhyncha (Campbell et al. 1994, 1995a). However, some morphological evidence supports a sister-group relationship between whiteflies and psyllids (e.g. Quaintance and Baker 1913; Goodchild 1966; Hennig 1981; Carver et al. 1991; Shcherbakov 2000; Shcherbakov and Popov 2002), though Evans' (1963)

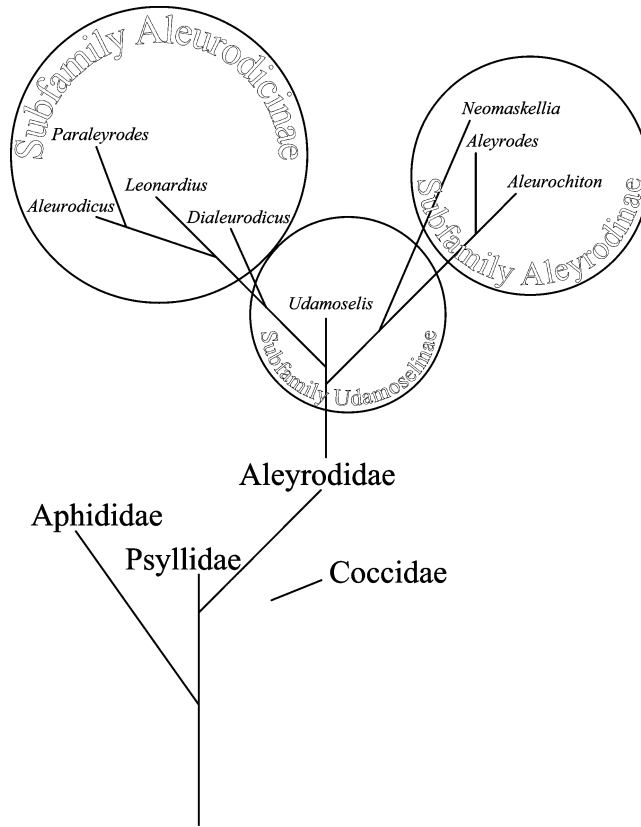


Figure 1. Genealogical diagram of the Aleyrodidae; after Quaintance and Baker (1913).

morphologically based conclusions are more in agreement with the available molecular studies. Some molecular studies of their primary bacterial endosymbionts (P-endosymbionts) generally concur with the available morphological evidence, with an analysis of 16S rDNA symbiont sequences supporting a sister relationship between primary endosymbionts of psyllids and whiteflies (Spaulding and von Dohlen 1998, 2001).

Efforts to determine the phylogenetic origin of whiteflies have been further impeded by gaps in the hemipteran fossil record, and especially a paucity of fossil whiteflies (Campbell et al. 1994). Owing to their size and fragility, adult aleyrodids would not be expected to be preserved as impression fossils often, and still less, their larvae (Evans 1963). The possibility that they arose in tropical latitudes may also account for their absence in fossil deposits now located in northern latitudes, where fossils of aphids are often found (Shcherbakov and Wegierek 1991). The oldest known fossil whiteflies, *Juleyrodes gilli* Shcherbakov and *J. visnyai* Shcherbakov, belong to an extinct subfamily, Bernaeinae, and are from the Late Jurassic or Early Cretaceous (more than 140 myr ago) (Shcherbakov 2000).

Monophyly of the Aleyrodoidea, and consequently of its single included family, Aleyrodidae, is well-supported by synapomorphies (Shcherbakov and Popov 2002). Five subfamilies have been established at various times within the extant Aleyrodidae, of which only two are now considered valid: the Aleurodicinae, found mainly in Central and South America, and the Aleyrodinae, which are more widespread (Mound and Halsey 1978; Gill 1990). The Udamoselinae was proposed on the basis of a single specimen of a South

American species, a male with a body length of 7 mm (Enderlein 1909). This subfamily is now considered dubious and the name is treated as *nomen dubium* (Mound and Halsey 1978), though Shcherbakov (2000) restored Udamoselinae in a broad sense (including Aleurodicinae). The type genera of the fourth and fifth subfamilies, Uraleyroinae and Siphonaleyroinae, i.e. *Uraleyrodes* Sampson & Drews and *Siphonaleyrodes* Takahashi, were synonymized with the aleyrodine genus *Aleurocerus* Bondar and the psyllid genus *Trioza*, respectively (Mound and Halsey 1978). These authors also synonymized Siphonaleyroinae with Triozinae. Russell (2000) provides a more detailed account of the authorships of family group names in the Aleyrodidae, and on the validities of the names of the current subfamilies within the Aleyrodidae. Recent molecular studies on their P-endosymbionts support five, or more, major groupings within the family (Baumann and Thao 2004), so perhaps a revised classification into more subfamilies may well ensue. However, the two main divisions found by Baumann and Thao (2004) correspond largely to the Aleurodicinae and Aleyrodinae.

The Aleyrodinae consists of more than 140 described genera, but their relationships are equivocal. Although 13 tribes have been proposed (Aleurocanthini Takahashi, Aleurochitonini Sampson, Aleurolobini Takahashi, Aleuroplatini David, Aleyrodini Sampson, Bemisini David, Dialeurodini Sampson, Lipaleyroini David, Neomaskellini Sampson, Siphoniniini Sampson, Tetraaleurodini David, Trialeurodini Russell and Zaphanerini David; see Sampson 1943; Russell 1947; Takahashi 1954; David 1990), at least half of the genera have not been allocated to tribes and there is little consensus about the composition of those tribes that have been proposed. For example, the well-known genus *Bemisia* Quaintance & Baker has been placed in three different tribes, viz. Dialeurodini, Aleurolobini and Bemisini by Sampson (1943), Takahashi (1954) and David (1990), respectively (Appendix A). Characters used previously to identify the tribes are shown in Appendix B.

The application of molecular techniques such as DNA nucleotide-sequencing for whitefly taxonomy has mainly been restricted to determining the biotypes of a few economically important species, especially *Bemisia tabaci* (Gennadius), and phylogenetic relationships among them rather than building a phylogenetic classification (e.g. Campbell et al. 1993; Perring et al. 1993; Wool et al. 1994; Guirao et al. 1997; Frohlich et al. 1999; Brown et al. 2000; Cervera et al. 2000; De Barro et al. 2000; Lima et al. 2000; Viscarret et al. 2003; Wu et al. 2003). Campbell et al. (1995a) presented a cladogram based on 18S rDNA sequences for 11 whitefly species belonging to seven genera, of which 10 species were from the subfamily Aleyrodinae and the last one from the Aleurodicinae. In this cladogram, the aleurodicine was sister to all aleyrodines, and among aleyrodines, members of the genus *Bemisia* formed a sister clade to all other species.

The present study is the first attempt to construct a phylogenetic classification at genus level for the Aleyrodinae on a world basis as well as testing the applicability of its tribe level classification. To avoid considering current described genera as monophyletic units, multiple representative species of each genera were selected where possible (obviously excepting monobasic genera) so as to represent a wide range of the morphological variation present, and these were treated separately in the analyses; making it possible to test the monophyly of genera and to evaluate the current whitefly classification.

Materials and methods

In total, 439 whitefly species, about one third of all described species, were examined and used to define character systems. These comprised 124 genera, of which 117 were

aleyrodines (including two undescribed genera mentioned as New genus 1 and New genus 2) and seven were aleurodicines, which were used as out-groups. All examined species were slide-mounted specimens and, in most cases, several specimens were studied for each species. Ninety-four morphological characters, 70 binary and 24 multistate, all of which were derived from the pupal case, were scored.

Choice of species examined was largely dictated by the material in the collection of The Natural History Museum, London (BMNH), which houses one the world's largest collections of whiteflies, and includes almost all described genera. Additional material was borrowed from the United States Department of Agriculture, Beltsville, Maryland, USA (Sternorrhyncha collections of the United States National Museum of Natural History, Washington DC) (USNM).

No specimens were available for the following 25 small genera (nearly all of which are monobasic): *Aleuroporosus* Corbett, *Anomaleyrodes* Takahashi & Mamet, *Axacalia* Danzig, *Dialeurotrachelus* Takahashi, *Dothioia* Dumbleton (the paratype puparium in the BMNH was entirely black (i.e. unbleached) and no characters recognisable), *Hesperaleyrodes* Sampson, *Hindaleyrodes* Meganathan & David, *Juglasaleyrodes* Cohic, *Keralaleyrodes* Meganathan & David, *Lankaleyrodes* David, *Luederwaldtiana* Hempel, *Metaleyrodes* Sampson, *Mexicaleyrodes* Sampson & Drews, *Mohanasundaramiella* David, *Nealeyrodes* Hempel, *Neoaleurodes* Bondar (all five specimens on the *Neoaleurodes* slide from USNM were constricted and not cleared), *Neoaleurolobus* Takahashi, *Plataleyrodes* Takahashi & Mamet, *Pseudaleurolobus* Hempel, *Pseudaleyrodes* Hempel, *Rosanovia* Danzig, *Shanthinia* David, *Trichoaleyrodes* Takahashi & Mamet, *Vasantharajiella* David, and *Xenobemisia* Takahashi. The monobasic genus *Laingiella* Corbett was erected by Corbett (1926) to accommodate specimens collected on bamboo. The syntype specimens of *L. bambusae* Corbett have been deposited in the USNM and BMNH (Mound and Halsey 1978). Those in the USNM were not examined but all syntypes in the BMNH were third instar larvae and agreed with its original description (one syntype lacked its ventral surface, so there was no way to determine whether it was a third instar larva or a pupal case). Most probably, therefore, the genus was described based on the characters of third instar larvae and for this reason it was omitted from the analysis. The genera and species included in the cladistic analysis are given in Appendix C.

The characters scored include nearly all those used to describe new genera/species in the literature. Different minor ornamentations of pupal cases of some species, which were almost impossible to code consistently for all taxa, were not included. We have attempted to make character state definitions as objective as possible to minimise miscoding of the characters used in this study.

Basic morphological terminology follows that of Russell (1943, 1947, 1948, 1965, 1986), Bink-Moenen (1983), Martin (1987) and Gill (1990). Figure 2 shows the stylized pupal case with major characters.

Phylogenetic analysis

The data matrix used is given in Appendix D. Multistate characters were treated as unordered and without *a priori* weighting. Inapplicable characters and missing data were coded as question marks (Lee and Bryant 1999; Strong and Lipscomb, 1999). Characters which the "part" (e.g. seta) and "character-variable" (e.g. seta shape) could be coded as separate characters and/or fused into a single character, were coded as separate characters to avoid loss of phylogenetic information (Lee and Bryant 1999). Multistate taxa were coded as

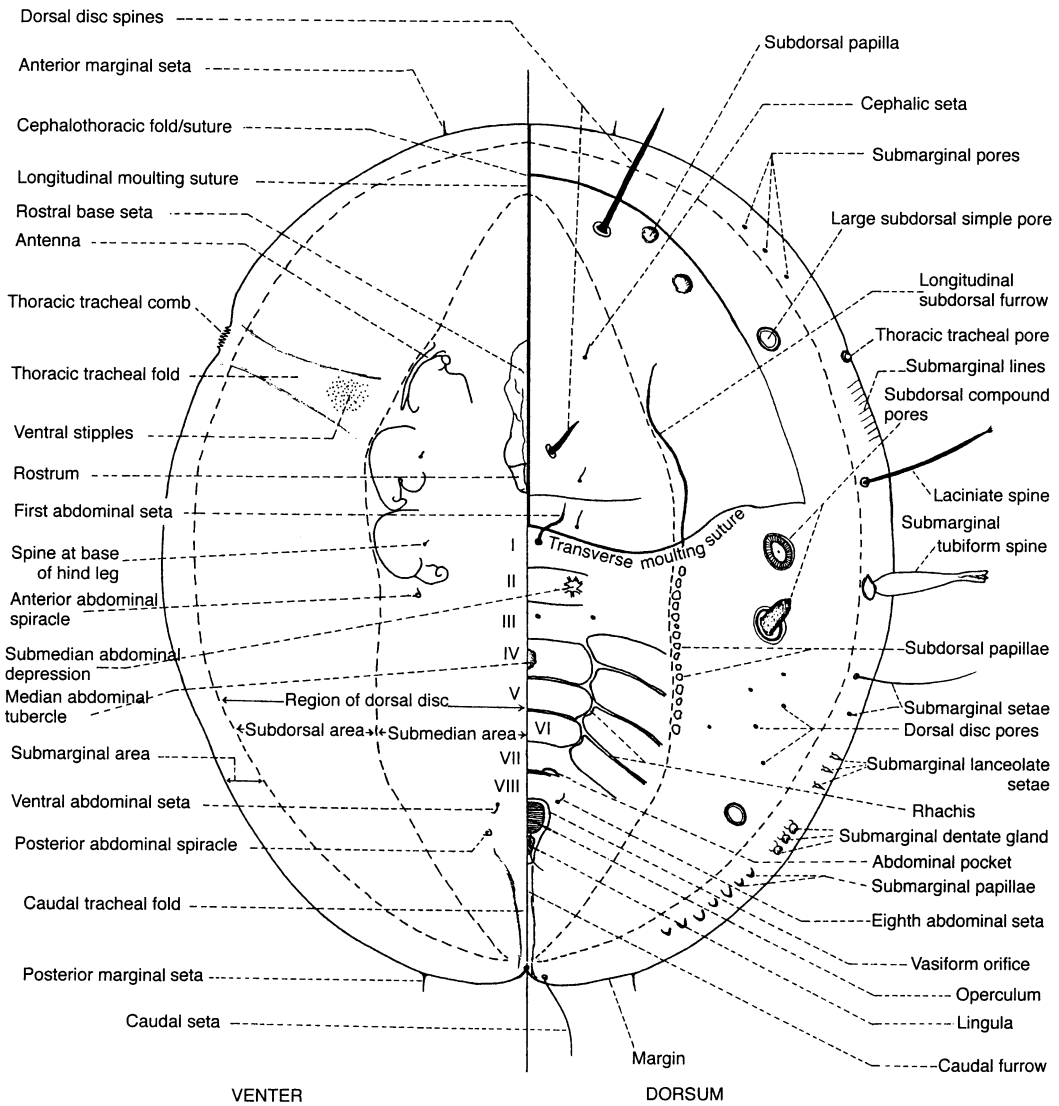


Figure 2. Stylized whitefly puparium with major morphological features annotated (from Martin 1987).

polymorphisms. Maximum parsimony analysis (MP) was carried out using PAUP*, version 4.0b10 (Swofford 1998). Heuristic searches were carried out with 1000 random additions followed by branch swapping using tree-bisection-reconnection (TBR) holding a single tree (NCHUCK=1, CHUCKSCORE=1) (Morris et al. 1999; Sallum et al. 2000; Hall and Harvey 2001; Skevington and Yeates 2001; Vardal et al. 2002). The resulting tree(s) were used for a final round of branch swapping with maxtrees 30 000. The new strategy for estimating large phylogenies (changing the landscape) was employed to help find more parsimonious trees (Quicke et al. 2001). The approach was used with maximum and minimum retention index (RI), and maximum and minimum consistency index (CI).

The bootstrap support for individual branches was estimated using 100 pseudoreplicates each of 100 random additions (Felsenstein 1985), though this will be very conservative due

to the difficult nature of this data set (e.g. Gauthier et al. 2000). Parsimony Jackknifing with 36.79% deletion, i.e. resampling 63.21% of characters, proposed by Farris et al. (1996) was also carried out to find support for branches as well as finding Bremer support (Bremer 1994) for some clades. To carry out Bremer support using PAUP*, a constraint tree description for the taxa of the clade in question was created and a search then performed by enforcing the topological constraint, but with the programme to only search for trees that were NOT compatible with that constraint. As a result, if the node in question is strongly supported, the best trees found that do not contain that node, i.e. are not compatible with the constraint, will be at least one step longer than the most parsimonious tree found from the initial unconstraint analysis. This discrepancy in tree length is the decay index or Bremer support value.

Additionally, successive approximation weighting (SAW) (Farris 1969) was applied for unweighted analysis using each of the four above-mentioned indices to increase resolution. Analyses were performed constraining some traditional whitefly genera to be monophyletic, and the resulting trees compared with the initial unconstrained ones using the Templeton (Wilcoxon signed-ranks) test implemented in PAUP*. The same procedure was used to find most-parsimonious trees (MPTs) for constraint analysis, i.e. heuristic searches with 1000 random additions, and employing the new strategy. Characters were traced on to cladograms using MacClade 4.0 (Maddison and Maddison 2000) and were optimised using the accelerated transformation algorithm (AccTran). In all cases, the obtained trees were rooted using the out-group taxa.

Characters and character coding using in the analysis

Pupal case

- [1] Pupal case margin: (0) smooth (Figure 3A); (1) undulate and/or indistinctly indentate (Figure 3B); (2) distinctly toothed (Figure 3C–L). Comments: in all cases, the word 'margin' is used to mean the real margin rather than the apparent margin, which may be submarginal in the case of marginal recurvature (see character 3). In other words, the real margin was used as a criterion to score the relevant characters.
- [2] If pupal case margin distinctly toothed: (0) crenate (Figure 3C); (1) dentate (Figure 3D); (2) lobulate (Figure 3E); (3) truncate–lobulate (Figure 3F, G); (4) serrate (Figure 3H); (5) each tooth with apical notch, which gives the teeth the appearance of being twinned and sometimes being triangular (Figure 3I–K); (6) close-set lanceolate processes, each arising from a blunt marginal tooth (Figure 3L).
- [3] Recurvature of margin: (0) margin not recurved; (1) margin partially recurved ventrally; (2) margin completely recurved ventrally. Comments: when the ventral surface of pupal case is much smaller than the dorsal surface the margin is evenly reflexed, a part of the submargin is recurved or the whole submargin is recurved. In some species the margin may easily be recurved by the mounting process and can cause misinterpretation when the character is coded.

Setae–spines

- [4] Anterior marginal setae: (0) absent; (1) present (Figure 2).

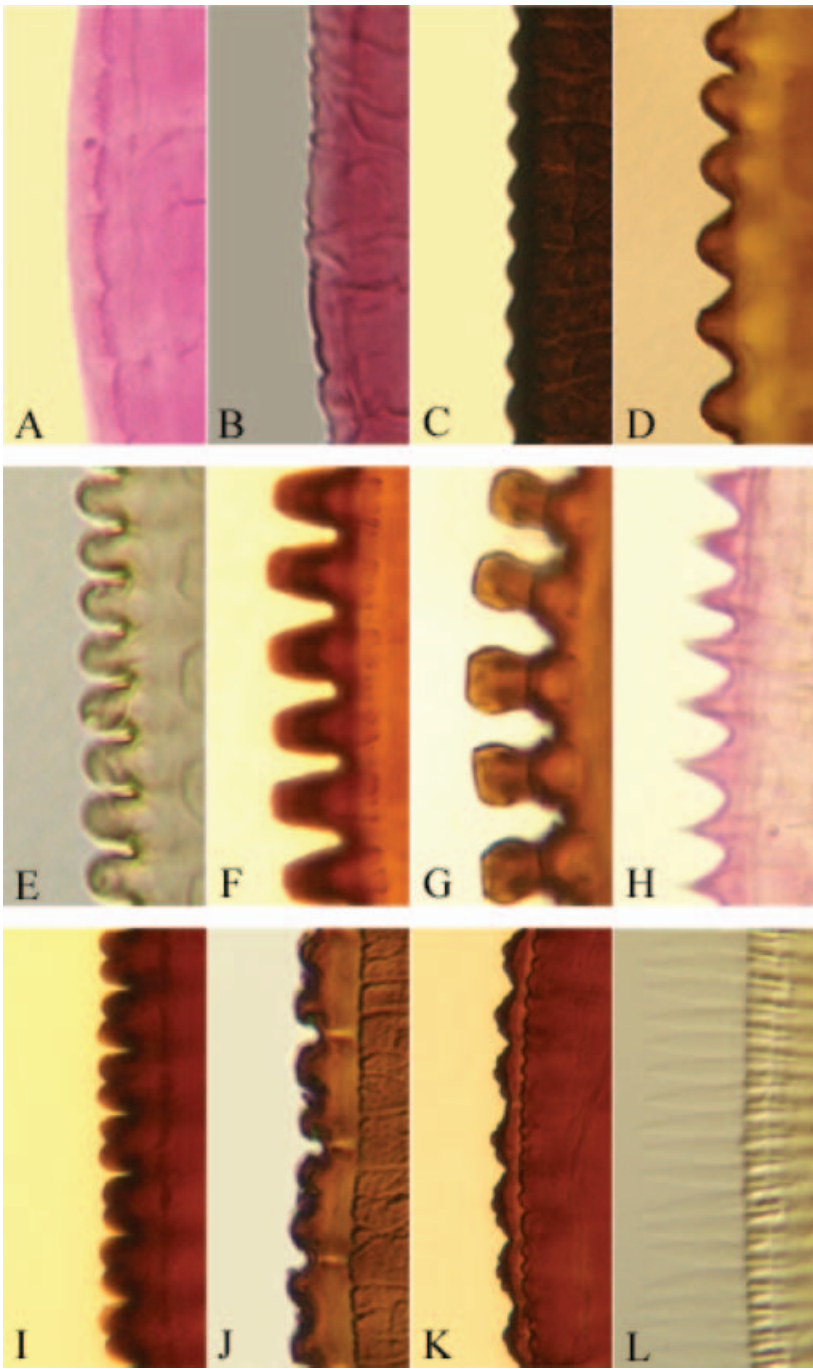


Figure 3. Different shapes of margin. (A) Smooth line; (B) undulate and/or indistinctly indentate; (C) crenate; (D) dentate; (E) lobulate; (F, G) truncate-lobulate; (H) serrate; (I–K), each tooth with apical notch, which gives the teeth the appearance of being twinned and sometimes being triangular; (L) close-set lanceolate processes, each arising from a blunt marginal tooth.

- [5] Posterior marginal setae: (0) absent; (1) present (Figure 2). Comments: although Bink-Moenen (1983) states they are probably always present, careful examination here at $\times 1000$ shows they are completely absent in some cases.
- [6] Cephalic setae: (0) absent; (1) present.
- [7] Position of cephalic setae: (0) nearly above, lateral or anterior to mouthparts; (1) far removed from mouthparts toward the front, margin, or both.
- [8] Mesothoracic setae: (0) absent; (1) present.
- [9] Metathoracic setae: (0) absent; (1) present.
- [10] First abdominal setae: (0) absent; (1) present.
- [11] Position of first abdominal setae: (0) nearly above, lateral or anterior to hind legs; (1) distinctly mediad of hind legs. Comments: the presence of the first abdominal setae can be of taxonomic value at genus level but in some genera there is interspecific variation (Bink-Moenen 1983). In the literature, the term 1st abdominal "pseudo-setae" is sometimes used for 1st abdominal setae when they are thickened, their bases submedial and often close-set (e.g. Martin 1999 for the genus *Aleuroduplidens* Martin). Their function then seems to be to keep the third-instar exuviae attached to the pupal case. Such setae are not always considered as 1st abdominal setae, e.g. for *Aleurocanthus* Quaintance & Baker, Bink-Moenen (1983) stated that "1st abdominal setae absent" but Martin (1999) stated "1st abdominal setae present but usually thickened, sometimes similar to glandular spines". In this study, these setae were accepted as 1st abdominal setae and mostly separated from the normal ones based on their position on the 1st abdominal segment.
- [12] Eighth abdominal setae: (0) absent; (1) present.
- [13] Position of eighth abdominal setae: (0) at the base of anterior margin and/or lateral to vasiform orifice (Figure 4A); (1) anterolateral to vasiform orifice (Figure 4B); (2) anterior to vasiform orifice (Figure 4C).
- [14] Caudal setae: (0) absent; (1) present.
- [15] Position of caudal setae: (0) close to margin; (1) on about half way from the posterior end of vasiform orifice to margin; (2) close to vasiform orifice.
- [16] Length of caudal setae: (0) distinctly shorter than vasiform orifice; (1) approximately as long as vasiform orifice; (2) distinctly longer than vasiform orifice. Comments: the length of caudal setae shows both interspecific and intraspecific variation in relation to the vasiform orifice, but appears to be fixed for the species of some genera.
- [17] Submarginal setae (excluding those in character 14): (0) absent; (1) present.
- [18] Submarginal setae (excluding those in character 14): (0) hair-like, short or long (Figure 5A); (1) stout, apical acute and/or rounded, short or long (Figure 5B); (2) somewhat swollen at base and pointed apically (lanceolate) (Figure 5C); (3) capitate (Figure 5D); (4) apical bended (Figure 5E); (5) laciniate (Figure 5F); (6) almost T-shaped (Figure 5H); (7) almost arrow-shaped (Figure 5I). Comments: see the comments for character 20.
- [19] Setae of dorsal disc area (except those in characters 6, 8, 9, 10 and 12): (0) absent; (1) present.
- [20] Setae of dorsal disc area (except those in characters 6, 8, 9, 10 and 12): (0) hair-like, short or long (Figure 5A); (1) stout, apical acute and/or rounded, short or long (Figure 5B); (2) somewhat swollen at base and pointed apically (lanceolate) (Figure 5C); (3) capitate (Figure 5D); (4) apical bended (Figure 5E); (5) laciniate (Figure 5F); (6) very stout, apical truncate-capitate (Figure 5G); (7) almost T-shaped (Figure 5H); (8) almost arrow-shaped (Figure 5I). Comments: the length

and position of setae as well as their number is variable within some species, e.g. *Aleurolobus gruvelli* Cotic (Bink-Moenen 1983). Except for a few genera, for which the position and number of setae are characteristic, e.g. *Corbettia* Dozier, their number and position have not usually been used as diagnostic characters at genus level (unlike for the aleurodicine genera). This is most probably due to variability so that even for a given number of setae for a species, it seems they are submarginal when small, but are located more subdorsally when they are enlarged. Therefore, the number of setae and their positions (except being on submargin or dorsal disc area) were not coded as separate characters. In the case of lack of submarginal line (see character 28), it is sometimes very difficult to assign setae as being submarginal or subdorsal when they are located far from the margin. In such cases, other features such as submarginal ridges radiating from the margin, colour differences or different ornamentations depending on the examined taxa were used to help define the submarginal limit. It is worth mentioning that according to Bink-Moenen (1983), the division of the dorsal surface into submargin, subdorsum, submedian and median areas (the three latter areas all together is called dorsal disc area) is artificial, as can be seen from the variable position of some setae in many species. Even when the submargin is separated by a furrow, the same setae may occur as often on the submargin as on the subdorsum.

- [21] Submarginal spines: (0) absent; (1) present. Comments: there is almost no distinct boundary between setae and spines in some cases in the literature and the same feature has been defined as a seta and/or a spine by different authors. In general, if there is no circle at the base of this character indicating a socket, it is considered as a spine. The length, position and number of spines are variable as mentioned above for the setae.
- [22] Submarginal spines: (0) stout and acute (Figure 5L); (1) stout and lacinate (Figure 5N); (2) tubiform (siphon-like) (Figure 5O).
- [23] Spines of dorsal disc area: (0) absent; (1) present.
- [24] Spines of dorsal disc area: (0) spines reduced to capitate seta-like structures (Figure 5K); (1) stout and acute (Figure 5L); (2) stout spines with a seta acutely pointed at the tip (Figure 5M); (3) stout and lacinate (Figure 5N); (4) tubiform (siphon-like) (Figure 5O).

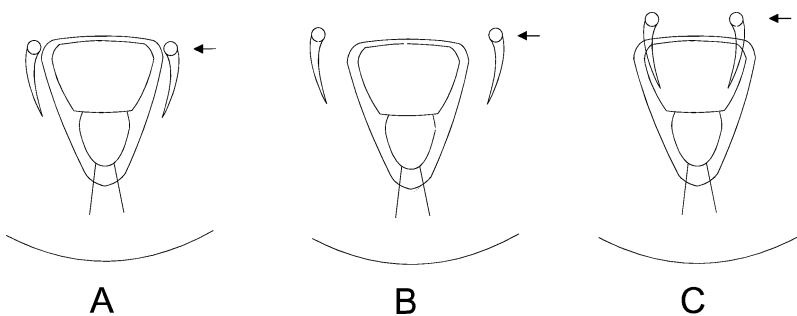


Figure 4. Position of eighth abdominal setae (arrowed). (A) At the base of anterior margin and/or lateral to vasiform orifice; (B) anterolateral to vasiform orifice; (C) anterior to vasiform orifice.

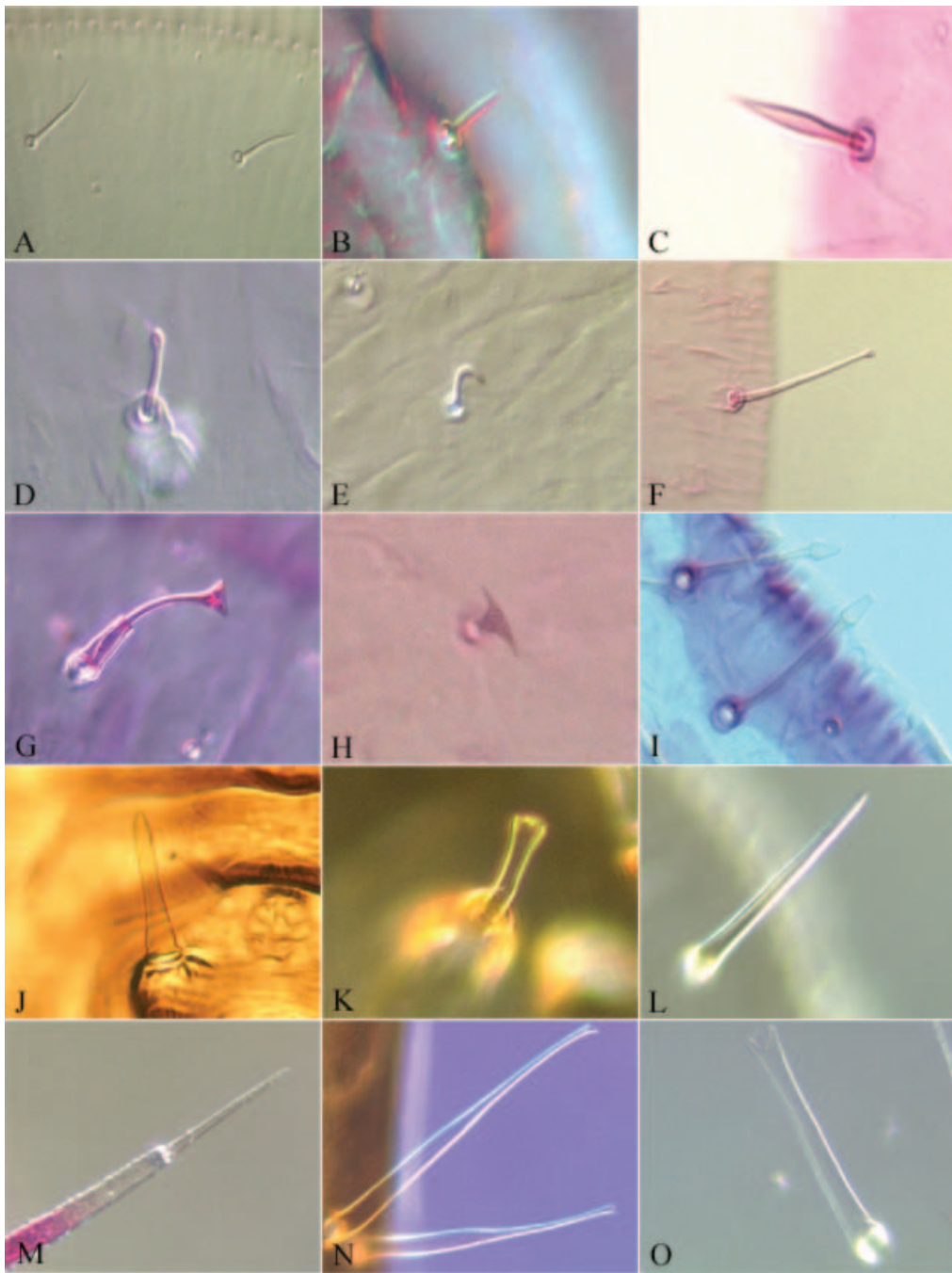


Figure 5. Different shapes of setae (A)–(J) and spines (K)–(O). (A) hair-like, short or long; (B) stout, apical acute and/or rounded, short or long; (C) somewhat swollen at base and pointed apically (lanceolate); (D) capitate; (E) apical bended; (F) lacinate; (G) very stout, apical truncate-capitate; (H) almost T-shaped; (I) almost arrow-shaped; (J) trumpet-like wax-secreting setae; (K) spines reduced to capitate seta-like structures; (L) stout and acute; (M) stout spines with a seta acutely pointed at the tip; (N) stout and lacinate; (O) tubiform (siphon-like).

Moulting sutures – major dorsal marking

- [25] Transverse moulting suture: (0) not reaching margin of pupal case; (1) reaching margin of pupal case.
- [26] Shape of transverse moulting suture: (0) like Figure 6A; (1) not bent near 90° in middle of each side (Figure 6B); (2) curves anteriorly at the junction of longitudinal moulting suture and posteriorly at the middle of each side (Figure 6C); (3) like Figure 6D; (4) each side curves anteriorly (Figure 6E); (5) like Figure 6F; (6) like Figure 6G; (7) each side bends to anterior and meets in the midline (Figure 6H); (8) each side curves anteriorly in the form of a pear together with a pair of membranous furrows (Figure 6I); (9) each sides curves anteriorly in the form of an almost triangular together with a pair of membranous furrows (Figure 6J). Comments: the shape of transverse moulting suture is quite variable and the 10 shapes chosen above were almost the main shapes and other intermediate states were assignable to one the main ones.
- [27] Longitudinal moulting suture: (0) not reaching margin of pupal case; (1) reaching margin of pupal case. Comments: only in one species studied, *Viennotaleyrodes curvisetosus* Martin, the longitudinal moulting suture was absent. For this species the character was coded as a question mark similar to those few species that the distal extremity of the suture was not clearly discernible due to slide quality.
- [28] Submarginal line or fold, which separates submargin from the dorsal disc area: (0) absent; (1) present.
- [29] If submarginal line or fold present: (0) parallel and longitudinal or almost bracket-shaped, not meeting at both anterior and posterior; (1) not parallel and longitudinal, meeting at both anterior and posterior, concentric with margin or nearly so; (2) like state 1, but with breaks at eyespot regions; (3) like state 1, but interrupted at caudal area; (4) like state 1, but with breaks at eyespot regions and interrupted at caudal area. Comments: in the literature, the shape of submarginal line or fold has sometimes been cited but almost never with clear diagnostic states separated or compared between taxa.
- [30] Extra submarginal lines encircling the pupal case: (0) absent; (1) present.
- [31] Longitudinal subdorsal furrow: (0) absent; (1) present.
- [32] Subdorsal line, which separates subdorsal area from submedian area: (0) absent; (1) present.
- [33] Cephalothoracic fold: (0) absent; (1) present. Comments: in some species, e.g. *Cockerelliella psidii* (Corbett), the cephalothoracic fold joins to the distal extremities of the transverse and longitudinal moulting sutures and forms a “trapdoor”. This feature is similar to character 26, state 7 but in the latter, each side of the transverse moulting suture curves anteriorly and is connected with the longitudinal moulting suture.

Vasiform orifice

- [34] Vasiform orifice shape: (0) semicircular; (1) nearly circular (Figure 7A); (2) nearly triangular (Figure 7E–J); (3) nearly tetrahedral (Figure 7K–M); (4) oval, truncate at the top (scutellate) (Figure 7B); (5) subcordate (Figure 7C); (6) transversely elliptical (Figure 7D). Comments: Maskell (1895) was the first to describe and name the vasiform orifice and its components the lingula, operculum and lingular setae. The shape of vasiform orifice is very variable and the same feature has been defined using different

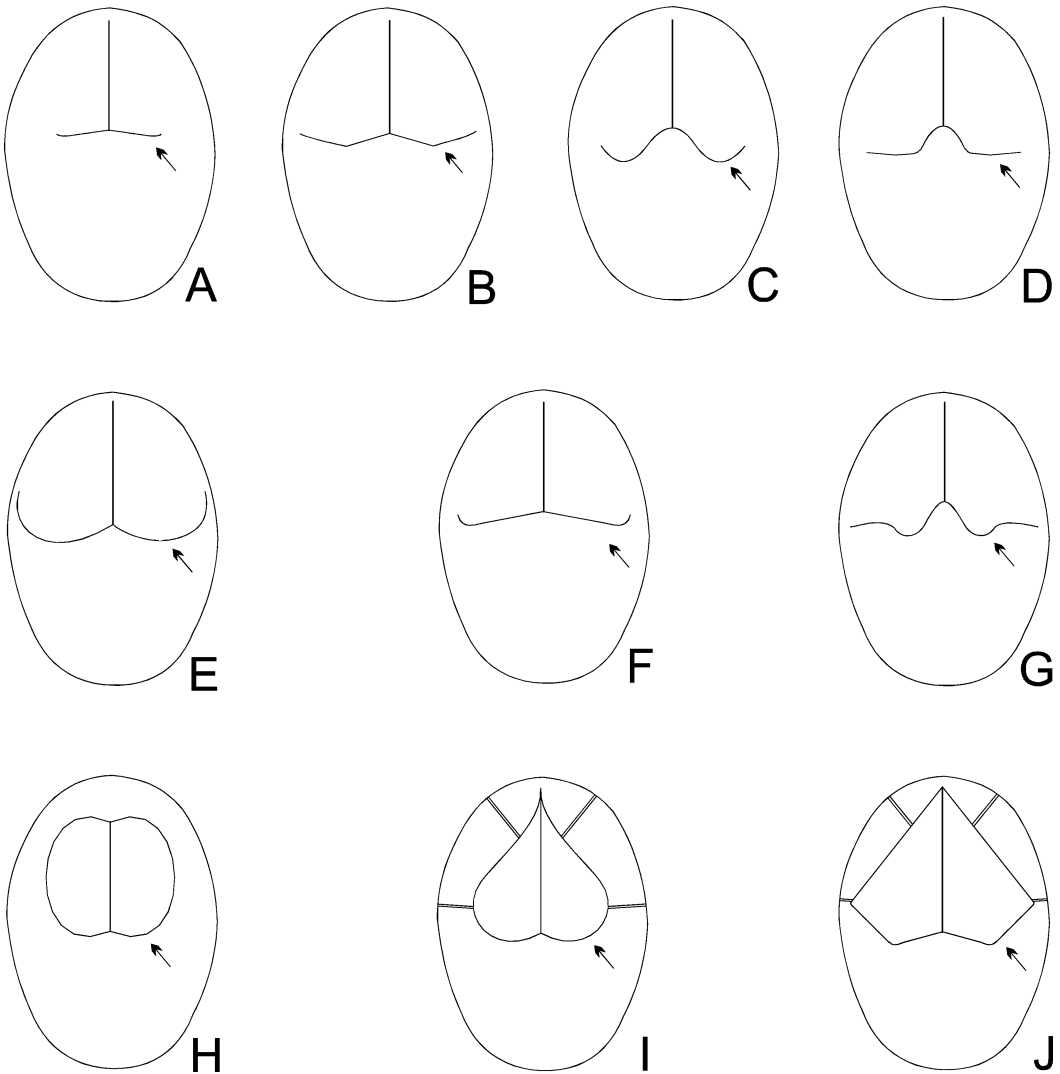


Figure 6. Different shapes of transverse moulting suture (arrow shows transverse moulting suture).

terms and/or short sentences by different authors. The shapes chosen in the characters 34–36 were the main shapes and other intermediate states were assignable to one of these states. The shape of vasiform orifice, at least in some cases, may be subjective but it has been used in many generic and species descriptions and so it has been included here and we have been as objective as possible in differentiating the states.

- [35] If vasiform orifice nearly triangular: (0) triangular, straight-sided (Figure 7E); (1) cordate (Figure 7F); (2) triangular, not very elongate, emarginate-sided (Figure 7G); (3) elongate-triangular, straight-sided (Figure 7H); (4) elongate–cordate (Figure 7I); (5) elongate-triangular, sinuous-sided (Figure 7J).
- [36] If vasiform orifice nearly tetrahedral: (0) squarish (Figure 7K); (1) rectangular (Figure 7L); (2) nearly trapezoidal (Figure 7M).

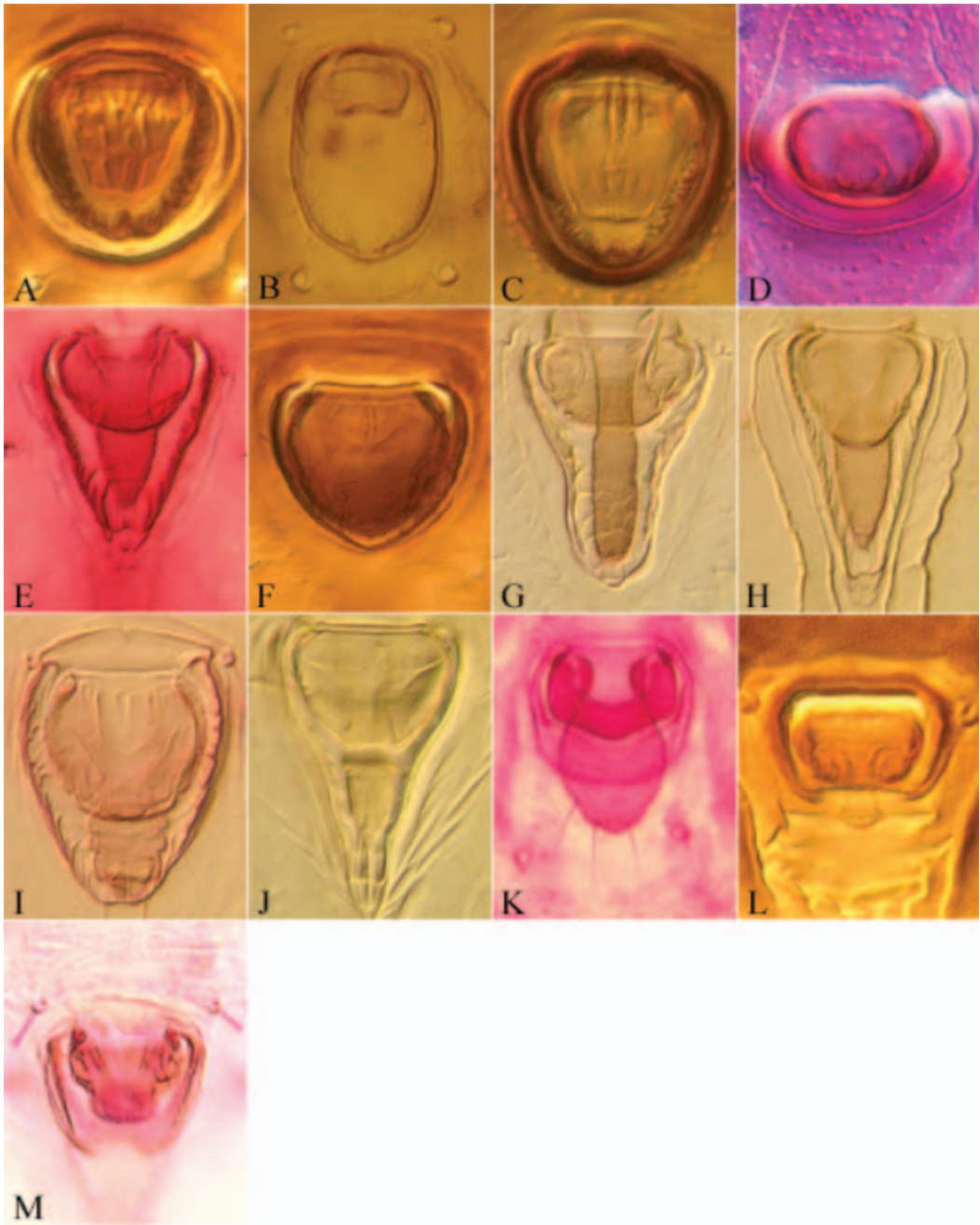


Figure 7. Vasiform orifice shape. (A) nearly circular; (B) oval, truncate at the top (scutellate); (C) subcordate; (D) transversely elliptical; (E) triangular, straight-sided; (F) cordate; (G) triangular, not very elongate, emarginate-sided; (H) elongate-triangular, straight-sided; (I) elongate-cordate; (J) elongate-triangular, sinuous-sided; (K) squarish; (L) rectangular; (M) nearly trapezoidal.

- [37] Vasiform orifice: (0) not elevated; (1) elevated. Comments: although in some genera e.g. *Aleurocanthus*, this character is easily recognisable, but it is sometimes less obvious. For example, for the genus *Siphoninus* Silvestri, Martin (1999) stated that “vasiform orifice not elevated”, which is the opposite of what Bink-Moenen (1983) stated, i.e. “vasiform orifice elevated”. The latter author defined two major types of vasiform orifice: “open” vasiform orifice and “elevated” vasiform orifice. To recognize the latter type she stated that “the elevated vasiform orifice is situated on an elevation; its margin is often surrounded by a sclerotized ring; caudal furrow and ridges are absent”. In this study, it has been found that the absence of caudal furrow and ridges is not always true and in some species for example, in *Dialeurolonga elongata* Dozier and *Zaphanera papyrocarpae* Martin, the vasiform orifice is elevated and the caudal furrow and ridges are present. Elevation of the vasiform orifice is probably not generically important but is important at the species level (J. H. Martin, BMNH, personal communication).
- [38] Vasiform orifice inner margin teeth (carinae): (0) absent; (1) present.
- [39] Vasiform orifice posterior notch: (0) absent; (1) present. Comments: although the shape of notch also shows variation, this variation appears to be continuous rather than discrete. As far it is known from the literature, with the exception of Jensen’s articles (1999, 2001), the shape of vasiform orifice notch has never been used as a diagnostic character, at least at genus level.
- [40] Position of vasiform orifice: (0) inset from posterior margin of pupal case by less than its own length; (1) inset from posterior margin of pupal case by about its own length; (2) inset from posterior margin of pupal case by distinctly more than its own length (sometimes two or more times).
- [41] Operculum size: (0) occupying about 0.25 of orifice; (1) occupying about 0.5 of orifice; (2) occupying about 0.75 of orifice; (3) occupying about whole of orifice. Comments: there are two types of vasiform orifice: the orifice is nearly or completely filled by the operculum and the lingula is normally not visible in dorsal view, and in the other type, the operculum does not fill the posterior part of the orifice and the lingula is usually visible (Gill 1990). Here, these two main states were divided into four states so as to cover all those used by various authors in generic and/or species descriptions to date.
- [42] Operculum shape: (0) transversely elliptical (Figure 8A); (1) nearly trapezoidal, rounded at angles (Figure 8B); (2) nearly rectangular, wide about twice as long as length, posterior angles toothed and/or slightly indentate (Figure 8C); (3) subcordate (Figure 8D); (4) semicircular (Figure 8E); (5) rectangular, convex-sided, hind margin straight and/or slightly curving anteriorly (Figure 8F); (6) rectangular, convex-sided, hind margin with a wide projection (Figure 8G); (7) cordate or slightly elongate-cordate (Figure 8H); (8) rectangular shaped pushed inward at posterior margin (Figure 8I); (9) squarish, convex-sided, narrowed posteriorly (Figure 8J). Comments: as with the vasiform orifice, the shape of the operculum is also variable and the same feature has been defined using different terms and/or short sentences used by different authors (see comments for character 34). This character has not been used as frequently as the shape of vasiform orifice in species or generic descriptions.
- [43] Opercular setae: (0) absent at posterior margin of the operculum; (1) present at posterior margin of the operculum (two setae).
- [44] Head of lingula: (0) completely obscured by opaque operculum, not visible; (1) completely obscured by transparent operculum, visible; (2) incompletely obscured by operculum, partly visible; (3) not obscured by operculum.
- [45] Head of lingula: (0) elongate (Figure 9H–J); (1) not elongate (Figure 9A–G, K).

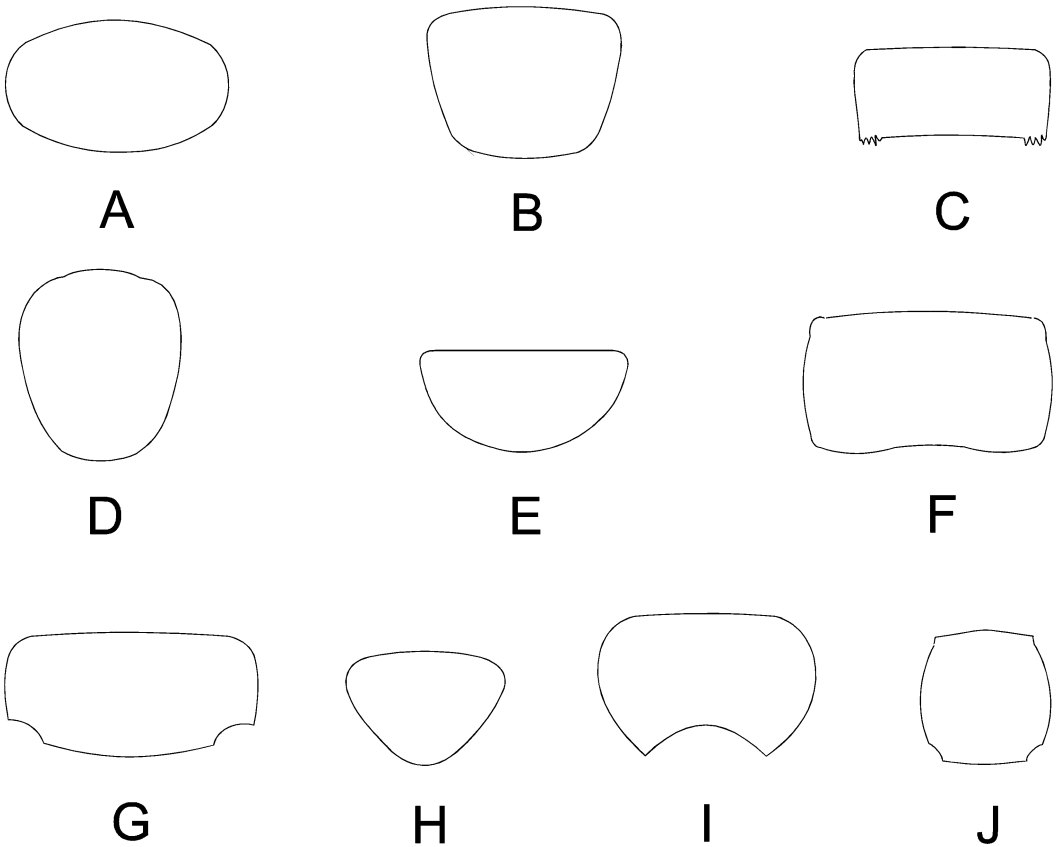


Figure 8. Different shapes of the operculum. (A) transversely elliptical; (B) nearly trapezoidal, rounded at angles; (C) nearly rectangular, wide about twice as long as length, posterior angles toothed and/or slightly indented; (D) subcordate; (E) semicircular; (F) rectangular, convex-sided, hind margin slightly curves anteriorly and/or straight; (G) rectangular, convex-sided, hind margin with a wide projection; (H) cordate or slightly elongate-cordate; (I) rectangular shaped pushed inward at posterior margin; (J) squarish, convex-sided, narrowed posteriorly.

[46] If head of lingula elongate: (0) lanceolate (Figure 9H); (1) cylindrical, rounded posteriorly (Figure 9I); (2) large and almost conical (Figure 9J).

[47] If head of lingula not elongate: (0) blunt apex exposed; (1) expanded apically (Figure 9B); (2) spatulate, sometimes nearly triangular (Figure 9G); (3) forked-shaped (Figure 9D); (4) lobular (Figure 9E); (5) large and tongue-shaped (Figure 9K); (6) bilobate (Figure 9C); (7) knobbed (Figure 9A); (8) transversely elliptical (Figure 9F). Comments: the same as for the vasiform orifice and operculum but the character shows some discrete states (e.g. states 3 and 6).

[48] Lingula: (0) not extending beyond posterior margin of vasiform orifice; (1) extending beyond posterior margin of vasiform orifice. Comments: sometimes the lingula may be protruded by the mounting process and can cause misinterpretation this character. It should be noted that in some species the length of lingula is longer than vasiform orifice but it folds inside the orifice. To avoid any error in determining its true length in such species, the normal state of lingula was chosen, i.e. the character was coded as having the first state (not extending beyond posterior margin of vasiform orifice).

[49] Lingular setae: (0) absent or not apparent; (1) present.

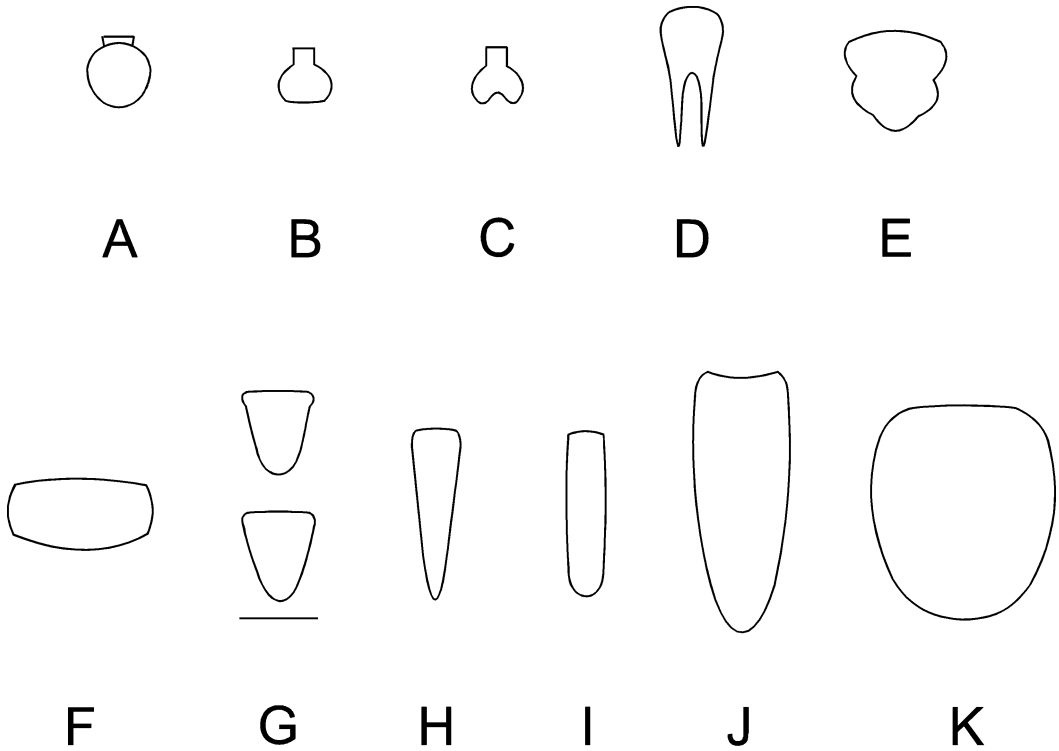


Figure 9. Different shapes of lingula. (A) knobbed; (B) expanded apically; (C) bilobate; (D) forked-shaped; (E) lobular; (F) transversely elliptical; (G) spatulate, sometimes nearly triangular; (H) lanceolate; (I) cylindrical, rounded posteriorly; (J) large and almost conical; (K) large and tongue-shaped.

[50] Number of lingular setae: (0) two (if two more setae are present, these are minute and often visible only at $\times 1000$ magnification with a light microscope); (1) four. Comments: the lingula is usually spinulose and the number of conspicuous setae generally varies from without any seta to having four setae. In the Aleurodicinae, some of whose members were used as out-groups, the lingula almost always has four conspicuous setae. In the Aleyrodinae, only two apical setae have usually been mentioned in original descriptions. Russell (1947) was the first to report the presence of two more small lateral lingular setae (for some species of *Trialeurodini* that she described as a new tribe including three genera (see Appendix A)). In this study, examination of thousands specimens belonging to about one third of all described whitefly species at $\times 1000$ magnification indicates that with the exception of lack of lingular setae in some species, these two small lateral setae are also present in other genera and quite visible when the lingula is in a right position. It seems that they have been neglected, even by Russell in her other publications, because they are small and difficult to see at least at $\times 400$. However, since these two small lingular setae somehow differ from those of the Aleurodicinae, the character was included here.

Tracheal pores–tracheal folds

[51] Thoracic tracheal pores: (0) no discernible pore; (1) not forming a distinct pore, margin slightly curved inwardly/outwardly or thickened (Figure 10A); (2) somewhat

invaginated and with slightly thickened cuticle (Figure 10B); (3) crenulated to a lesser degree than the rest of margin (Figure 10C); (4) in the form of a comb of modified teeth, sometimes only two or three teeth (Figure 10D–F); (5) C-shape pore without teeth (Figure 10G, H); (6) C-shape with teeth (Figure 10I, J); (7) deeply emarginate and covered by a smooth rounded lobe (Figure 10K); (8) gland spines forming a comb (Figure 10L). Comments: each state shows variations even within some species, so for such a large-scale study it was almost impossible to score all numerous visible states that did not seem to be generic and even sometimes species level characters. The eight states coded are as objective as possible.

- [52] Thoracic tracheal folds: (0) absent; (1) present. Comments: in some species tracheal folds are finely stippled or defined by microtubercles.
- [53] Caudal tracheal fold: (0) absent; (1) present. Comments: the same as for the character 52.
- [54] Secondary tracheal folds underline abdominal sutures III/IV: (0) absent; (1) present. Comments: this character was only observed in one monobasic genus, *Gagudjuia* Martin. In the future the character may be seen in other species, so it was added to the character system to help for further studies, though in this study it is parsimony-uninformative.
- [55] Caudal furrow: (0) absent; (1) present.
- [56] Caudal furrow: (0) narrow, half-width or less than half-width of vasiform orifice, sometimes suture-like; (1) wide at base (near vasiform orifice), the rest part narrow; (2) wide, more than half-width of vasiform orifice; (3) narrow at base, the rest part wide.
- [57] Caudal ridges: (0) absent; (1) present.

Abdominal segments

- [58] Lengths of seventh abdominal segment: (0) length of abdominal segment VII not significantly (half-length or less than half-length of segment VI) reduced medially; (1) length of abdominal segment VII significantly reduced medially. Comments: the length of abdominal segment VII, comparing to abdominal segment VI, is quite variable but the reduction of its length medially has been used as a diagnostic character for some genera, e.g. *Bemisia* and *Pealius* Quaintance & Baker, without clearly defining that how much reduction is significant. To make it clear and practical to score the character and subsequently to justify its validity as being a diagnostic character, the above-mentioned criterion was used.
- [59] Submedian abdominal depressions: (0) absent; (1) present.
- [60] Median abdominal tubercles: (0) absent; (1) present; (2) tubercle-like markings.
- [61] Abdominal rhachis: (0) absent; (1) present.
- [62] Lateral arms arising from abdominal rhachis: (0) absent; (1) present.
- [63] Eighth abdominal bifid process: (0) absent; (1) present (Figure 11A).
- [64] Segmentation between the abdominal segments: (0) normal; (1) first three abdominal segments joined together, apparently forming a single structure.

Papillae–tubercles–glands–pores

- [65] Submarginal papillae: (0) absent; (1) present (Figure 11B). Comments: there is no clear distinction between papillae and tubercles in most cases in the literature. Their

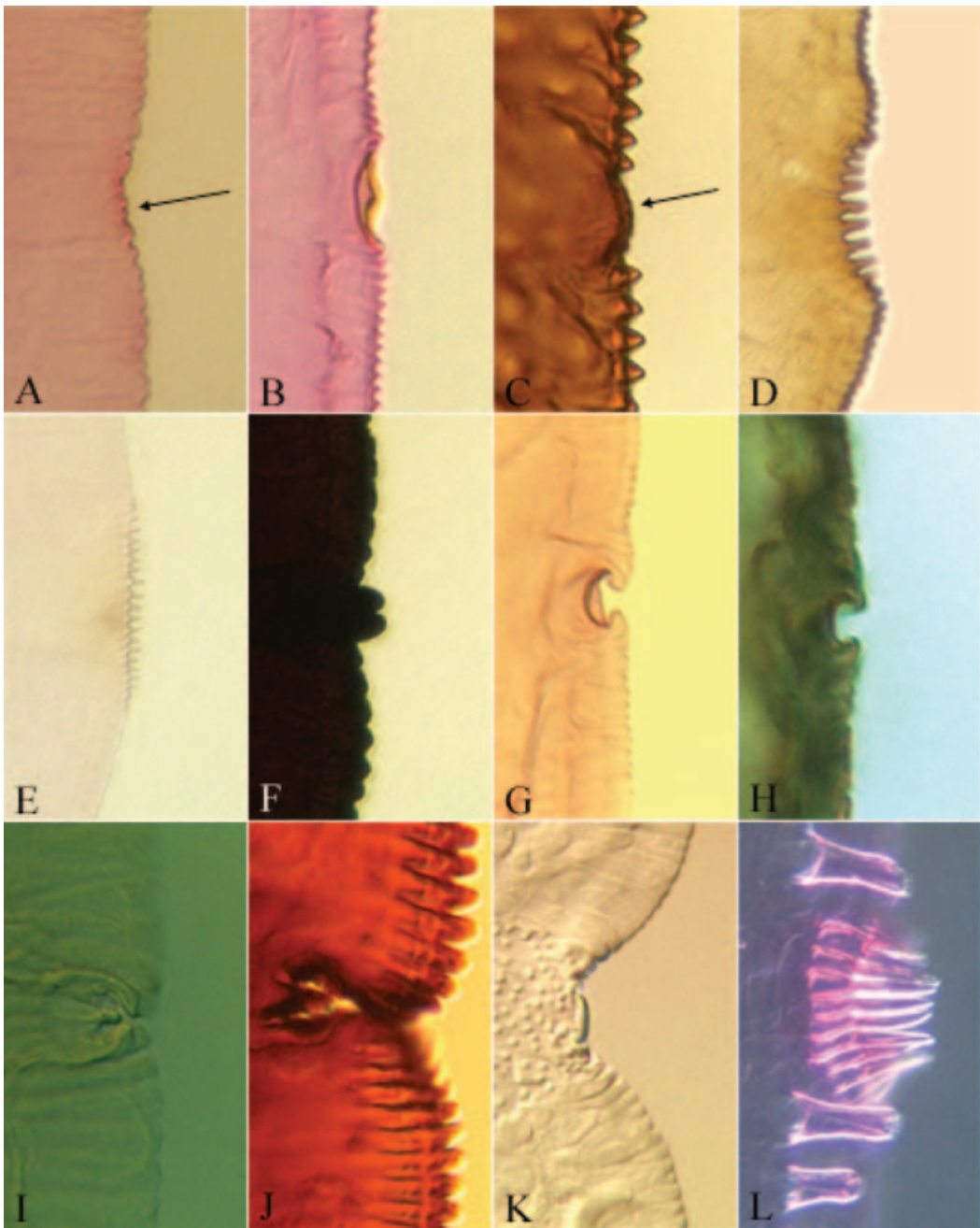


Figure 10. Different forms of the tracheal pore. (A) Not forming a distinct pore, margin slightly curved inwardly/ outwardly or thickened (arrowed); (B) somewhat invaginated and with slightly thickened cuticle; (C) crenulated to a lesser degree than the rest of margin (arrowed); (D)–(F) a comb of modified teeth, sometimes only two or three teeth; (G, H) C-shape pore without teeth; (I, J) C-shape with teeth; (K) deeply emarginate and covered by a smooth rounded lobe; (L) gland spines forming a comb.

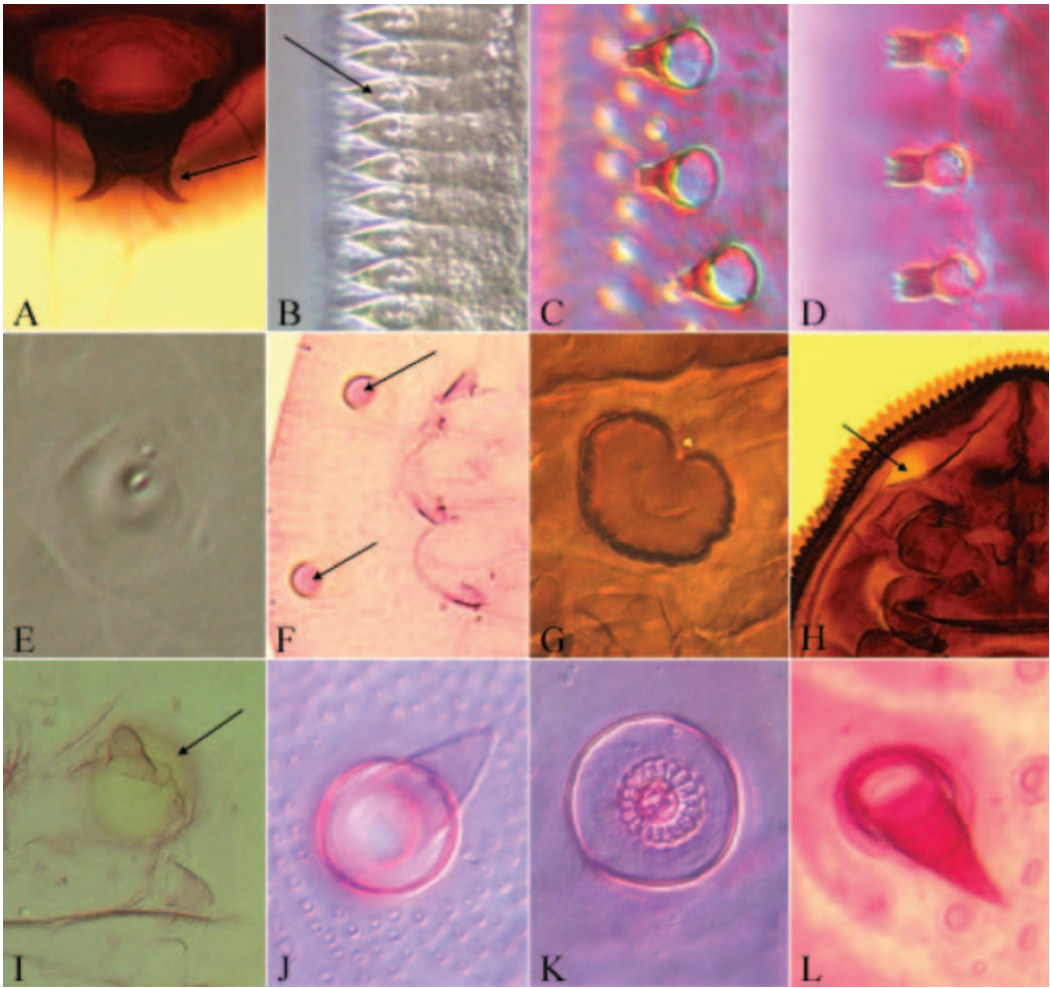


Figure 11. Different characters of pupal cases. (A) eighth abdominal segment showing bifid process (arrowed); (B) submarginal papillae (arrowed); (C) submarginal glands (tips of glands smooth); (D) submarginal glands (tips of glands dentate); (E) geminate pore/porettes; (F) large subdorsal simple pores (arrowed); (G) modified papillae of dorsal disc area, which are slightly elevated, nearly flat and plate-like; (H) eyespot (arrowed); (I) dorsal gland (arrowed); (J)–(L) compound wax pores.

presence, size and structure are very variable even within a species. Therefore, only the presence and absence of these characters on the submargin (this character and character 66) or on the dorsal disc area (characters 74 and 75) was scored. Compared with papillae, tubercles are more shapeless, usually irregularly distributed and have no associated pore. According to J. H. Martin (BMNH, personal communication), tubercles are often environmentally induced characters.

[66] Submarginal tubercles (excluding the tubercle between caudal setae, when is present, and those at the bases of the larger setae): (0) absent; (1) present. Comments: see character 65.

[67] Submarginal glands: (0) absent; (1) present (Figure 11C, D). Comments: these structures were named by Russell (1958), thought they have also been called papillae, pores, or tubercles by other writers.

- [68] Shape of submarginal glands: (0) tips of glands smooth (Figure 11C); (1) tips of glands dentate (Figure 11D). Comments: the tips of the glands in the ten species described/redescribed by Russell (1958) were dentate. Later, Bink-Moenen (1983) described such glands for the monobasic genus, *Arachnaleyrodes* Bink-Moenen, but in that case their tips were smooth, and was used as one of the diagnostic character for the genus. Glands with smooth tip were only observed in one taxon, so this parsimony-uninformative character was automatically ignored by the analyses.
- [69] Submarginal simple pores: (0) absent; (1) present.
- [70] Submarginal geminate pore/porettes: (0) absent; (1) present (Figure 11E). Comments: these structures are also called combined pores and sometimes pores and porettes are widely spaced from each other. According to Russell (1948) and Bink-Moenen (1983), their distribution is of some value at the species level.
- [71] Simple pores of dorsal disc area: (0) absent; (1) present.
- [72] Geminate pore/porettes of dorsal disc area: (0) absent; (1) present (Figure 11E). Comments: the same as for the character 70.
- [73] Large subdorsal simple pores: (0) absent; (1) present (Figure 11F). Comments: these structures are usually a generic level diagnostic feature but at least in *Dialeuropora decempuncta* (Quaintance & Baker) these large simple pores are sometimes absent or much reduced in size. It is considered likely that this is a modification arising from environmental factors which remain obscure (Martin 1999).
- [74] Subdorsal papillae: (0) absent; (1) present. Comments: the same as for character 65.
- [75] Tubercles of dorsal disc area (except the tubercles mentioned in the character 60): (0) absent; (1) present. Comments: the same as for character 65.
- [76] Modified papillae of dorsal disc area, which are slightly elevated, nearly flat and plate-like: (0) absent; (1) present (Figure 11G).
- [77] Eyespots: (0) absent; (1) present (Figure 11H). Comments: according to Bink-Moenen (1983), the presence of eyespots can be of taxonomic value at the species level, but presence and expression are often variable interspecifically. Both states were scored for those species that at least some individuals with them.
- [78] Dorsal glands: (0) absent; (1) present (Figure 11I). Comments: these glands are easily distinguished from those mentioned in the characters 67 and 68 (see Figure 11C, D, I).
- [79] Subdorsal compound wax pores and/or agglomerate pores: (0) absent; (1) present (Figures 11J–L and 12A, B). Comments: only members of the subfamily Aleurodicinae that have been chosen as out-groups for this study have compound wax pores and/or agglomerate pores. For this reason, these characters were not treated separately.
- [80] Glandular bases to marginal teeth: (0) absent; (1) present (Figure 12C). Comments: in the literature, this structure has sometimes been described as “a double row of teeth” or rarely “a translucent membranous area at the bases of the marginal teeth”. According to Bink-Moenen (1983), these are probably wax-secreting pores or papillae.
- [81] Trumpet-like wax-secreting setae: (0) absent; (1) present (Figure 5J). Comments: these setae were considered by Tremblay and Iaccarino (1978) as a generic character for the monobasic genus, *Aleurotuba* Tremblay & Iaccarino. Because they were observed only in this single taxon, it is parsimony-uninformative.
- [82] Chain-like design along some abdominal and thoracic sutures: (0) absent; (1) present (Figure 12D).
- [83] Submedian conspicuous invaginations; sometimes hemispherical: (0) absent; (1) present (Figure 12E, F).

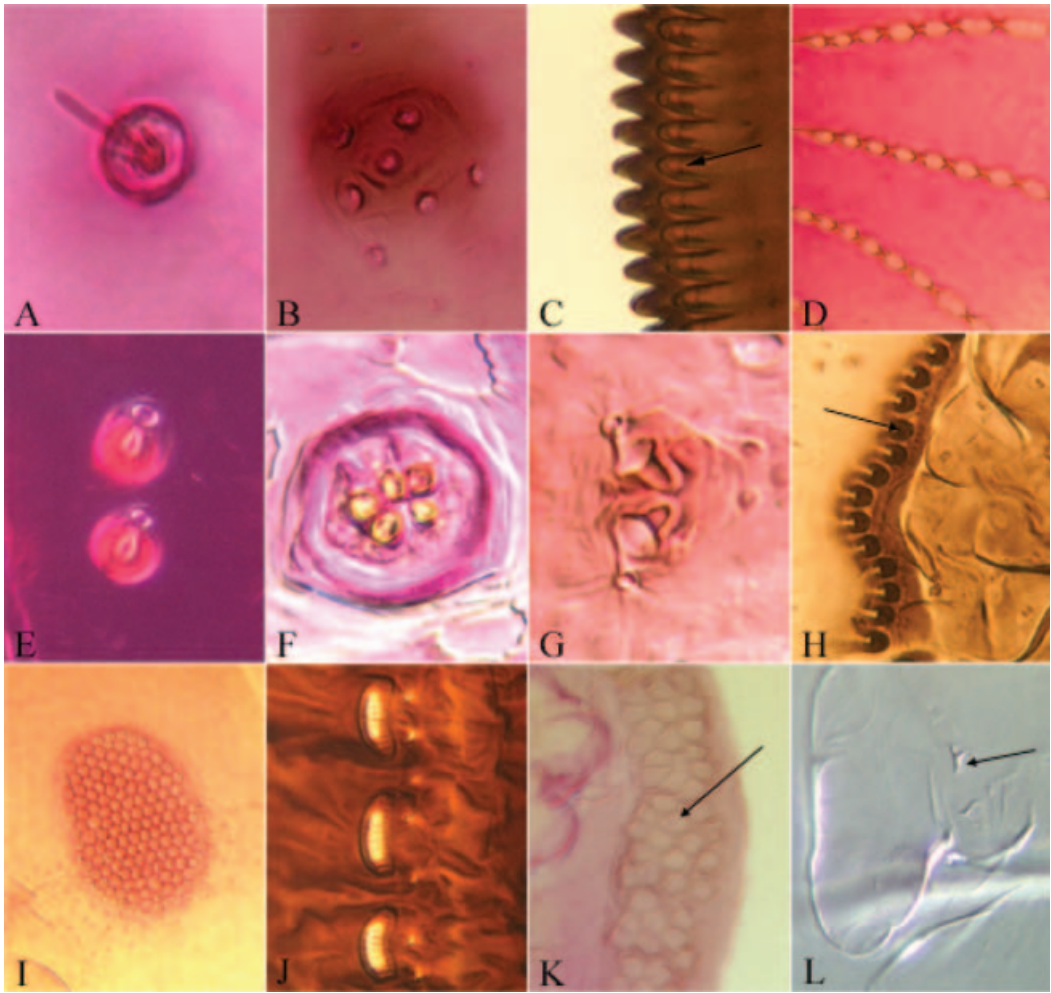


Figure 12. Different characters of pupal cases. (A, B) Compound wax pores and/or agglomerate pores; (C) glandular bases to marginal teeth (arrowed); (D) chain-like design along some abdominal and thoracic sutures; (E, F) submedian conspicuous invaginations; sometimes hemispherical; (G) subdorsal plate with conical pores; (H) a row of scallop-shaped thickenings along the longitudinal subdorsal furrow (arrowed); (I) prominent glandular areas with polygonal reticulate pattern, sometimes resembling compound eyes; (J) crescent-shaped pores; (K) submarginal wax-plat cluster (arrowed); (L) basal leg spine (arrowed).

- [84] Subdorsal plate with one or two conical pores connected with tracheal pore area: (0) absent; (1) present (Figure 12G).
- [85] A row of scallop-shaped thickenings along the longitudinal subdorsal furrow: (0) absent; (1) present (Figure 12H).
- [86] Prominent glandular areas with polygonal reticulate pattern, sometimes resembling compound eyes: (0) absent; (1) present (Figure 12I).
- [87] Submarginal crescent-shaped pores: (0) absent; (1) present (Figure 12J).
- [88] Submarginal wax-plat cluster: (0) absent; (1) present (Figure 12K).

Legs and antennae

- [89] Apical adhesion pad of legs: (0) absent; (1) present.
- [90] Tarsal claw: (0) absent; (1) present.
- [91] Basal mid- and hind leg seta: (0) absent; (1) present. Comments: this character has sometimes been used in species and/or generic descriptions, especially whenever the seta(e) have been long enough to be easily recognisable. In some cases, there is no clear distinction between setae and spines (character 92) and, apparently, the terms have been used synonymously by some authors. For example, for *Acaudaleyrodes africana* (Dozier), Mound (1965) stated “*A. africanus* can be distinguished from the other species in *Acaudaleyrodes* by the absence of setae from the base of legs, ...”. Based on the criterion given in the comments for the character 21, what are absent from the base of legs in this species are spines not setae. Here, examination of thousands of specimens at $\times 1000$ magnification, clearly showed that at least one microseta is always present at the base of each mid- and hind leg. Sometimes one or rarely two or more setae can be seen in some species but because of slide quality it was almost impossible to count them precisely for each species for such a large-scale study. It seems that the existence of extra seta(e) at most can be important at species level, but it is sometimes variable even for the same species. This character was automatically ignored due to being always present, i.e. a constant character.
- [92] Basal mid- and hind leg spine: (0) absent; (1) present (Figure 12L).
- [93] Antenna length: (0) limited to front legs or slightly more; (1) reaching middle of mid legs or more. Comments: sexual dimorphism occurs in some genera and species as a difference in the length of the antenna. In this case, the antennae of males are comparatively longer than the antennae of females. Such species were scored as polymorphic.
- [94] Position of antenna relative to the prothoracic legs: (0) mesal to the prothoracic legs; (1) lateral to the prothoracic legs.

Results*Parsimony analysis*

Maximum parsimony holding a single tree yielded one MPT (length=2759, RI=0.668, CI=0.136). This shortest tree was hit only once during the 1000 random additions, showing that this is a rather difficult data set to analyse. Branch swapping with maxtrees set at 30 000 using this tree as the starting tree gave 30 000 equally MPTs with a length of 2743 (RI=0.671, CI=0.137). Application of the new strategy (changing the landscape) for these trees using maximum RI to weight characters found 30 000 (limited by computer memory) equally MPTs with a length of 2730 (RI=0.672, CI=0.137) (employing the same strategy using maximum CI, minimum CI or minimum RI to weight characters found no shorter trees). The strict consensus tree of these is shown in Figure 13. Although the strict consensus tree contains several monophyletic clades, the relationships among most of these are unresolved. Ignoring 44 of the 117 examined genera that are monobasic and/or only one species were available for study, the following 33 genera (the author’s names of all genera are given in Appendix C) were found to be monophyletic: *Acanthaleyrodes*, *Acaudaleyrodes*, *Africaleurodes*, *Aleurocerus*, *Aleurocybotus*, *Aleuroduplidens*, *Aleuroglandulus*, *Aleuroparadoxus*, *Aleuropteridis*, *Bellitudo*, *Combesaleyrodes*, *Corbettia*, *Dialeuropora*, *Fascaleyrodes*, *Filicaleyrodes*, *Fippataleyrodes*, *Indoaleyrodes*, *Leucopogonella*, *Metabemisia*,

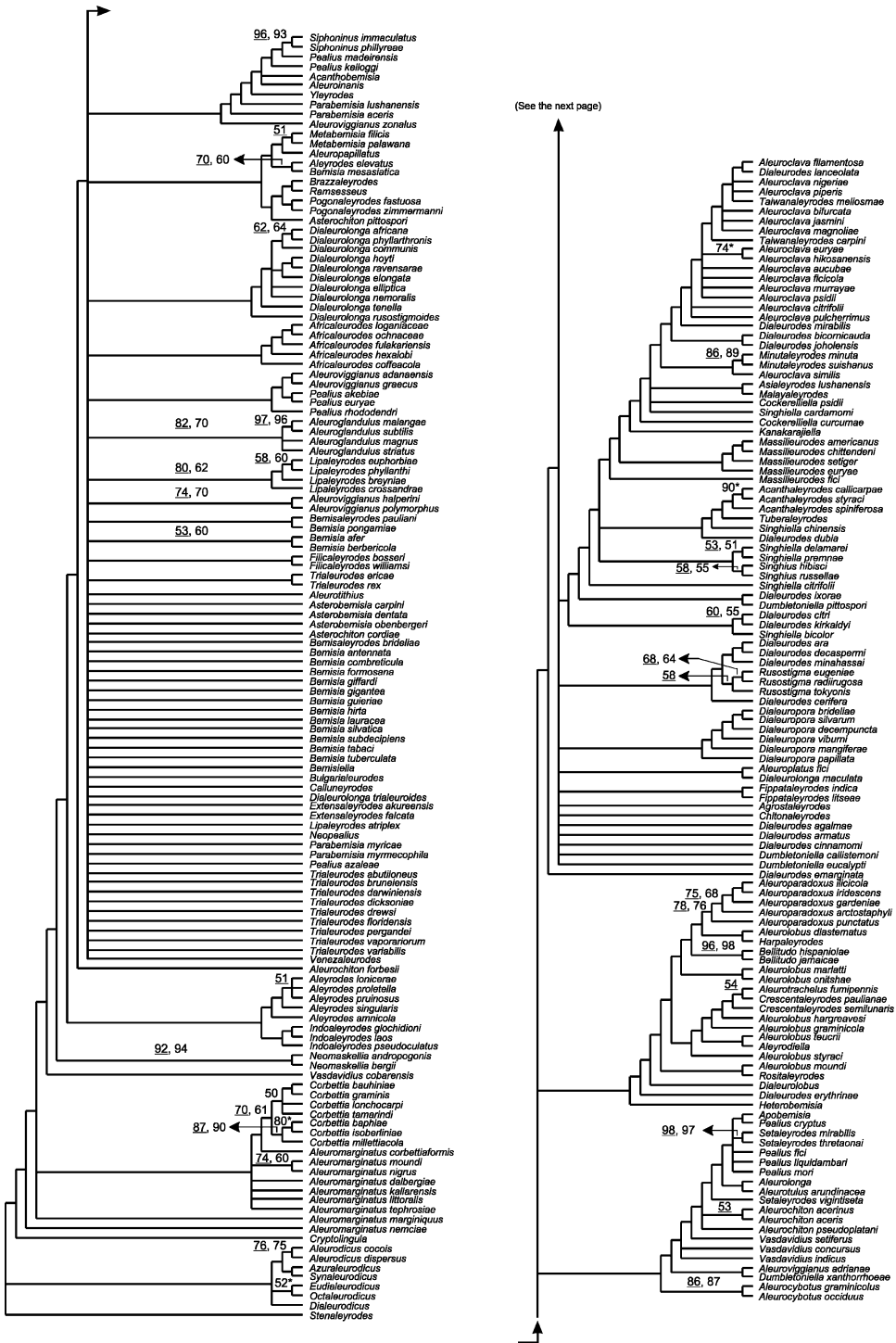


Figure 13. Strict consensus tree of 30 000 UW-MPTs resulting from analyses of morphological data. Figures above branches show bootstrap and Jackknife (underlined figures) values. Figures with an asterisk indicate the same value for both bootstrap and Jackknife.

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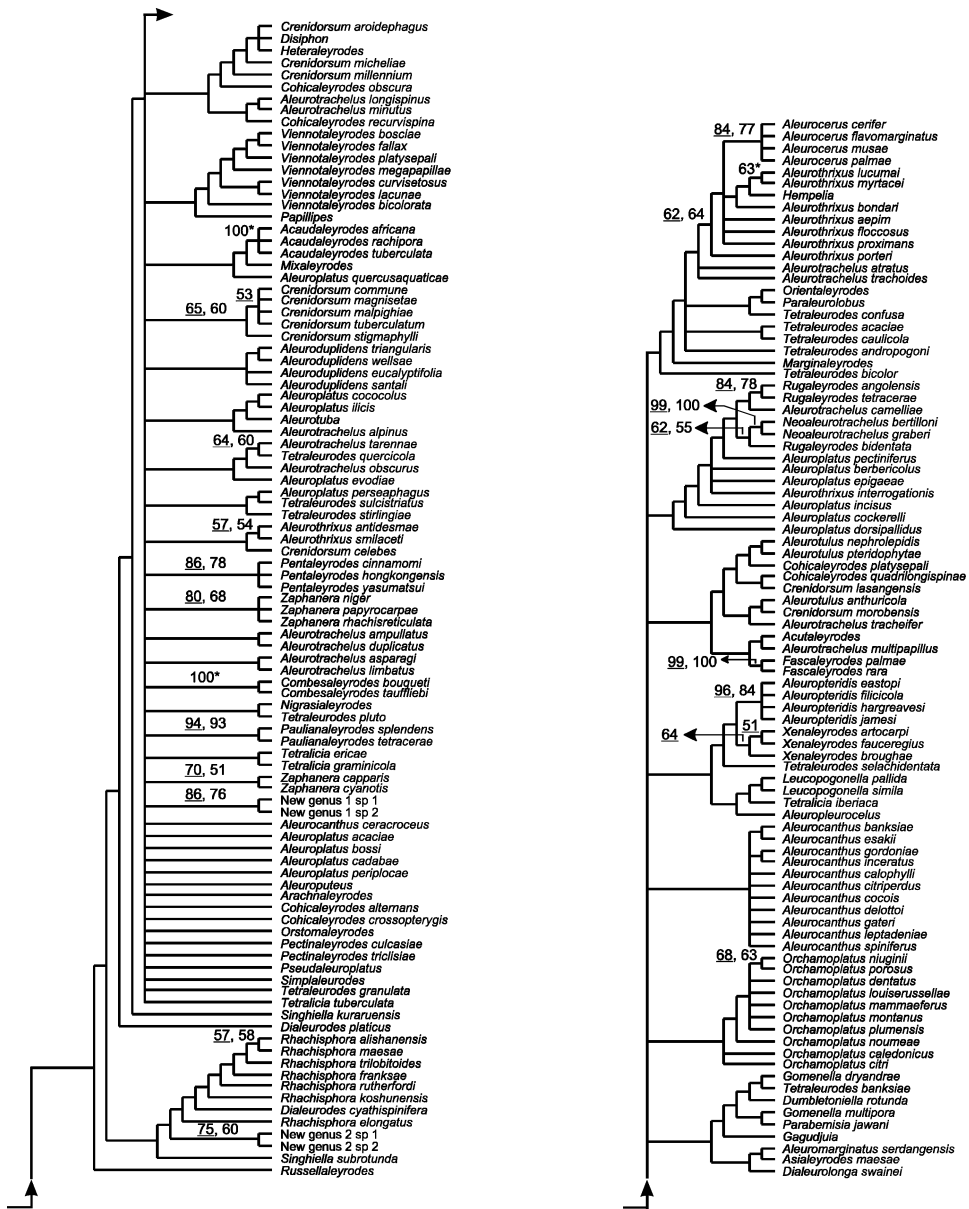


Figure 13. (Continued).

Minutaleyrodes, *Neoaleurotrachelus*, *Neomaskellia*, New genus 1, New genus 2, *Orchamoplatus*, *Paulianaleyrodes*, *Pentaleyrodes*, *Pogonaleyrodes*, *Rusostigma*, *Siphoninus*, *Singhius*, *Viennotaleyrodes*, *Xenaleyrodes*. This means that 40 genera were not recovered to be monophyletic. However, some of these were rendered non-monophyletic by a single species or very small groups of aberrant ones, viz. *Aleurocanthus* excluding *A. ceracroceus* Martin, *Aleyrodes* excluding *A. elevatus* Silvestri, *Lipaleyrodes* excluding *L. atriplex* (Froggatt), *Massilieurodes* excluding *M. fici* (Takahashi), *Rhachisphora* excluding *R. elongatus* Regu & David, *Rugaleyrodes* excluding *R. bidentata* Bink-Moenen, *Setaleyrodes*

excluding *S. vigintiseta* Martin, and most of the species of the genus *Dialeurologa* including the type-species as well as those species of *Crenidorsum* described by Russell (1945), including the type-species formed monophyletic clades. None of the mentioned excluded species are the type-species of their genera.

Successive approximation weighting

Applying SAW gave more resolution for unresolved nodes of the tree. More than 10% of the 30 000 MPTs obtained from the unweighted analysis (UW-MPTs) were selected so that to have at least 100 representatives from each 1000 MPTs. Then, the approach was used with four different indices, i.e. maximum RI, maximum CI, minimum RI, and minimum CI to weight characters, and four, eight, four, and eight cycles of reweighting were needed, respectively, before stability was reached. When all characters were set back to unit weight, the lengths of the trees were 2733, 2953, 2734 and 2927, respectively. Comparing these lengths with the length of UW-MPTs, i.e. 2730, indicates that the trees obtained from applying SAW using maximum and minimum RI (MPTs-SAW-MaxRI and MPTs-SAW-MinRI) are only three and four steps longer, respectively, and here the strict consensus tree of these trees (MPTs-SAW-MaxRI) is presented (Figure 14) and discussed. It is worth mentioning that the topology of the strict consensus trees of MPTs-SAW-MaxRI and MPTs-SAW-MinRI was almost the same as each other, and both were more in agreement with the strict consensus tree of UW-MPTs (Table I). The strict consensus trees obtained from applying SAW using maximum and minimum CI (MPTs-SAW-MaxCI and MPTs-SAW-MinCI) showed two different topologies and were somewhat less in agreement with the strict consensus tree of UW-MPTs (Table I).

In the strict consensus tree of MPTs-SAW-MaxRI (Figure 14), of the 33 genera recovered as monophyletic in the initial analyses, two, viz. *Dialeuropora* and *Pogonaleyrodes*, were no longer recovered, but the former excluding *D. silvarum* (Corbett) (not the type-species) did form a monophyletic clade. However, *Pectinaleyrodes* was monophyletic in this tree, whereas it was not in the initial tree. The monophyly of other mentioned genera were the same.

Monophyly of large genera

The evidence against monophyly of two economically important genera, *Bemisia* and *Trialeurodes*, as well as several large genera, i.e. *Aleuoplatus*, *Aleurotrachelus*, *Dialeurodes*, and *Tetraleurodes*, which were not recovered to be monophyletic in either unweighted or SAW analyses was investigated. Separate analyses were performed constraining each of these to be monophyletic, and the resulted trees compared statistically with the initial unconstrained ones using the non-parametric ranked-sign test of Templeton.

Constraining *Bemisia* to be monophyletic resulted in one MPT (length=2772) and using the new tree searching strategy yielded more than 1000 MPTs (length=2756). The Templeton test showed that there were significantly longer than the best unconstrained ones ($P=0.0196-0.0419$) (Table II). Constraining *Dialeurodes* to be monophyletic yielded trees of length 2771 which were not different from unconstrained ones based on the Templeton test ($P>0.07$).

In all constraint analyses, only one MPT found after performing a heuristic search with 1000 random additions, which were significantly different from the UW-MPTs except for *Bemisia* and *Trialeurodes*. The final results changed for some genera after finding shortest

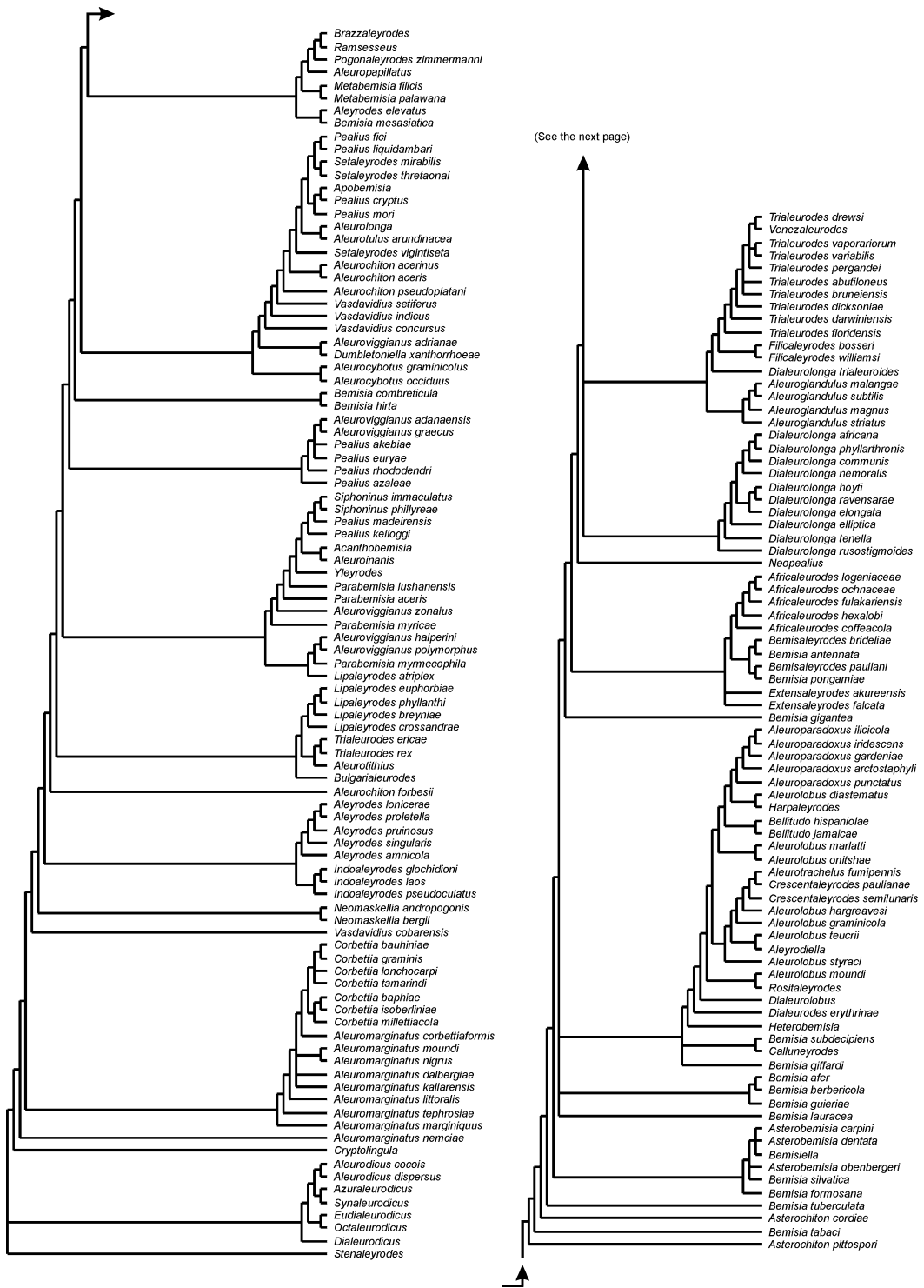


Figure 14. Strict consensus tree of more than 10 000 MPTs-SAW-MaxRI.

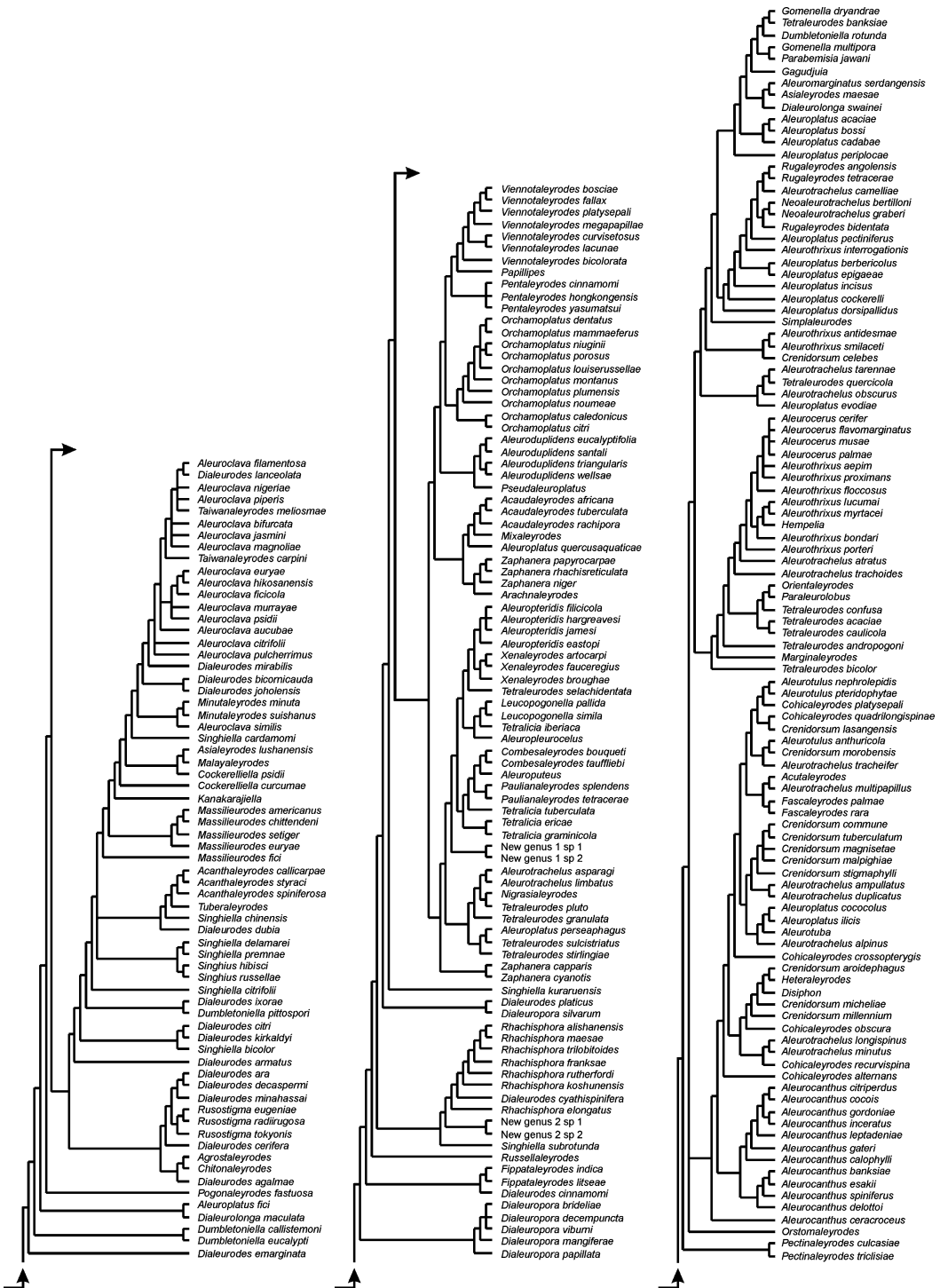


Figure 14. (Continued).

Table I. Symmetric-difference distances between different trees obtained from unweighted analysis and by applying successive approximation weighting.

Between trees	Symmetric-difference distances
MPTs-SAW-MaxRI-MPTs-SAW-MinRI	42–50
MPTs-SAW-MaxRI-UW-MPTs	73–98
MPTs-SAW-MinRI-UW-MPTs	100–124
MPTs-SAW-MaxCI-MPTs-SAW-MinCI	437–444
MPTs-SAW-MaxCI-UW-MPTs	611–620
MPTs-SAW-MinCI-UW-MPTs	614–624

possible trees applying the new strategy (at least 500 trees were compared), which are shown in Table II. Generally, the monophyly of *Aleurotrachelus* and *Bemisia* was statistically rejected, while that of *Aleuroplatus*, *Dialeurodes*, *Tetraleurodes*, and *Trialeurodes* was not (Table II). These results will be discussed in detail later.

The finding that statistical significance can increase when difference in tree length decreases (as a result of finding most parsimonious trees) indicates that it is important to reduce the variance, i.e. noise, in the estimate of numbers of character steps, and the test is only meaningful if the shortest possible trees to be compared. If they are not the MPTs, there will be a degree of “randomness” in the character lengths and such random variation might occasionally be on the side of increasing significance, sometimes not.

Evaluating the current tribe classification

As already mentioned, 13 tribes have been proposed so far within the Aleyrodinae. In addition to the problems posed by unplaced genera, there is little agreement about the composition of some proposed tribes (Sampson 1943, 1947; Russell 1947; Takahashi 1954; David 1990). Before starting to evaluate the current tribe classification, it should be mentioned that some of the genera in Appendix A have been synonymized and these will be indicated in the discussion of each tribe. Furthermore, except the tribe proposed by Russell (1947), Trialeurodini, the other tribes were only defined in identification keys by the authors, who erected them (see above). Tribal level characters used by these authors for each tribe are given in Appendix B to avoid repeating them when each tribe is discussed.

Table II. Results of the Templeton test obtained by comparing the shortest trees for those genera constrained to be monophyletic, with the UW-MPTs.

Constrained genus	Constrained tree length	Tree length of UW-MPTs	Templeton test <i>P</i> values
<i>Aleuroplatus</i>	2757	2730	0.0513–0.2019
<i>Aleurotrachelus</i>	2780	2730	0.0085*–0.0186*
<i>Bemisia</i>	2756	2730	0.0196*–0.0419*
<i>Dialeurodes</i>	2771	2730	0.0785–0.1120
<i>Tetraleurodes</i>	2764	2730	0.0469*–0.0948
<i>Trialeurodes</i>	2741	2730	0.5590–0.7035

*Significant difference at $P < 0.05$.

Tribe Aleurocanthini

Three genera, viz. *Aleurocanthus*, *Aleurotrachelus*, and *Pentaleyrodes*, were placed in the Aleurocanthini by Takahashi (1954), and only *Aleurocanthus* by David (1990) for the whiteflies of Japan and India, respectively (Appendix A). As mentioned before, the latter excluding *A. ceracroceus* was found to be monophyletic and the monophyly of *Aleurotrachelus* even statistically rejected (Table II). However, the type-species of *Aleurotrachelus*, *A. tracheifer* (Quaintance), was never recovered in the same clade either with *Aleurocanthus* or *Pentaleyrodes* in the strict consensus tree of UW-MPTs (Figure 13), whereas in the strict consensus tree of MPTs-SAW-MaxRI (Figure 14) it was found in a different clade from the *Aleurocanthus* clade within a big clade.

Tribe Aleurochitonini

Sampson (1943) proposed this tribe on the basis of a single genus, *Aleurochiton* (Appendix A). Two of the four studied species of this genus, *A. acerinus* Haupt and *A. aceris* (Modeer) (the type-species), were recovered as forming a monophyletic group in this study (Figures 13 and 14). Although in the strict consensus trees of MPTs-SAW-MaxCI and MPTs-SAW-MinCI (trees not presented), *A. pseudoplatani* Visnya also came out with the former two species, *A. forbesii* (Ashmead) was not recovered with them in the same clade in any analysis. Interestingly, Sampson (1943) erected a new subgenus, *Nealeurochiton* Sampson, to accommodate *A. forbesii*, but this was subsequently synonymized with *Aleurochiton* by Mound and Halsey (1978).

Tribe Aleurilobini

This tribe was proposed by Takahashi (1954), for five genera based on Japanese species (*Acanthobemisia*, *Aleurolobus*, *Apobemisia*, *Bemisia*, *Parabemisia*). All studied species of *Aleurolobus* including the type-species, *A. marlatti* (Quaintance), were recovered in the same clade but not as a monophyletic group. The only examined species of *Apobemisia*, *A. kuwanai* (Takahashi) (the type-species), was not found with *Aleurolobus* in the same clade, and the type-species of *Bemisia* (not recovered as a monophyletic genus), *B. tabaci*, was never recovered either with *Aleurolobus* or *Apobemisia*. The type-species of *Parabemisia* was not studied but the five examined species did not form a clade. *Acanthobemisia* and the four species of *Parabemisia* were only recovered in the same clade in the strict consensus tree of MPTs-SAW-MaxRI (Figure 14). David (1990) later transferred *Bemisia* to a separate tribe, Bemisini (see below), but Takahashi's Aleurolobini, even excluding *Bemisia*, was not supported in this study.

David (1990) and Regu and David (1993) classified six genera from India in this tribe (Appendix A). Only the type-species of *Aleuropapillatus* and *Orientalleyrodes* were studied and these appeared in two distantly separated clades. *Africaleurodes* was found as a monophyletic genus, but did not form a clade with the other five genera. Two examined species of *Asterochiton* (but not the type-species) were recovered on two different branches. All species of *Aleurolobus* and *Crescentaleyrodes* were found in the same clade (Figures 13, 14 and 19), and the possible monophyly of the latter will be discussed later.

Tribe Aleuroplatini

David (1990) placed two genera in this tribe (Appendix A), of which *Moundiella* David was later synonymized with *Viennotaleyrodes* (David et al. 1994). Unlike the latter genus,

Aleuroplatus was found to be non-monophyletic, though its monophyly was not rejected statistically using the Templeton test (Table II) (see also “Discussion”). The type-species of *Aleuroplatus*, *A. quercusaquaticae* (Quaintance), and *Viennotaleyrodes* gathered in two different clades within a relatively big monophyletic clade in the strict consensus tree of MPTs-SAW-MaxRI (Figure 14).

Tribe Aleyrodini

The Aleyrodini originally comprised 24 genera (Sampson 1943), and the author added another genus in 1947. Five more genera were added later (Drews and Sampson 1956; Sampson and Drews 1956) (Appendix A). Of these *Aleuromigada* Singh is considered as *nomina nuda* and the name *Frauenfeldtiella* Gomez-Menor is not available in Aleyrodidae as it is preoccupied in the *Cecidomyiidae*; the genus is now known as *Aleurotuba* (Mound and Halsay 1978; Tremblay and Iaccarino 1978). There were no specimens available for *Hesperaleyrodes*, *Luederwaldtiana*, *Mexicaleyrodes*, *Nealeyrodes*, and *Neoaleyrodes*, and *Laingiella* was also omitted from the analysis (see “Materials and methods”). Furthermore, *Aleurocanthus*, *Aleurotrachelus*, and *Pentaleyrodes* were transferred to Aleurochantini (see above), but David (1990) later reassigned *Aleurotrachelus* to Aleyrodini. Russell (1947) transferred *Aleurotithius* to Trialeurodini, and David (1990) relocated *Acaudaleyrodes*, *Tetraleurodes*, and *Zaphanera* in Neomaskellini, Tetraleurodini, and Zaphanerini, respectively (see below). Monophyly of Aleyrodini based on Sampson’s classification even excluding *Acaudaleyrodes*, *Aleurocanthus*, *Aleurotithius*, *Aleurotrachelus*, *Pentaleyrodes*, *Tetraleurodes*, and *Zaphanera* was not supported in this study.

Takahashi (1954) placed only five genera from Japan in the Aleyrodini (Appendix A), of which two genera are now considered as synonyms: *Odontaleyrodes* Takahashi was synonymized with *Pealius* by Martin (1999). This tribal classification, too, failed to be monophyletic. Except *Aleyrodes*, David (1999) transferred all genera placed by Takahashi in Aleyrodini, to Bemisini and placed two more genera in the latter tribe (see Bemisini below).

David’s (1999) classification of Aleyrodini with five genera from India, viz. *Aleurocybotus*, *Aleuromarginatus*, *Aleurotrachelus*, *Aleurotulus* and *Aleyrodes*, was not supported by this study.

Tribe Bemisini

Four of the six genera placed in this tribe by David (1990) (Appendix A) were briefly discussed above. The type-species of *Pealius* was not studied but the 10 examined species did not form a monophyletic clade. *Setaleyrodes* including the type-species, *S. mirabilis* Takahashi, also appeared non-monophyletic, but they were recovered together with some species of *Pealius* in the same clade (Figures 13 and 14). *Indoaleyrodes* (three species examined including the type-species, *I. laos* (Takahashi)) was recovered as monophyletic but in a separate clade. The type-species of *Neopealius* (the only examined species of this small genus) was on its own branch. As already mentioned, the monophyly of *Bemisia* was refuted statistically (Table II) (see also “Discussion”) and its type-species, *B. tabaci*, was on its own branch.

Tribe Dialeurodini

Sampson (1943, 1947) placed 32 genera in the tribe Dialeurodini and 13 more genera were later added by Sampson and Drews in 1956 (Appendix A). Of these, *Aleuroclava* and

Aleurotuberculatus Takahashi are now treated under the former name (Martin 1999), as well as *Corbettella* Sompson, *Neobemisia* Visnya and *Roucasia* Goux, which were synonymized with *Pealius*, *Asterobemisia* (Mound and Halsey 1978) and *Bemisia* (Danzig 1964), respectively. Also, *Nipaleyrodes* and *Stenaleyrodes* have since been classified as members of the Aleurodicinae (Mound and Halsey 1978) (see also "Discussion" for *Stenaleyrodes*). For *Aleuroporosus*, *Anomaleyrodes*, *Dialeurotrachelus*, *Neoleurolobus*, *Metaleyrodes*, *Plataleyrodes*, *Pseudaleurolobus*, *Pseudaleyrodes*, *Trichoaleyrodes* and *Xenobemisia* there were no specimens available for study. Although more than half of the 29 remaining genera were not found to be monophyletic, considering only their type-species, they never form a monophyletic clade, and the picture is not improved by excluding those genera that were later transferred to other tribes, viz. *Acanthobemisia*, *Africaleurodes*, *Aleurolobus*, *Aleuroparadoxus*, *Aleuroplatus*, *Asterochiton*, *Bemisia*, *Parabemisia*, *Pealius*, *Setaleyrodes*, and *Trialeurodes* (Russell 1947; Takahashi 1954; David 1990; Regu and David 1993).

Takahashi (1954) placed only four genera from Japan in this tribe (Appendix A), of which *Rhachisphora* was recovered in a separate clade from the other three genera in all analyses (Figures 13 and 14) but not as a monophyletic genus. All examined species of *Taiwanaleyrodes* and *Aleuroclava* (= *Aleurotuberculatus*) (see above), and also a few species of *Dialeurodes* including the type-species, *D. citri* (Ashmead) were in the same clade together with some other genera in both strict consensus trees (Figures 13, 14 and 17). *Aleuroclava* and *Taiwanaleyrodes* will be discussed in detail later.

David (1990) and David and Sundararaj (1993) classified 19 genera from India in the Dialeurodini (Appendix A), of which three genera have now been synonymized (*Martiniella* Jesudasan & David and *Aleurotuberculatus* were synonymized with *Aleuroclava* by Martin (1999)). Also, the three subgenera of *Dialeurodes*, viz. *Dialeuronomada*, *Gigaleurodes*, and *Rabdostigma* of Quaintance & Baker, which were raised to the generic level by Indian authors (David and Sundararaj 1993; Sundararaj and David 1994), were here treated under *Dialeurodes* (the four examined species of the latter genus, viz. *D. cerifera* Quaintance & Baker, *D. cinnamomi* Takahashi, *D. ixorae* Singh and *D. minahassai* Martin are assignable to *Gigaleurodes*, *Gigaleurodes*, *Dialeuronomada* and *Rabdostigma*, respectively, but none of them is the type-species). *Kanakarajiella* (only the type-species examined), *Minutaleyrodes*, and *Singhius* gathered in the same clade with *Aleuroclava*, *Dialeurodes*, and *Taiwanaleyrodes* (Figure 17). The type-species of *Asialeurodes* and *Cockerelliella* were not studied but one of the two examined species of the former and all examined species of the latter genus as well as six of the eight examined species of *Singhiella* including the type-species, *S. bicolor* (Singh), were also recovered in this clade, while *Rusostigma* was only found in this clade in the MPTs-SAW-MaxRI (Figure 14). Furthermore, except *D. cinnamomi*, the other three species of *Dialeurodes*, all were found in this clade but only in the MPTs-SAW-MaxRI (*D. ixorae* was also found in the UW-MPTs), i.e. recovering the representatives of all three subgenera. The four remaining genera, *Dialeurolobus*, *Dialeuropora*, *Fippataleyrodes*, and *Rhachisphora* also appeared in four different clades.

Tribe Lipaleyrodini

David (1990) included only *Lipaleyrodes* for this tribe (Appendix A) and the genus excluding *L. atriplex* was recovered to be monophyletic (including the type-species, *L. phyllanthi* Takahashi). *L. atriplex* had originally been described in *Aleyrodes*, and was subsequently transferred to *Lipaleyrodes* by Martin (1999), though it did not group even with the species of *Aleyrodes* in this study.

Tribe Neomaskellini

Sampson (1943) based the monotypic Neomaskellini on *Neomaskellia*, and later *Acaudaleyrodes* was added by David (1990) (Appendix A). Both genera were recovered to be monophyletic but located in two distantly separated clades in all analyses (Figures 13 and 14).

Tribe Siphoninini

This tribe includes only the monophyletic genus *Siphoninus* (Sampson 1943; David 1990).

Tribe Tetraleurodini

The only included genus, *Tetraleurodes*, was not found to be monophyletic in any analyses, though its monophyly was not rejected statistically (Table II) (see also “Discussion”).

Tribe Trialeurodini

Russell (1947) erected this tribe and stated “only the known genera *Aleuroparadoxus* and *Aleurotithius* of Quaintance and Baker and *Trialeurodes* Cockerell appear to be assignable to this tribe”. Subsequently, Russell (1967) added her new monobasic genus, *Venezaleurodes*. In the strict consensus tree of UW-MPTs (Figure 13), nine of the 11 examined species of *Trialeurodes* including the type-species, *T. pergandei* (Quaintance), as well as *Aleurotithius* (only the type-species studied), *Venezaleurodes* and several different genera formed a polytomy, and only two species of *Trialeurodes* were recovered in the same clade. *Aleuroparadoxus* was found to be monophyletic and formed a separate clade with several different genera. In the MPTs-SAW-MaxRI (Figure 14), the nine species of *Trialeurodes*, and *Venezaleurodes* were recovered in the same clade, and *Aleurotithius* formed a separate small clade with the two other species of *Trialeurodes*.

Tribe Zaphanerini

This tribe was proposed by David (1990) for *Zaphanera*, (Appendix A). Five species of this genus including the type-species, *Z. cyanotis* Corbett, were included in the present study. These formed two separate monophyletic clades which were supported by relatively high Jackknife values (Figure 13). The African and oriental species (*Z. capparisi* Bink-Moenen and *Z. cyanotis*, respectively) were recovered in one, and the three Australian species in another (Figures 13 and 14). Thus, the Australian species (*Z. niger* (Maskell), *Z. papyrocarpae* Martin, and *Z. rhachisreticulata* Martin) do not appear to be congeneric with the type-species and a new genus needs to be erected for them (see “Nomenclatural changes”).

Whitefly–host plant relationships

Host plant information at family, order and higher group levels for all whitefly species included in this study is given in Appendix E. All botanical names follow the system of the Angiosperm Phylogeny Group (APG) (Bremer et al. 1998, 2003; Soltis et al. 2000). Whitefly host plant information was obtained from Mound and Halsey (1978), Bink-Moenen (1983) and Martin (1999), as well as from more recent original descriptions, and from unpublished information available on slides examined.

Except for the only gymnosperm host record, *Dioon spinulosum* (Zamiaceae) for *T. vaporariorum* (Westwood), and also those few species recorded from Pteridophyta (see

below), whiteflies are found mainly on angiosperms. Furthermore, few whitefly species are known to be monophagous, most being oligo- or polypahgous. The number of host plant orders attacked by the species studied here varies from one to 31 (probably the true number is higher) (Appendix E).

Twenty described, and three or four undescribed, species are reported from fern hosts (Pteridophyta) (Mound and Halsey 1978; Martin and Camus 2001), of which the fern host record for *T. vaporariorum* (which is highly polyphagous on Angiospermae) is uncertain, and the identification of one of these species as *B. tabaci* (also highly polyphagous on Angiospermae) is not definitive (Martin and Camus 2001). Here, 14 described species recorded from ferns (ignoring *B. tabaci* and *T. vaporariorum*) were scored. Of these, all described species of *Aleuopteridis* and *Filicaleyrodes*, were recovered as monophyletic, but they were distantly separated in trees from both analyses (Figures 13 and 14). The results were the same for the two fern-feeding species of *Metabemisia* (*M. filicis* Mound and *M. palawana* Martin), as well as the two fern-feeding species of *Aleurotulus* (*A. nephrolepidis* (Quaintance) (the type-species) and *A. pteridophytiae* Martin). Neither the type-species of *Metabemisia* (not examined) nor the other species of *Aleurotulus* have been recorded from ferns. The latter genus appears not to be monophyletic and, apparently, its non-fern-feeding species are not congeneric with the type-species (it should be noted that one of its non-fern-feeding species, *A. arundinacea* Singh, is considered as species *incertae sedis* (Mound et al. 1994)). For *Trialeurodes*, three of the 11 examined species, *T. bruneiensis* Martin, *T. dicksoniae* Martin, and *T. rex* Martin, colonise ferns, but they were not recovered in the same clade. The only studied species of *Mixaleyrodes* were not recovered as a sister group to any of the fern-feeding species. In general, the position of these 14 species on the both cladograms did not support a clear Pteridophyta–whitefly relationship for the all fern-feeding species, although separate correlations were found for some species at genus level.

Almost the same results were obtained after mapping angiosperm hosts on both strict consensus trees. In general, some small monophyletic clades appear to be specialized on particular host plant orders, but in most cases, there was little of any indication of correlation between the species of a particular clade and host plant order.

Three genera for which a clear relationship with host plant order was apparent were *Aleuromarginatus*, *Corbettia*, and *Viennotaleyrodes*. The last two appeared to be monophyletic, but *Aleuromarginatus* was not found to be so, though most of its examined species formed a monophyletic clade with *Corbettia* (Figure 15). Virtually all species of these three genera have been recorded from Fabaceae (the exception being *C. graminis* Mound on Poaceae, *V. bosciae* Bink-Moenen recorded from Brassicaceae, and *C. milletticola* Dozier which has additionally been recorded from Apocynaceae) (there is no host record for *A. serdangensis* Takahashi but this species most probably does not belong to *Aleuromarginatus*; see “Discussion”) (Appendix E). Furthermore, the monobasic genus *Papillipes*, which is also recorded from Fabaceae, formed a sister group to *Viennotaleyrodes* (Figures 13 and 14), but this clade did not form a sister group to *Aleuromarginatus*+*Corbettia*.

Discussion

The family Aleyrodidae is taxonomically difficult. The genera usually have no reliable suite of morphological characters to allow them to be recognised unambiguously, and this is obvious from many of the original descriptions. Many of the larger genera have some species included which differ in at least one of the characters given in the description of the genus (e.g. *Aleurotrachelus asparagi* (Lewis) and *A. tarennae* Bink-Moenen have a

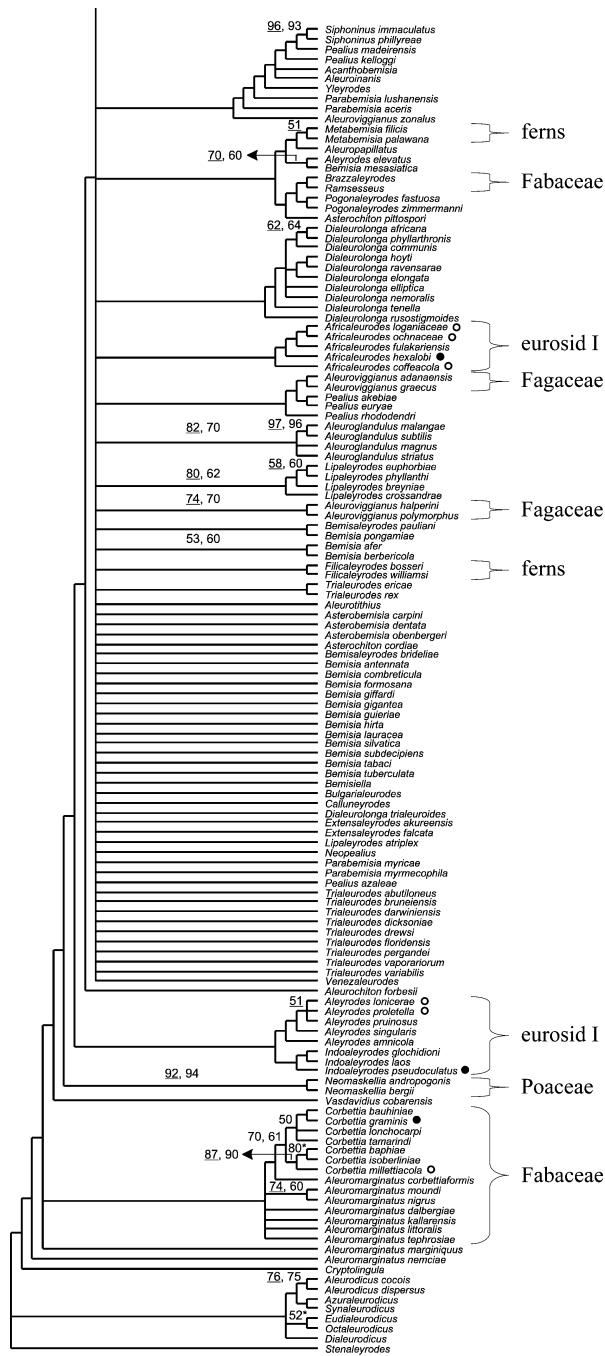


Figure 15. Correlation of the species in the basal part of the strict consensus tree of UW-MPTs with host plants. Figures above branches show bootstrap and Jackknife (underlined figures) values. Figures with an asterisk indicate the same value for both bootstrap and Jackknife. A black circle indicates that the species was not recorded from the same host family/higher group in the relevant clade. An empty circle indicates additional host family/higher group record for the species in the relevant clade.

submarginal line or fold, whereas, based on the original description of the genus, the submarginal area is not separated from the dorsal disc (Quaintance and Baker 1914) and, sometimes, there is really no reason for a species to have been placed in its current genus at all (e.g. *Aleuromarginatus serdangensis*; see below).

These and other problems created some difficulties in choosing possible genus level characters for the cladistic analyses. We attempted to code all puparial characters used by authors in generic descriptions, although in most cases there are no distinct boundaries between genus level characters and those of species level. For example, Bink-Moenen (1983) while describing three new genera, *Arachnaleyrodes*, *Papillipes* and *Yleyrodes*, and comparing them with their morphologically closely related genera, i.e. *Orchamoplatus*, *Viennotaleyrodes* and *Africaleurodes* respectively, used the absence and/or presence of first abdominal setae as one of the diagnostic characters to separate the new genera from their allied genera. This character, however, has also been used to differentiate two species of a particular genus, e.g. *Dialeurolobus rhamni* Bink-Moenen and *D. pulcher* Danzig (Bink-Moenen and Gerling 1990), and *Crenidorsum millennium* Martin and *C. celebes* Martin (Martin 1999). As stated by Martin (2003), there is nothing inherently wrong with the use of immature stages taxonomically, but the problem is that, with our current poor understanding of the true significance of many puparial characters, we have yet to maximise their value. The lack of such information has impeded our understanding of relationships between aleyrodids, rather than any lack of characters for use (immature stages have been used taxonomically in other insects such as Psylloidea (White and Hodkinson 1982) and Lepidoptera (Kitching 2002, 2003; Willmott 2003)). For example, as mentioned in comments for character 80 (glandular bases to marginal teeth), this structure has sometimes been described as “a double row of teeth” or rarely “a translucent membranous area at the bases of the marginal teeth”, and according to Bink-Moenen (1983), these are probably wax-secreting pores or papillae. In the latter case, characters 65 (submarginal papillae) and 80, apart from their shape differences, can be considered the same in a broad sense. Table III shows some examples regarding the opinions of different authors about the relationships between some genera. In fact, with whitefly puparial taxonomy lacking objective criteria (it seems to be partly a matter of guess-work at genus level), there is likely to be quite a lot of apparent homoplasy (in addition to some true homoplasy) due to misinterpreting similar character states during the coding stage.

Owing to the high level of homoplasy in the data set, the maximum consistency index may be the best choice for reweighting characters for SAW, because it cannot weight characters as zero, so potentially useful characters would not be excluded from the analysis (Quicke et al. 2001). It should be noted that when stability was reached in SAW analyses and all characters then set back to unit weight, the lengths of the MPTs-SAW-MaxCI and MPTs-SAW-MinCI were much higher (223 and 197 steps, respectively) than that of UW-MPTs, while MPTs-SAW-MaxRI and MPTs-SAW-MinRI were only a few steps longer (see “Results”).

Retention indices (ri) for each non-constant character were calculated to evaluate how much of the variation displayed by the characters could be attributed to true synapomorphy on trees, as well as calculating consistency indices (ci) to evaluate the amount of homoplasy of each character, and length. The results are shown in Table IV. In all UW-MPTs, the retention indices of characters 5, 14, and 86 were zero, i.e. contributing no synapomorphy to the cladograms. In contrast, retention index had the maximum value for the characters 22, 24, 30, 46 (only the best fit), 63, 64, 76, 78/79 (only the best fit), 82–84, 87/88 and 90, indicating the contribution of maximum synapomorphy to the cladograms (Table IV). Of

Table III. Relationships between whitefly genera proposed by various authors.

Genus	Allied genera	Reference
<i>Aleuroduplidens</i>	<i>Aleuroplatus</i> , <i>Zaphanera</i> , <i>Tetraleurodes</i>	Martin (1999)
<i>Aleuroglandulus</i>	<i>Aleyrodes</i>	Russell (1944)
<i>Aleuropapillatus</i>	<i>Aleurolobus</i> , <i>Crescentaleyrodes</i>	Regu and David (1993)
<i>Aleuroparadoxus</i>	<i>Trialeurodes</i>	Russell (1947)
<i>Aleyrodiella</i>	<i>Aleuroparadoxus</i> , <i>Aleurotithius</i> , <i>Trialeurodes</i>	Danzig (1966)
<i>Asialeyrodes</i>	<i>Africaleurodes</i> , <i>Aleurolobus</i> , <i>Dialeurodes</i>	Corbett (1935)
<i>Belliudo</i>	<i>Aleuroparadoxus</i> , <i>Pseudaleurolobus</i> , <i>Africaleurodes</i> , <i>Aleurolobus</i> , <i>Paraleurolobus</i>	Russell (1943)
<i>Brazzaleyrodes</i>	<i>Aleurolonga</i> , <i>Ramsesseus</i>	Bink-Moenen (1983)
<i>Bulgarialeurodes</i>	<i>Corbettia</i>	Corbett (1936)
<i>Chitonaleyrodes</i>	<i>Zaphanera</i> , <i>Gagudjuia</i>	Martin (1999)
<i>Cockerelliella</i>	<i>Asialeyrodes</i>	Sundararaj and David (1992)
<i>Cockerelliella</i>	<i>Asialeyrodes</i> , <i>Dialeurodes</i>	Martin (1999)
<i>Crenidorsum</i>	<i>Aleuroplatus</i> , <i>Aleurotrachelus</i> , <i>Aleurotulus</i> , <i>Aleuroputeus</i> , <i>Laingiella</i>	Russell (1945)
<i>Crenidorsum</i>	<i>Aleurotrachelus</i> , <i>Aleurotulus</i>	Martin et al. (2001)
<i>Dumbletoniella</i>	<i>Aleuroputeus</i> , <i>Aleuroclava</i>	Jesudasan and David (1990)
<i>Dumbletoniella</i>	<i>Aleuroclava</i> , <i>Aleuroplatus</i> , <i>Pseudaleuroplatus</i>	Martin (1999)
<i>Extensaleyrodes</i>	<i>Dialeurolonga</i> , <i>Aleuromarginatus</i>	Bink-Moenen (1983)
<i>Fascaleyrodes</i>	<i>Aleurocanthus</i>	Bink-Moenen (1983)
<i>Filicaleyrodes</i>	<i>Trialeurodes</i>	Takahashi (1962); Mound (1966)
<i>Gomenella</i>	<i>Indoaleyrodes</i> , <i>Asialeyrodes</i>	Martin (1999)
<i>Heterobemisia</i>	<i>Bemisia</i>	Takahashi (1957)
<i>Leucopogonella</i>	<i>Aleurotrachelus</i>	Dumbleton (1961)
<i>Malayaleyrodes</i>	<i>Africaleurodes</i> , <i>Asialeyrodes</i> , <i>Aleurolobus</i>	Corbett (1935)
<i>Marginalaleyrodes</i>	<i>Aleurotrachelus</i>	Takahashi (1961)
<i>Metabemisia</i>	<i>Parabemisia</i> , <i>Apobemisia</i>	Takahashi (1963)
<i>Mixaleyrodes</i>	<i>Aleurotrachelus</i>	Takahashi (1936a); Jesudasan and David (1991)
<i>Neopealius</i>	<i>Pealius</i> , <i>Bemisia</i>	Takahashi (1954)
<i>Nigrasialeyrodes</i>	<i>Asialeyrodes</i> , <i>Gomenella</i>	Martin (1999)
<i>Papillipes</i>	<i>Viennotaleyrodes</i>	Bink-Moenen (1983)
<i>Pseudaleuroplatus</i>	<i>Aleuroduplidens</i> , <i>Aleuroplatus</i> , <i>Trialeurodes</i> , <i>Zaphanera</i>	Martin (1999)
<i>Rositalyrodes</i>	<i>Aleurolobus</i>	Meganathan and David (1994)
<i>Rugaleyrodes</i>	<i>Neoaleurotrachelus</i> , <i>Cohicaleyrodes</i>	Bink-Moenen (1983)
<i>Setaleyrodes</i>	<i>Pealius</i>	Martin (1999)
<i>Taiwanaleyrodes</i>	<i>Aleurotuberculatus</i> , <i>Aleurocybotus</i>	Takahashi (1932)
<i>Tetralicia</i>	<i>Aleuropleurocelus</i>	Drews and Sampson (1956); Mound (1966)
<i>Tetralicia</i>	<i>Aleuropleurocelus</i> , <i>Tetraleurodes</i>	Martin (1999)
<i>Trialeurodes</i>	<i>Aleuroparadoxus</i> , <i>Aleurotithius</i>	Russell (1948)
<i>Trialeurodes</i>	<i>Filicaleyrodes</i>	Mound (1966)
<i>Trialeurodes</i>	<i>Dialeurolonga</i>	Bink-Moenen (1983)
<i>Venezaleurodes</i>	<i>Trialeurodes</i>	Russell (1967)
<i>Xenaleyrodes</i>	<i>Aleuroplatus</i>	Takahashi (1936b)
<i>Xenaleyrodes</i>	<i>Aleurocanthus</i>	Martin (1999)
<i>Yleyrodes</i>	<i>Africaleurodes</i>	Bink-Moenen (1983)
<i>Zaphanera</i>	<i>Aleurotrachelus</i>	Corbett (1926)
<i>Zaphanera</i>	<i>Dialeurolobus</i> , <i>Tetraleurodes</i>	Martin (1999)

Table IV. Range of retention indices, consistency indices and lengths for each non-constant character calculated for the UW-MPTs. B=Best; Ch=Character; Ma=Maximum; Mi=Minimum; W=Worst. Characters 54, 68 and 81 are parsimony uninformative.

Ch	Retention index		Consistency index		Length		Ch	Retention index		Consistency index		Length	
	B	W	B	W	Mi	Ma		B	W	B	W	Mi	Ma
[1]	0.73	0.71	0.07	0.07	54	58	[48]	0.40	0.35	0.73	0.72	45	46
[2]	0.58	0.57	0.10	0.10	58	60	[49]	0.84	0.83	0.08	0.08	36	37
[3]	0.54	0.53	0.15	0.15	33	34	[50]	0.80	0.80	0.50	0.50	2	2
[4]	0.64	0.64	0.09	0.09	11	11	[51]	0.56	0.55	0.09	0.08	106	109
[5]	0.00	0.00	0.50	0.50	2	2	[52]	0.61	0.59	0.05	0.05	64	67
[6]	0.61	0.57	0.05	0.04	22	24	[53]	0.52	0.50	0.03	0.03	67	70
[7]	0.80	0.80	0.17	0.17	12	12	[54]	0/0	0/0	1.00	1.00	1	1
[8]	0.69	0.68	0.07	0.07	29	30	[55]	0.90	0.89	0.05	0.04	21	23
[9]	0.73	0.71	0.03	0.03	32	34	[56]	0.64	0.64	0.12	0.12	42	43
[10]	0.79	0.77	0.12	0.11	43	46	[57]	0.90	0.89	0.04	0.04	23	25
[11]	0.86	0.86	0.17	0.17	6	6	[58]	0.70	0.70	0.08	0.07	40	41
[12]	0.33	0.33	0.33	0.33	3	3	[59]	0.68	0.66	0.04	0.03	56	59
[13]	0.62	0.60	0.06	0.06	84	88	[60]	0.43	0.40	0.25	0.23	57	60
[14]	0.00	0.00	0.20	0.20	5	5	[61]	0.69	0.67	0.02	0.02	48	51
[15]	0.42	0.40	0.04	0.04	50	52	[62]	0.45	0.42	0.05	0.04	22	23
[16]	0.63	0.62	0.27	0.26	111	114	[63]	1.00	1.00	1.00	1.00	1	1
[17]	0.68	0.65	0.03	0.03	58	62	[64]	1.00	1.00	1.00	1.00	1	1
[18]	0.60	0.57	0.42	0.41	52	54	[65]	0.61	0.58	0.22	0.21	18	19
[19]	0.70	0.68	0.07	0.07	54	57	[66]	0.54	0.54	0.20	0.20	20	20
[20]	0.58	0.53	0.52	0.49	50	53	[67]	0.90	0.90	0.50	0.50	2	2
[21]	0.50	0.38	0.20	0.17	5	6	[68]	0/0	0/0	1.00	1.00	1	1
[22]	1.00	1.00	1.00	1.00	2	2	[69]	0.73	0.69	0.04	0.03	28	32
[23]	0.72	0.72	0.17	0.17	6	6	[70]	0.71	0.71	0.03	0.03	40	40
[24]	1.00	1.00	1.00	1.00	4	4	[71]	0.62	0.60	0.03	0.02	39	41
[25]	0.59	0.57	0.03	0.03	34	36	[72]	0.64	0.60	0.05	0.05	20	22
[26]	0.61	0.59	0.10	0.10	88	92	[73]	0.75	0.75	0.75	0.75	4	4
[27]	0.62	0.59	0.08	0.07	40	43	[74]	0.63	0.60	0.27	0.25	15	16
[28]	0.71	0.69	0.04	0.04	23	24	[75]	0.48	0.45	0.11	0.11	82	85
[29]	0.69	0.69	0.27	0.27	15	15	[76]	1.00	1.00	1.00	1.00	1	1
[30]	1.00	1.00	1.00	1.00	1	1	[77]	0.48	0.44	0.08	0.07	39	42
[31]	0.78	0.76	0.08	0.08	12	13	[78]	1.00	0.50	1.00	0.50	1	2
[32]	0.35	0.30	0.07	0.07	14	15	[79]	1.00	0.80	1.00	0.50	1	2
[33]	0.44	0.33	0.09	0.08	11	13	[80]	0.64	0.63	0.03	0.03	38	40
[34]	0.68	0.68	0.09	0.09	79	80	[81]	0/0	0/0	1.00	1.00	1	1
[35]	0.66	0.63	0.14	0.13	37	40	[82]	1.00	1.00	1.00	1.00	1	1
[36]	0.69	0.69	0.33	0.33	6	6	[83]	1.00	1.00	1.00	1.00	1	1
[37]	0.88	0.88	0.04	0.04	27	27	[84]	1.00	1.00	1.00	1.00	1	1
[38]	0.75	0.74	0.04	0.04	49	52	[85]	0.67	0.67	0.17	0.17	6	6
[39]	0.65	0.64	0.07	0.07	30	31	[86]	0.00	0.00	0.20	0.20	5	5
[40]	0.59	0.57	0.03	0.03	68	72	[87]	1.00	1.00	1.00	1.00	1	1
[41]	0.77	0.75	0.09	0.09	53	56	[88]	1.00	1.00	1.00	1.00	1	1
[42]	0.67	0.65	0.10	0.09	94	97	[89]	0.90	0.90	0.33	0.33	3	3
[43]	0.86	0.86	0.50	0.50	2	2	[90]	1.00	1.00	1.00	1.00	1	1
[44]	0.79	0.79	0.25	0.25	71	72	[91]	-	-	-	-	-	-
[45]	0.57	0.43	0.14	0.11	7	9	[92]	0.41	0.33	0.06	0.05	17	19
[46]	1.00	0.50	1.00	0.67	2	3	[93]	0.53	0.50	0.56	0.54	36	37
[47]	0.75	0.75	0.25	0.25	48	48	[94]	0.73	0.71	0.22	0.21	18	19

Character 91 is a constant character; see the comments of this character.

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these 15 characters, characters 79 (subdorsal compound wax pores and/or agglomerate pores) and 90 (tarsal claw) are restricted to the Aleurodicinae, although *Dialeurodicus* lacks character 79 and *Stenaleyrodes* lacks both characters.

Excluding *Stenaleyrodes* from out-group taxa, puts it in the position of being a sister group to all ingroup taxa. Although Takahashi (1938) gave no information about the subfamily placement in his description of the monobasic genus *Stenaleyrodes* – as well as Mamet (1952) giving supplementary notes on the type-species, *S. vinsoni* Takahashi – but at the end of the description, the author stated “related to *Trialeurodes* Laing, but differs in the elongate vasiform orifice, Resembles *Bemisia* Quaint. et Baker in the characters of vasiform orifice, ...” (Takahashi 1938). In other words, Takahashi was indicating similarities to two aleyrodine genera. As already mentioned, Sampson (1943, 1947) placed *Stenaleyrodes* in the Aleyrodinae, tribe Dialeurodini (Appendix A). Mound and Halsey (1978) listed the genus under Aleurodicinae and synonymized *Dialeurodicus elongatus* Dumbleton, which had already been transferred to *Stenaleyrodes* by Cohic (1968), with *S. vinsoni*. The former authors also stated “the material in the BMNH from Tanzania listed here as *Stenaleyrodes* sp. indet. represents a distinct species, which as in the case of *vinsoni*, can only be distinguished from *Dialeurodicus* at present by the elongate shape of its pupal case”. No syntype materials of *S. vinsoni* were examined, but a paratype of *D. elongatus* and another specimen of *S. vinsoni* collected from the type locality (Reunion Island) and from the host of *S. vinsoni* (palm) as well as Tanzanian samples were studied. These samples lacked both the tarsal claw and apical adhesion pad. The illustration of hind leg in the original description of *S. vinsoni* (Takahashi 1938, Figure 5) is in agreement with this observation. It seems that Mound and Halsey (1978) did not notice the absence of the tarsal claw in *Stenaleyrodes* while comparing it with *Dialeurodicus*, in which the character is present, and only mentioned the elongation of pupal case to distinguish these genera from each other.

From a consideration of puparial characters, there is no reason for *Stenaleyrodes* to be in Aleurodicinae due to its lack of the tarsal claw and lack of compound wax pores and/or agglomerate pores, i.e. the two basic puparial characters of the Aleurodicinae, though the vasiform orifice and the general size are similar to those of aleurodicine members. However, the examination of recently collected material of an undescribed species of *Stenaleyrodes* in BMNH has revealed aleurodicine characters in adults. Therefore, it would seem expedient to retain *Stenaleyrodes* in Aleurodicinae.

All 13 remaining characters with maximum retention indices ($ri=1$) (except characters 22, 24, and 46) are autapomorphic characters of certain genera, e.g. characters 63 (eighth abdominal bifid process) and 82 (chain-like design along some abdominal and thoracic sutures) support monophyly of *Aleurocerus* and *Bellitudo* respectively. Thus, these characters did not provide any phylogenetic information about relationships between genera.

According to Bink-Moenen (1983), character 3 (recurvature of margin) is mostly of value at the genus level, although in our analyses it had an intermediate ri and pretty low ci (Table IV). Furthermore, Mound (1961), who erected the genus *Aleuropteridis*, mentioned that “this genus is near *Tetralicia* and *Aleuropleurocelus* in that the true margin is deflexed ventrally, i.e. the dorsal disc is larger than the ventral surface”. Although in these three genera the character is quite typical, it can also be seen in several species putatively belonging to other genera such as *Bemisia hirta* Bink-Moenen and *Tetraleurodes caulicola* Nakahara, which it is not a specific character of the latter genera. All examined species currently placed in *Tetralicia*, in spite of their sharing this character, were not found to be

monophyletic. *Aleuropteridis* formed a sister group to *Xenaleyrodes* (also shows the character), and among the taxa gathered with *Aleuropteridis* in the same clade (Figure 16), two examined species of *Leucopogonella* lacked this character. In this clade, *Tetraleyrodes selachidentata* Bink-Moenen was described as a species *incertae sedis* (Bink-Moenen 1983). Incidentally, some examined species of *Corbettia* and *Viennotaleyrodes* (both genera were recovered as monophyletic) lacked the character. Thus, character 3 seems to be far less reliable than past authors have suggested.

Submarginal line or fold (character 28) have been thought to be an important character at genus level and has been used by authors both in generic descriptions and identification keys (e.g. Mound 1965; Bink-Moenen 1983; Jesudasan and David 1991; Martin et al. 2000). It separates the submargin from the dorsal disc area and can appear in different forms (see character 29). In general, this character could not gather the genera possessed it in a separate clade, at least those that it appears in an identical shape. It should be noted that *Aleurotuberculatus*, in spite of having the character, was synonymized with *Aleuroclava* (Martin 1999). Although the 15 examined species (seven species had the character) of the latter large genus were not found to be monophyletic in this analysis (Figure 17), but

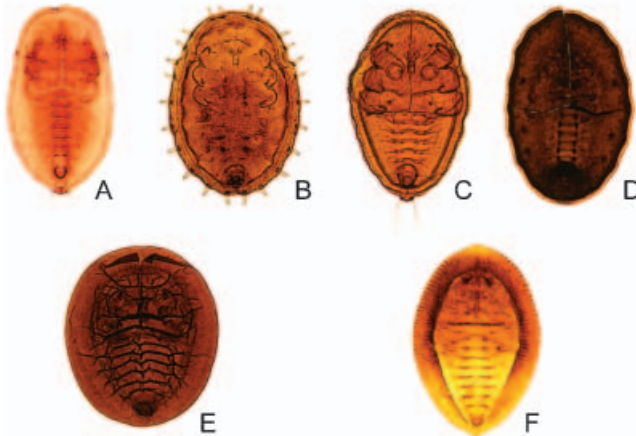
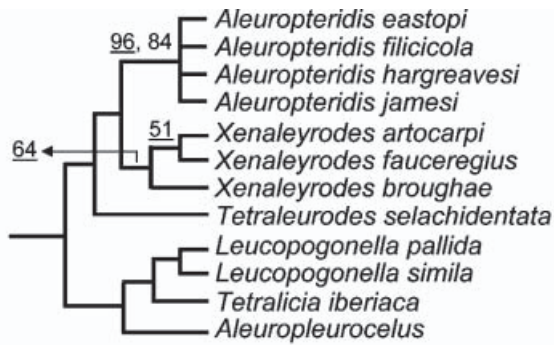


Figure 16. One of the clades of the strict consensus tree of UW-MPTs. Figures above branches show bootstrap and Jackknife (underlined figures) values. (A) *Aleuropteridis filicicola* (Newstead); (B) *Xenaleyrodes artocarpi* Takahashi; (C) *Tetraleyrodes selachidentata* Bink-Moenen; (D) *Leucopogonella simila* Dumbleton; (E) *Tetraticia iberiaca* Bink-Moenen; (F) *Aleuropleurocelus nigrans* (Bemis).

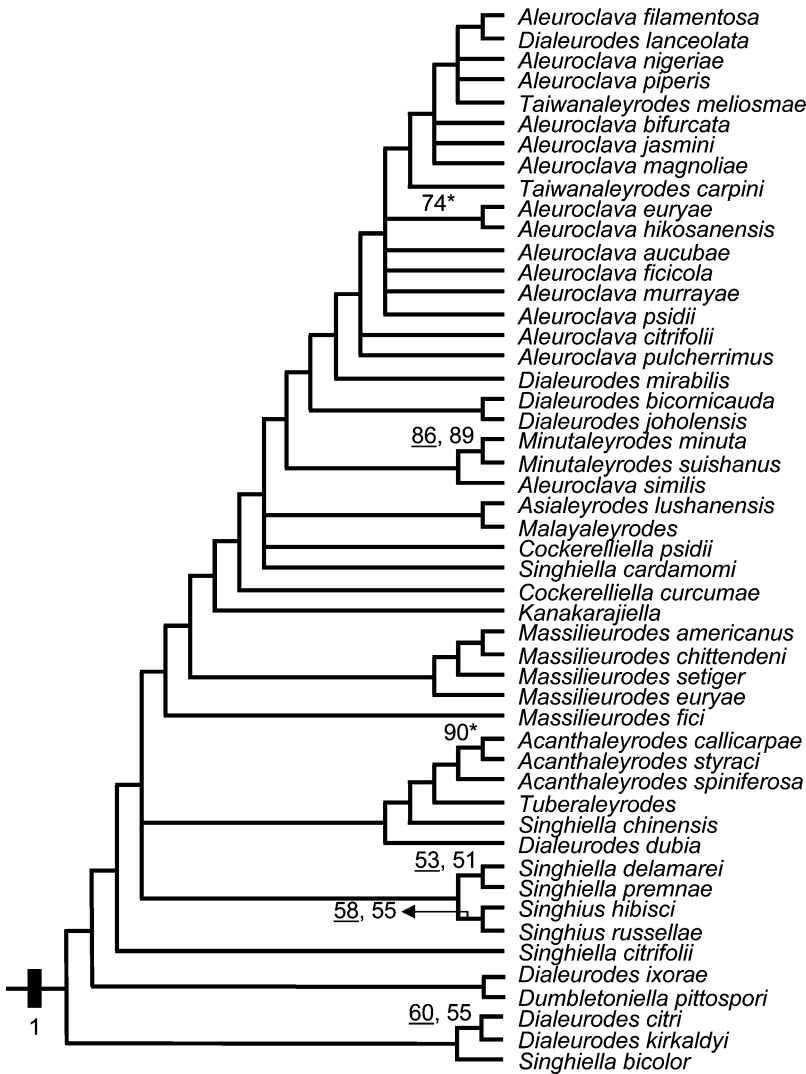


Figure 17. One of the clades of the strict consensus tree of UW-MPTs. Figures above branches show bootstrap and Jackknife (underlined figures) values. Figures with an asterisk indicate the same value for both bootstrap and Jackknife. Black rectangles and the figure below indicate the Bremer support.

Aleuroclava can be considered as a monophyletic genus if some new combinations and synonymy are accepted (see “nomenclatural changes”).

Some binary characters, e.g. 7 (position of cephalic setae), 11 (position of first abdominal setae), 37 (elevation of vasiform orifice), 43 (opercular setae), 49 (lingular setae), 50 (number of lingular setae), 55 (caudal furrow), 57 (caudal ridges), 67 (submarginal glands) and 89 (apical adhesion pad of legs), which have had little attention paid to some of them by aleyrodid systematists, had high retention indices ($0.8 < ri < 1$), of which the four characters, 37, 49, 55 and 57 were important in supporting main clades in the strict consensus tree of UW-MPTs. The general topology of this tree is shown in Figure 18. With a few exceptions all taxa included in clade B (Figure 18) have elevated vasiform orifice (character 37), and considering its presence in some other small clades in the remaining



Figure 18. The general topologies of the strict consensus trees of UW-MPTs (A) and MPTs-SAW-MaxRI (B). See text for the discussion of indicated clades.

part of the tree, e.g. clades II and VI, it can be hypothesized that the character has independently evolved several times. The same hypothesis probably applies to characters 49, 55 and 57. Most taxa in the clade C (Figure 18) lack lingular setae (or the setae are not apparent) (character 49), and those in the clade A lack both caudal furrow (character 55) and caudal ridges (character 57), but as for character 37, some other small clades in the other part of the tree lack these characters too, e.g. clades III and V for character 49, and clades I and IV for both characters 55 and 57.

The applicability of any tribe level classification for the Aleyrodinae seems to be premature. Sampson (1943) was the first to classify aleyrodine genera to five tribes (Appendix A). Whereas, in 1956, in their identification key for the genera of the subfamily, Sampson and Drews (1956) placed almost all described genera by that time in two tribes, Aleyrodini and Dialeurodini, and the other three tribes was each comprised of a single genus. The authors also did not include the sixth tribe proposed by Russell (1947) in the key because of difficulty of finding characters to separate it from the others, and nothing was mentioned about the two tribes proposed by Takahashi (1954). It seems that erecting more tribes, as done by David (1990), with our current poor understanding the significance of puparial characters will not help. For example, none of the genera placed in the Aleyrodini by three different authors (Appendix A) was recovered in the same clade as *Aleyrodes* in this study, and *Indoaleyrodes*, which was placed in the Bemisini by David (1990), formed a sister group to *Aleyrodes* (Figure 15). Bearing in mind that defining a reliable suite of morphological characters at genus level is still problematic and many genera lack objective definitive, it will obviously be difficult to find some synapomorphic characters to define different tribes confidently.

Unfortunately, some strikingly obvious structural apomorphies have probably been independently derived in a number of quite closely related species. Such characters can easily confuse attempts at phylogenetic reconstruction as they are difficult to differentiate from genuine synapomorphies. Often such striking characters are considered to be of “generic worth” and in practice many species which show these kinds of characters are placed in separate monobasic or oligobasic genera (Gauld and Mound 1982). It seems that the ability to develop apomorphies of this type independently in closely related taxa can be found in Aleyrodidae. The taxa of the clade shown in Figure 19 may be good examples. In this clade, three genera, *Aleuroparadoxus*, *Bellitudo*, and *Crescentaleyrodes*, have obvious apomorphies, i.e. characters 76 (modified papillae of dorsal disc area, which are slightly elevated, nearly flat and plate-like), 82 (chain-like design along some abdominal and thoracic sutures) and 87 (submarginal crescent-shaped pores) respectively. Of these three genera, *Crescentaleyrodes* was not recovered as monophyletic and *Aleurotrachelus fumipennis* (Hempel) made it paraphyletic (Figure 19). David and Jesudasan (1987) proposed *Crescentaleyrodes* for *Tetraleurodes semilunaris* Corbett, which had already been transferred to *Aleurolobus* by Bink-Moenen (1983), as well as transferring two species of *Aleurolobus*, *A. monodi* Cohic and *A. paulianae* Cohic to *Crescentaleyrodes*. The main diagnostic character of *Crescentaleyrodes*, as the name implies, is the presence of crescent-shaped (semilunate) pores (Figure 12J) along the submargin. It should be noted that the syntype materials of *A. fumipennis* were not examined and its original description (Hempel 1899) provides no illustration and is inadequate for the positive identification of the species. The examined specimens (from Brazil collected on grasses, the same locality and host plant as the syntypes) identified as *A. fumipennis* in BMNH had submarginal crescent-shape pores (character 87) and were in agreement with the redescription and illustration of *A. fumipennis* provided by Bondar (1923), i.e. belonging to *Crescentaleyrodes* (see also

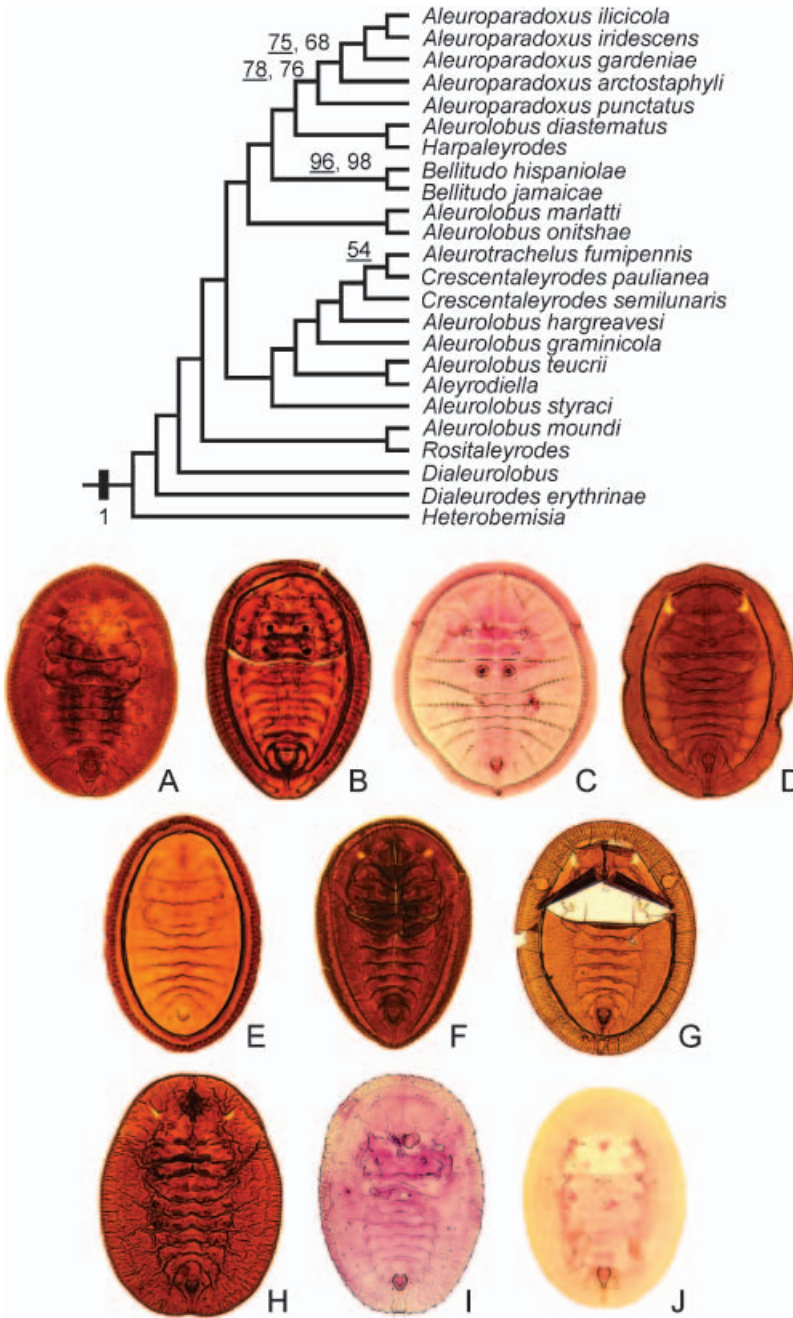


Figure 19. One of the clades of the strict consensus tree of UW-MPTs. Figures above branches show bootstrap and Jackknife (underlined figures) values. Black rectangular and the figure below indicate the Bremer support. (A) *Aleuoparadoxus arctostaphyli* Russell; (B) *Harpaleyrodes tuberculata* Bink-Moenen; (C) *Bellitudo jamaicae* Russell; (D) *Aleurolobus marlatti* (Quaintance); (E) *Crescentaleyrodes semilunaris* (Corbett); (F) *Aleyrodiella lamellifera* Danzig; (G) *Rositaleyrodes oplismeni* (Takahashi); (H) *Dialeurolobus rhammi* Bink-Moenen; (I) *Dialeurodes erythrinae* Corbett; (J) *Heterobemisia alba* Takahashi.

“Nomenclatural changes”). Furthermore, there are three monobasic genera, *Aleyrodiella*, *Harpaleyrodes* and *Rositaleyrodes*, in this clade. The latter was proposed for *Aleurolobus oplismeni* Takahashi by Meganathan and David (1994), who erroneously proposed a new holotype and six paratypes from their samples collected for an extant species of another author. The authors stated “this genus resembles *Aleurolobus* Quaintance & Baker in all the characters, but it differs from it, by having the submargin entirely demarcated from dorsum by a submarginal furrow and thoracic and caudal tracheal combs differentiated from margin by teeth and a pouch like structure”. The entire submarginal furrow can be seen in other species of *Aleurolobus*, e.g. *A. hargreavesi* Dozier, and also the differentiation of tracheal area from the margin is variable in this genus. It seems that there is no reason for proposing a separate genus (see also “Nomenclatural changes”). Danzig (1966) erected *Aleyrodiella* and stated “an extremely distinctive genus whose taxonomic position is uncertain”. The author then discussed its similarity to the three genera, *Aleuoparadoxus*, *Aleurotithius*, and *Trialeurodes* (Table III), as well as its differences from them. These three genera are those that Russell (1947) placed in the *Trialeurodini* (Appendix A) but only *Aleuoparadoxus* was recovered here in the same clade as *Aleyrodiella*. The latter two genera were found here to be more closely related to *Aleurolobus* than to *Trialeurodes* and/or *Aleurotithius* (Table III). The third monobasic genus in this clade, *Harpaleyrodes*, was erected by Bink-Moenen (1983) and, as the author stated, it resembles *Aleurolobus*. One of its differences from *Aleurolobus* is its strongly recurved submargin (Bink-Moenen 1983) (character 3), and as discussed above, this is variable in some genera. It is possible that *Harpaleyrodes* and *Aleurolobus* are congeneric, in spite of the alleged differences. It is also worth mentioning that according to Danzig (1964), *Dialeurolobus* (with two described species so far), which formed a sister group to all discussed taxa in this clade (Figure 19), is allied to *Aleurolobus* and differs from the latter in having no submarginal fold (character 28). The variability of this character has been discussed, above. *Dialeurodes erythrinae* Corbett has also much in common with the members of this clade (Figure 19) rather than with *Dialeurodes*, and its placement in *Dialeurodes* is doubtful (species *incertae sedis*).

As is common in phylogenetic analyses, the sister groups of some clearly monophyletic genera are often paraphyletic assemblages, e.g. *Aleurocerus* rendering *Aleurothrixus* paraphyletic (Figure 20). Seven of the 10 examined species, including the type-species, *Aleurothrixus floccosus* (Maslell), as well as *Hempelia* (see “Nomenclatural changes” for proposed synonymy of *Hempelia* with *Aleurothrixus* as well as the discussion of the other three species of *Aleurothrixus*) were recovered in the same clade as *Aleurocerus*. This clade was relatively highly supported in both bootstrap and parsimony Jackknifing analyses (Figure 20). A close relationship between *Aleurocerus* and *Aleurothrixus* was also postulated by Russell (1986) and the presence of the eighth abdominal bifid process (character 63) (Figure 11A) in *Aleurocerus* its only diagnostic character.

The same was found for *Corbettia*, recovered as monophyletic here, but rendering *Aleuromarginatus* paraphyletic (Figure 15). Interestingly, *Aleuromarginatus* and *Corbettia* also colonize the same host plant family, Fabaceae (see “Results”). Seven of the 10 examined species, including the type-species, *A. tephrosiae* Corbett, were found in the same clade as *Corbettia*. The two other species scored, *A. marginiquus* Martin and *A. nemciae* Martin, are apparently not congeneric with the type-species and this uncertainty was alluded to by the author (Martin 1999). Based on the results of this study, the placement of these species in *Aleuromarginatus* is uncertain (species *incertae sedis*). *A. serdangensis*, as already mentioned, seems to be a misplaced species differing in several characters (e.g. 4, 8, 9, 44 and 94) from the type and the other included species. *A. serdangensis* was recovered in

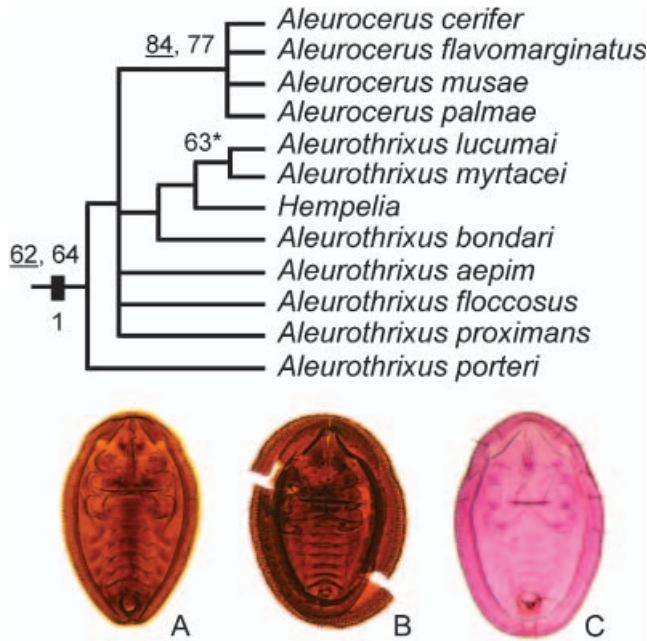


Figure 20. The terminal clade of the strict consensus tree of UW-MPTs. Figures above branches show bootstrap and Jackknife (underlined figures) values. Figure with an asterisk indicate the same value for both bootstrap and Jackknife. Black rectangular and the figure below indicate the Bremer support. (A) *Aleurocerus musae* Russell; (B) *Hempelia chivelensis* Sampson & Drews; (C) *Aleurothrixus proximans* Bondar.

a quite distantly separated clade that includes several diverse species putatively belonging to different genera (Figures 13 and 14).

Monophyly of large genera

Aleuroplatus, *Aleurotrachelus*, and *Tetraleurodes* are genera with many included species and worldwide distribution (Mound and Halsey 1978). Currently each includes many diverse species and lacks a suite of distinguishing characters. Therefore, there is no reason to assume that species currently classified in any of these, especially from different geographical regions, are related to one another. As expected, these genera were not found to be monophyletic, but only for *Aleurotrachelus*, was monophyly rejected statistically. The reason for this may be the lack of distinct generic characters. In other words, the species of these genera just have combinations of characters shared with various other genera. Regarding *Aleurotrachelus*, it should be noted that the specimens checked as *A. fumipennis*, as discussed above, had the typical character of *Crescentaleyrodes* and in the constraint analysis it was not excluded from *Aleurotrachelus*. Thus, it is quite likely that this species contributed largely to the statistical rejection of monophyly of *Aleurotrachelus*. To test this, another constraint analysis was performed for this genus excluding *A. fumipennis*. The length of the MPTs was 2773 compared with 2730 for the UW-MPTs. The result of the Templeton test (500 trees were compared) showed that the constrained trees were not consistently significantly longer with *P* values for the majority of the trees above 0.05 ($P=0.0390-0.0793$; see also Table II). Such interpretation may also apply to *Dialeurodes*, which in spite of many of its some species having been transferred to *Massilieurodes* and *Singhiella* by Jensen (2001), it is still a large and heterogeneous genus.

In contrast, although monophyly of *Trialeurodes* was not also rejected statistically, the same interpretation does not seem to apply. *Trialeurodes* is quite well characterised and most of its species are certainly of New World origin (Mound 1984). *T. vaporariorum*, which has long been known as a worldwide pest, almost certainly evolved in the south western part of North America (Russell 1948). In the strict consensus tree of MPTs-SAW-MaxRI (Figure 14), nine of the 11 examined species including the type-species, *T. pergandei*, as well as the closely related monobasic genus *Venezaleurodes* (probably should be synonymized with *Trialeurodes*) were recovered as a monophyletic group. These species were unresolved in the strict consensus tree of UW-MPTs (Figure 13). The probabilities that this genus is monophyletic (Templeton test: $P=0.5590-0.7035$) are much higher than for the other constrained genera (Table II). The factors mentioned above for *Trialeurodes* may also be applicable to *Bemisia*. According to Mound (1984), it seems probable that the *Bemisia* group primarily originated in the Palaearctic, though it was previously suggested that *B. tabaci* was Indian region in origin (Mound 1965). The main problem with this genus is that in spite of including the well-known pest, *B. tabaci*, it is not well defined. For this reason, it is expected that the genus includes at least several misplaced species which resulted its monophyly to be rejected statistically.

Host plant relationship

Relationships between whiteflies and their host plants are unclear. According to Mound and Halsey (1978), since the majority of aleyrodids are known only from the Angiosperms, it seems unlikely that this family evolved until after the radiation of the flowering plants. These authors compared the Cronquist's (1968) plant classification with the host plants of whiteflies and deduced that most whitefly species were recorded from plants belonging to four smaller, less highly evolved subclasses. The authors then stated: "As the Asteridae and Rosidae are the most advanced groups of higher plants, it may be that the aleyrodids did not evolve at the same rate but retained a relationship with the more primitive groups of plants. Alternatively the relationship may be a reflection of the geographical distributions of the organisms, the Asteridae and Rosidae being most common in temperate regions, whereas the Aleyrodidae are mainly a tropical group."

Based on the system of the Angiosperm Phylogeny Group (APG) (Bremer et al. 2003), 328 of the 439 whitefly species examined in this study are known from the two advanced groups, i.e. asterids (including euasterids I and euasterids II) and rosids (including eurosids I and eurosids II). In this system, euasterids II and euasterids I are two of the most recently derived groups. More than 100 of the examined whitefly species are known from these groups (21 from euasterids II, 78 from euasterids I, and 17 from both) (Appendix E). The positions of these species in the strict consensus tree of UW-MPTs as well in the MPTs-SAW-MaxRI show no pattern. Thus, parallel diversification with host plants and whiteflies is not apparent and it appears that widespread host switching has obscured any basic associations. Having said this, the evolution of aleyrodid host plant affiliations does not seem to be totally random as some groups have species feeding on related plants.

Nomenclatural changes

The results obtained from phylogenetic analyses presented above, indicate the great need for reclassification of the Aleyrodinae. It is quite possible that those species whose pupal cases differ considerably from each other, nevertheless have adults showing apparently

unique apomorphies and therefore they should be placed in the same genus (Bink-Moenen 1992, discussing the genus *Aleuroviggiatus*). On the other hand, adults can differ markedly despite of great similarities in their pupal cases, and so are placed in different genera (Russell 2000, discussing the genera *Aleurocybotus* and *Vasdavidius*). Since adults for the majority of described species are unknown, it is not desirable to change the current genus concepts defined based on pupal case character only, unless the true significance of the latter is known. This will be the next necessary step in improving the taxonomy of whiteflies.

Checking such a large number species, belonging to more than 100 genera, revealed the necessity for proposing some nomenclatural changes. These are only proposed for those taxa where the necessity of changing their current taxonomic positions has been supported by the phylogenetic analysis, in addition to the convictions obtained during the visual examination of specimens. Those genera mentioned here as New genus 1 and New genus 2 will be described somewhere else.

Aleuroclava Singh, 1931

Type-species: *Aleuroclava complex* Singh, 1931.

Aleurotuberculatus Takahashi, 1932. Type-species: *Aleurotuberculatus gordoniae* Takahashi, 1932 [synonymized by Martin 1999].

Martiniella Jesudasan & David, 1990. Type-species: *Aleurotuberculatus canangae* Corbett, 1935 [synonymized by Martin 1999].

Taiwanaleyrodes Takahashi, 1932. Type-species: *Taiwanaleyrodes meliosmae* Takahashi, 1932. **Syn. n.**

Comments: Takahashi (1932) erected *Aleurotuberculatus* mentioning that it was closely allied to *Aleuroclava* and may be a synonym of it. In the same article, he erected *Taiwanaleyrodes* and stated “allied to *Aleurotuberculatus* Takah., but differs from it in lacking dorsal tubercles and thoracic tracheal clefts, but in possessing a distinct marginal rim on the venter, and also distinguishable from ...” (Takahashi 1932). Takahashi (1939) in the original description of *Taiwanaleyrodes carpini* Takahashi stated “cephalothorax with a pair of large rounded tubercle-like markings on the pronotum, and also on the mesonotum, and 2 pairs of similar ones on the metanotum, ..., thoracic tracheal folds very short, the clefts indistinct, very shallow, wide, without teeth”. In this article, the author transferred *Aleurotuberculatus pyracanthae* Takahashi to *Taiwanaleyrodes* (apparently Mound and Halsey (1978) did not notice this transformation and listed the latter species under *Aleurotuberculatus* and/or did not accept it, but mentioned nothing), although in the original description of *A. pyracanthae* stated “venter with no rim” (Takahashi 1933). It is clear that the two species of *Taiwanaleyrodes*, which placed in this genus by the author of the genus, are not quite agree with the type-species, *T. meliosmae* Takahashi, in having the three diagnostic characters. Examination of many specimens of *Aleuroclava*, in addition to those included in this analysis, indicated that the dorsal tubercles and thoracic tracheal clefts definition is so variable and these characters should not be used for the separation of these genera. In the current study the type-species of *Taiwanaleyrodes*, as well as its one other included species were recovered within *Aleuroclava* (Figure 17). *Taiwanaleyrodes* is, therefore, here regarded as a junior synonym of *Aleuroclava* **syn. n.**

Jesudasan and David (1990) proposed *Minutaleyrodes* for four species formerly included within *Aleurotuberculatus*. The genus was provisionally retained for its included species by

Martin (1999), who synonymized *Aleurotuberculatus* and *Martiniella* with *Aleuroclava*. *Aleuroclava similis* (Takahashi) formed a sister group to *Minutaleyrodes* in this study (Figure 17). Of the characters specified for *Minutaleyrodes* by Jesudasan and David (1990), demarcation of the submargin on the ventral surface, can be seen in some species of *Taiwanaleyrodes* (discussed above). The shape of transverse moulting suture is the same that of in *A. similis*, but the size of this species is not similar to *Minutaleyrodes* and submargin not ventrally demarcated in *A. similis*. Unlike *Aleuroclava*, the first abdominal setae in *Minutaleyrodes* are absent (not mentioned by the authors) but some species of the former genus, e.g. *A. similis* and *A. nigeriae* (Mound), also lack these setae. In the latter species, these setae are sometimes present (Bink-Moenen 1983). Therefore, *Minutaleyrodes* seems to be congeneric with *Aleuroclava*, but because of the three species of *Dialeurodes* recovered between these two genera (see Figure 17), it is not synonymized. These three species currently accommodated in *Dialeurodes*, are not quite typical for their current genus nor for *Aleuroclava*, and it appears they have much in common with *Aleuroclava*, especially *D. mirabilis* Takahashi, rather than *Dialeurodes*.

***Aleuroclava lanceolata* (Takahashi) comb. n.**

Dialeurodes lanceolata Takahashi, 1949. Holotype, Riouw (Riau) Islands (south of Singapore): Rempang [not examined].

Material examined: Singapore: Botanic Gardens, 15.ix.1980, (D.H. Murphy), ex: *Bhesa panicula* (2 slides).

Comments: Specimens checked were in agreement with the original description and illustration, and indicate that the inclusion of this species in *Aleuroclava* is appropriate. As stated by Takahashi (1949), this species is not a typical form of *Dialeurodes*. *A. lanceolata* is closely related to *A. filamentosa* (Corbett) (Figure 17).

***Aleurolobus* Quaintance & Baker, 1914**

Type-species: *Aleurodes marlatti* Quaintance, 1903.

Rositaleyrodes Meganathan & David, 1994. Type-species: *Aleurolobus oplismeni* Takahashi, 1931. **Syn. n.**

Comments: As discussed before, according to Meganathan and David (1994), *Rositaleyrodes* resembles *Aleurolobus* in all the characters, but it differs from it, by having the submargin entirely demarcated from dorsum by a submarginal furrow and thoracic and caudal tracheal combs differentiated from margin by teeth and a pouch like structure. The latter character is variable in *Aleurolobus* and the entire submarginal furrow can be seen in other species of this genus, e.g. *A. hargreavesi*. *Rositaleyrodes* is therefore here regarded as a junior synonym of *Aleurolobus* **syn. n.**

***Aleurothrixus* Quaintance & Baker, 1914**

Type-species: *Aleyrodes howardi* Quaintance, 1907, a junior synonym of *Aleurodes floccosa* Maskell, 1895.

Hempelia Sampson & Drews, 1941. Type-species: *Hempelia chivelensis* Sampson & Drews, 1941. **Syn. n.**

Comments: As already discussed, seven of the 10 examined species of *Aleurothrixus* including the type-species, *A. floccosus* (Maslell), as well as *Hempelia* were recovered in a monophyletic clade with *Aleurocerus* (Figure 20). The syntype material of the type-species of monobasic genus *Hempelia* showed no clear differences from *Aleurothrixus* that appear to be of generic significance and here *Hempelia* is regarded as a junior synonym of *Aleurothrixus* **syn. n.**

Aleurothrixus antidesmae Takahashi and *A. smilaceti* Takahashi are closely related to each other and do not seem to be congeneric with the type-species. A different genus is needed for them; almost certainly a new one. The situation is the same for the American species, *A. interrogationis* (Bemis), but it differs from these two Asian species. It was recovered within some species of *Aleuroplatus* and its placement in *Aleurothrixus* is doubtful (species *incertae sedis*).

***Crescentaleyrodes* David & Jesudasan, 1987**

Type-species: *Tetraleurodes semilunaris* Corbett, 1926.

***Crescentaleyrodes fumipennis* (Hempel) comb. n.**

Aleurodes fumipennis Hempel, 1899. Syntypes, Brazil: São Paulo [not examined].

Aleurotrachelus fumipennis (Hempel) Quaintance & Baker, 1914.

Material examined: Brazil: Mato Grosso, 25.iii.1968, (O.W. Richards), ex: undetermined grass (2 slides).

Comments: as already mentioned in detail, the redescription and illustrations provided by Bondar (1923) as well as the specimens examined, clearly indicate the appropriate inclusion of this species in *Crescentaleyrodes*. This species is very close to *C. paulianae* (Cohic) and maybe they are synonym, but this need to check the type material of the latter species. The specimens of *C. paulianae* studied for this study were from the same locality and host plant of type materials and showed slightly differences from *C. fumipennis*.

***Pseudozaphanera* Manzari gen. n.**

Type-species: *Aleurodes niger* Maskell, 1895.

Description: Pupal case. Cuticle black. Margin toothed, thoracic and caudal tracheal pore areas not differentiated from margin. Submargin not separated from dorsal disc area by a line or fold. Longitudinal moulting suture reaching margin, transverse moulting suture curving anteriorly and reaching margin. Abdomen with exaggerated intersegmental sutures extending into outer subdorsum. Abdominal rhachis present or absent. Dorsum usually punctuated by geminate pore/porettes. Vasiform orifice cordate or subcordate, often elevated. Operculum almost filling vasiform orifice. Caudal furrow absent or faintly indicated. Eyespots absent or present.

Chaetotaxy: Cephalic and first abdominal setae absent, eight abdominal and caudal setae present. A row of submarginal setae present but their total number as well as their number on abdomen and cephalothorax variable (6–11, usually 7/8 pairs excluding the caudal setae), occasionally with a few pair subdorsal setae. All setae very small and often capitate.

Venter: Thoracic and caudal tracheal folds present. Antennae long, reaching middle of middle legs or more, male antennae much longer.

Distribution: Australia.

Hosts: Fabaceae.

Etymology: This genus is named to reflect its superficial similarity to *Zaphanera*.

Species included: *Pseudozaphanera niger* (Maskell) **comb. n.**; *Pseudozaphanera papyrocarpae* (Martin) **comb. n.**; *Pseudozaphanera rhachisreticulata* (Martin) **comb. n.**; *Pseudozaphanera splendida* (Martin) **comb. n.**; *Pseudozaphanera wariensis* (Martin) **comb. n.**

Comments: except the type-species, all above species included in this genus were originally been described in *Zaphanera*. The type-species has already been transferred to *Aleurolobus*, *Tetraleurodes* and *Zaphanera* by Quaintance and Baker (1914), Dumbleton (1956) and Martin (1999), respectively. *Pseudozaphanera* differs from *Zaphanera* in the number of submarginal setae as well as their shape, absence of cephalic setae, position of eight abdominal setae, shape of transverse moulting suture, and length of antennae. In *Zaphanera*, there are five pairs of setae (excluding the caudal setae), two pairs on the cephalothorax and three pairs on the abdomen, and these setae are not capitate. Antennae do not reach beyond front legs, and the eighth abdominal setae are located adjacent to postero-lateral margin of vasiform orifice. In three of the four described species (examined) retained in *Zaphanera*, including the type-species, *Z. cyanotis*, cephalic setae are present, and based on the redescription provided by David and Subramaniam (1976), they are not discernible in *Z. publicus* (Singh) (not examined). In *Zaphanera* abdominal segmentation is distinct only in the submedian area in contrast to *Pseudozaphanera* (see also tribe Zaphanerini).

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Appendix A: the tribes of the Aleyrodinae and their included genera according to different authors

Based on	Tribe	Included genera
Sampson (1943)	Aleurochitonini	<i>Aleurochiton</i> Tullgren
	Aleyrodini	<i>Acaudaleyrodes</i> Takahashi ¹ , <i>Aleurocanthus</i> Quaintance & Baker, <i>Aleurocerus</i> Bondar, <i>Aleurocybotus</i> Quaintance & Baker, <i>Aleuromarginatus</i> Corbett, <i>Aleuromigda</i> Singh, <i>Aleuropleurocelus</i> Drews & Sampson ² , <i>Aleuroputeus</i> Corbett, <i>Aleurothrixus</i> Quaintance & Baker, <i>Aleurotithius</i> Quaintance & Baker, <i>Aleurotrachelus</i> Quaintance & Baker, <i>Aleurotulus</i> Quaintance & Baker, <i>Aleyrodes</i> Latreille, <i>Bulgarialeyrodes</i> Corbett, <i>Corbettia</i> Dozier, <i>Crenidorsum</i> Russell ³ , <i>Frauenfeldiella</i> Gomez-Menor ⁴ , <i>Hempelia</i> Sampson & Drews, <i>Hesperaleyrodes</i> Sampson, <i>Heteraleyrodes</i> Takahashi ⁵ , <i>Laingiella</i> Corbett, <i>Luederwaldtiana</i> Hempel, <i>Mexicaleyrodes</i> Sampson & Drews, <i>Nealeyrodes</i> Hempel, <i>Neoleurodes</i> Bondar, <i>Pentaleyrodes</i> Takahashi, <i>Simplaleyrodes</i> Goux ⁶ , <i>Tetraleyrodes</i> Cockerell, <i>Tetralicia</i> Harrison, <i>Zaphanera</i> Corbett
	Dialeurodini	<i>Acanthaleyrodes</i> Takahashi, <i>Acanthobemisia</i> Takahashi, <i>Africaleyrodes</i> Dozier, <i>Aleuroclava</i> Singh, <i>Aleuoglandulus</i> Bondar, <i>Aleurolobus</i> Quaintance & Baker, <i>Aleuroparadoxus</i> Quaintance & Baker, <i>Aleuroplatus</i> Quaintance & Baker, <i>Aleuroporosus</i> Corbett, <i>Aleurotuberculatus</i> Takahashi, <i>Anomaleyrodes</i> Takahashi & Mamet ⁷ , <i>Asialeyrodes</i> Corbett, <i>Asterobemisia</i> Trehan, <i>Asterochiton</i> Maskell, <i>Bellitudo</i> Russell ⁸ , <i>Bemisia</i> Quaintance & Baker, <i>Corbettella</i> Sampson, <i>Dialeurodes</i> Cockerell, <i>Dialeurolonga</i> Dozier ⁹ , <i>Dialeuropora</i> Quaintance & Baker, <i>Dialeurotrachelus</i> Takahashi ¹⁰ , <i>Malayaleyrodes</i> Corbett, <i>Metaleyrodes</i> Sampson, <i>Mixaleyrodes</i> Takahashi, <i>Neoleurolobus</i> Takahashi ¹¹ , <i>Neoleurotrachelus</i> Takahashi & Mamet ¹² , <i>Neobemisia</i> Visnya ¹³ , <i>Nipaleyrodes</i> Takahashi ¹⁴ , <i>Parabemisia</i> Takahashi ¹⁵ , <i>Paraleurolobus</i> Sampson & Drews, <i>Pealius</i> Quaintance & Baker, <i>Plataleyrodes</i> Takahashi & Mamet ¹⁶ , <i>Pseudaleurolobus</i> Hempel, <i>Pseudaleyrodes</i> Hempel, <i>Rhachisphora</i> Quaintance & Baker ¹⁷ , <i>Roucasia</i> Goux ¹⁸ , <i>Setaleyrodes</i> Takahashi, <i>Singhiella</i> Sampson, <i>Stenaleyrodes</i> Takahashi, <i>Taiwanaleyrodes</i> Takahashi, <i>Triaurodes</i> Cockerell, <i>Trichoaleyrodes</i> Takahashi & Mamet ¹⁹ , <i>Tuberaleyrodes</i> Takahashi, <i>Xenaleyrodes</i> Takahashi, <i>Xenobemisia</i> Takahashi ²⁰
	Neomaskellini	<i>Neomaskellia</i> Quaintance & Baker
Russell (1947)	Siphoninini	<i>Siphoninus</i> Silvestri
	Trialeurodini	<i>Aleuroparadoxus</i> , <i>Aleurotithius</i> , <i>Triaurodes</i> , <i>Venezaleyrodes</i> Russell ²¹
Takahashi (1954)*	Aleurocanthini	<i>Aleurocanthus</i> , <i>Aleurotrachelus</i> , <i>Pentaleyrodes</i>
	Aleurolobini	<i>Acanthobemisia</i> , <i>Aleurolobus</i> , <i>Apobemisia</i> Takahashi, <i>Bemisia</i> , <i>Parabemisia</i>
	Aleyrodini	<i>Aleyrodes</i> , <i>Neopealius</i> Takahashi, <i>Odontaleyrodes</i> Takahashi, <i>Pealius</i> , <i>Setaleyrodes</i>
	Dialeurodini	<i>Aleurotuberculatus</i> , <i>Dialeurodes</i> , <i>Rhachisphora</i> , <i>Taiwanaleyrodes</i>
David (1990)**	Aleurocanthini	<i>Aleurocanthus</i>
	Aleurolobini	<i>Africaleyrodes</i> ²² , <i>Aleurolobus</i> , <i>Aleuropapillatus</i> Regu & David ²³ , <i>Asterochiton</i> , <i>Crescentaleyrodes</i> David & Jesudasan, <i>Orientalleyrodes</i> David ²⁴
	Aleuroplatini	<i>Aleuroplatus</i> , <i>Moundiella</i> David
	Aleyrodini	<i>Aleurocybotus</i> , <i>Aleuromarginatus</i> , <i>Aleurotrachelus</i> , <i>Aleurotulus</i> , <i>Aleyrodes</i>
	Bemisini	<i>Bemisia</i> , <i>Indoaleyrodes</i> David & Subramaniam, <i>Neopealius</i> , <i>Odontaleyrodes</i> , <i>Pealius</i> , <i>Setaleyrodes</i>
	Dialeurodini	<i>Aleuroclava</i> , <i>Aleurotuberculatus</i> , <i>Asialeyrodes</i> , <i>Cockerelliella</i> Sundararaj & David ²⁵ , <i>Dialeurodes</i> , <i>Dialeurolonga</i> , <i>Dialeuromada</i> Quaintance & Baker ²⁶ , <i>Dialeuropora</i> , <i>Fippataleyrodes</i> Sundararaj & David ²⁷ , <i>Gigaleyrodes</i> Quaintance & Baker ²⁸ , <i>Kanakarajiella</i> David & Sundararaj ²⁹ , <i>Martiniella</i> Jesudasan & David ³⁰ , <i>Minualeyrodes</i> Jesudasan & David ³¹ , <i>Rabdstigma</i> Quaintance & Baker ³² , <i>Rhachisphora</i> , <i>Rusostigma</i> Quaintance & Baker, <i>Singhiella</i> , <i>Singhius</i> Takahashi, <i>Taiwanaleyrodes</i>
	Lipaleyrodini	<i>Lipaleyrodes</i> Takahashi
	Neomaskellini	<i>Acaudaleyrodes</i> , <i>Neomaskellia</i>
	Siphoninini	<i>Siphoninus</i>
	Tetraurodini	<i>Tetraurodes</i>
Trialeurodini	<i>Triaurodes</i>	
Zaphanerini	<i>Zaphanera</i>	

*The whitefly genera of Japan only. **The whitefly genera of India only.

1, 4–7, 9–20: Added by Sampson and Drews in 1956; 2: Added by Drews and Sampson in 1956; 3, 8: Added by Sampson in 1947; 21: Added by Russell in 1967; 22–24: Added by Regu and David in 1993; 25–32: Added by David and Sundararaj in 1993.

Appendix B: characters used to identify the tribes of Aleyrodinae according to different authors

Based on	Tribe	Characters
Sampson ¹ (1943); corrected in 1947	Aleurochitonini	Lingula elongate, much longer than wide. Dorsum completely covered with simple pores. Adults with radial ₁ , radial sector, and cubital veins in fore wing.
	Aleyrodini	Lingula elongate, much longer than wide. Dorsum with relatively few simple pores. Thoracic tracheal folds, combs, pores and anal fold absent. Adults with radial sector and cubital veins in fore wing and with tarsal paronychium.
	Dialeurodini	Lingula elongate, much longer than wide. Dorsum with relatively few simple pores. Thoracic tracheal folds and combs or pores, or pores or combs only, and anal fold, or only anal fold, present. Adults with radial sector and cubital veins in fore wing and with tarsal paronychium.
	Neomaskellini	Lingula extremely short, hardly longer than wide. Adults with radial sector vein only in fore wing.
	Siphoninini	Lingula elongate, much longer than wide. Dorsum with elongate, siphon-like wax tubes and with relatively few simple pores. Adults with radial sector and cubital veins in fore wing and lacking tarsal paronychium.
Russell ² (1947)	Trialeurodini	Main points: a submarginal row of disc pores and porettes with which are associated variously conspicuous papillae. Submargin and subdorsum not separated by a furrow. Submarginal setae shorter than the width of a submarginal ridge or without distinguishable setae. Vasiform orifice ending abruptly, its sides without minute spines. Thoracic tracheal folds without angular markings and usually without spines. Antennae not reaching to posterior thoracic spiracles.
Takahashi ² (1954)	Aleurocanthini	Tracheal pores or clefts absent, tracheal combs sometimes developed. Vasiform orifice not notched at the hind end, rounded, not elongated, sometimes elevated; lingula concealed under the operculum. Caudal furrow absent. Eyespots present in some species. Seventh abdominal segment nearly as long as, or a little shorter than, the six.
	Aleurolobini	Tracheal pores or clefts absent, tracheal combs sometimes developed. Vasiform orifice not notched at the hind end, subcordate, triangular or truncated at the hind end, elongated in some species; lingula exposed, knobbed and with a pair of long setae, sometimes large, knobbed part of lingula elongate, much longer than wide. Caudal furrow sometimes developed. Eyespots present in some species. Seventh abdominal segment much shortened at the median area in many genera.
	Aleyrodini	Tracheal pores or clefts absent, tracheal combs sometimes developed. Vasiform orifice not notched at the hind end, subcordate, triangular or truncated at the hind end, elongated in some species; lingula exposed, knobbed and with a pair of long setae, sometimes large, knobbed part of lingula globular, not distinctly longer than wide. Caudal furrow sometimes developed. Eyespots present in some species. Seventh abdominal segment much shortened at the median area in many genera.
	Dialeurodini	Tracheal pores or clefts usually present, if absent, vasiform orifice definitely notched at the hind end and caudal furrow distinctly defined, or lateral ridges (rhachis) developed on the abdomen and dorsum with many short spine-like setae, many of which are capitate. Lingula usually small, wanting long setae. Tracheal combs and eyespots lacking.

Appendix B. (Continued).

Based on	Tribe	Characters
David ² (1990)	Aleurocanthini	Thoracic and caudal tracheal combs, clefts, or pores or furrows absent. Vasiform orifice elevated, rounded; lingula concealed by operculum. Margin with a single row of teeth or smooth. Submargin not separated from dorsal disc. Many prominent spines on the dorsum, often longer.
	Aleurolobini	Thoracic and caudal tracheal combs, clefts, or pores or furrows present. Submargin separated from dorsal disc. Margin generally toothed. Seventh abdominal segment much shortened at the median area in many genera. Eighth abdominal segment often trilobed. Caudal furrow sometimes discernible. Vasiform orifice nearly triangular or subcordate; lingula knobbed and exposed.
	Aleuroplatini	Tracheal pores, clefts, folds, furrows absent but thoracic and caudal tracheal combs indicated. Submargin generally not separated from dorsal disc, if separated, eighth abdominal segment not trilobed. Margin toothed. Pores on dorsum variously distributed. Dorsum with a prominent central ridge terminating cephalad in a more or less arrow-shaped figure, or elevated and fringed with rounded protrusions around. Lingula usually concealed.
	Aleyrodini	Thoracic and caudal tracheal combs, clefts, or pores or furrows absent. Vasiform orifice not elevated; lingula generally exposed, knobbed, but in some concealed. Margin toothed or crenulate. Dorsum with rhachis and a pair of longitudinal fold in some genera. Seventh abdominal segment much shortened at the median area in many genera. Caudal furrow sometimes discernible.
	Bemisini	Tracheal pores, clefts, folds, furrows present but combs absent. Submargin generally not separated from dorsal disc, if separated, eighth abdominal segment not trilobed. Rhachis present or absent. Dorsum generally with setae/tubercles. Submargin with row of papillae. Margin with series of setae in some genera. Vasiform orifice subcordate or triangular or subrectangular, sometimes in a ribbed pyriform pit; lingula long or spatulate, setose and exposed.
	Dialeurodini	Tracheal pores, clefts, folds, furrows present but combs absent. Submargin generally not separated from dorsal disc, if separated, eighth abdominal segment not trilobed. Rhachis present or absent. Dorsum generally with setae/tubercles. Submargin with row of papillae. Margin with series of setae in some genera. Vasiform orifice cordate or subcordate; lingula usually small, concealed and wanting setae.
	Lipaleyrodini	Thoracic and caudal tracheal combs, clefts, or pores or furrows present. Submargin with wax plates in cluster arranged in a row.
	Neomaskellini	Thoracic and caudal tracheal combs, clefts, or pores or furrows absent. Vasiform orifice elevated, elongately elliptic or transversely oval; operculum transversely rectangular and extremely short, exposing lingula. Margin with a single row of teeth or smooth. Submargin not separated from dorsal disc. Rhachis sometimes present. Submargin in some with a row of setae.
	Siphoninini	Thoracic and caudal tracheal combs, clefts, or pores or furrows absent. Dorsum with elongate siphon-like wax tubes.
	Tetraleurodini	Thoracic and caudal tracheal combs, clefts, or pores or furrows absent. Vasiform orifice elevated, subcordate or rounded; lingula concealed by operculum. Margin with a single row of teeth or smooth. Submargin separated from dorsal disc.
	Trialeurodini	Tracheal pores, clefts, folds, furrows absent. Submargin generally not separated from dorsal disc, if separated, eighth abdominal segment not trilobed. Margin dentate, smooth or crenulate. Submargin with a row of disc pores and porettes with variously associated conspicuous papillae. Lingula exposed with a pair of setae. Caudal furrow discernible.
	Zaphanerini	Thoracic and caudal tracheal combs, clefts, or pores or furrows absent. Vasiform orifice elevated; lingula concealed by operculum. Margin with two rows of teeth. Submargin not separated from dorsal disc.

1. Pupal case and adult characters.

2. Pupal case characters only.

Appendix C: checklist of species examined for cladistic analysis

Note: Underlined words and/or figures from slide labels were interpreted with some uncertainty. The list is arranged alphabetically based on the most recent name

Aleyrodinae**1. *Acanthaleyrodes* Takahashi**

A. callicarpae Takahashi

Taiwan: Taichung, 18.xii.1968, (Y.C. Chang), ex: Grape, 2/69 (5 slides).

Taiwan: Taichung, 2.xi.1968, (Y.C. Chang), ex: *Vitis vinifera* (3 slides).

A. spiniferosa (Corbett)

Syntype, Malaya: Kuala Lumpur, 16.ii.1928, (G.H. Corbett), ex: *Diospyros* sp., B.M. 1932-511.

A. styraci Takahashi

Hong Kong: Pok Fu Lam, Reservoir Road, at Peak Road, 12.xii.2001, (J.H. Martin 7563), ex: *Rubus reflexus* (3 slides).

2. *Acanthobemisia* Takahashi

A. distylii Takahashi

Japan: Nagasaki, 27.viii.1957, (R. Takahashi), ex: *Distylium racemosum*, 86/62, B.M. 1962-401 (4 slides).

3. *Acaudaleyrodes* Takahashi

A. africana (Dozier)

Syntype, Belgian Congo: Barumbu, i.1926, (J. Ghesquière), ex: *Desmodium* sp., B.M. 1933.

Nigeria: Ibadun, 20.i.1914, (W.A. Lambourn), ex: herbaceous plant, 204/61, B.M. 1916-190 (2 slides).

A. rachipora (Singh)

Syntypes, Egypt: Meadi, 12.viii.1931, ex: Pomegranate.

Syntype, Egypt: Myniah, 14.xi.1931, ex: *Alhagi* sp.

Cyprus: 10.x.1928, (H.M. Morris), ex: Pomegranate, I.B.E. 2052, 36/61, B.M. 1961-4 (3 slides).

Iraq: Abu-Ghraib, 23.xii.1961, ex: *Zizyphus* sp., C.I.E. 6763, 71/62, B.M. 1962-3 (2 slides).

Sudan: Red Sea Prov. nr Mismar, Atbara-Haiya Jct., 31.iii.1981, (J.H. Martin 3226), ex: *Calotropis procera*.

Sudan: S. Darfur, Ed Da'ain, 3.v.1981, (J.H. Martin 3373), ex: *Calotropis procera*.

India: Madras, Coimbatore, 23.viii.1962, ex: *Prosopis* sp., C.I.E. 15456.

A. tuberculata Bink-Moenen

Holotype, Chad: Bebedjia, 395 m, 16.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Anogeissus leiocarpus*, B.M. 1977-471.

Paratypes, Chad: Boro, 453 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Bridelia ferruginea*, B.M. 1977-471.

4. *Acutaleyrodes* Takahashi

A. palmae Takahashi

Syntypes, Réunion Island: Saint Philippe, xii.1955, (R. Paulian), ex: *Acanthophoenix rubra*.

5. *Africaleurodes* Dozier

A. coffeacola Dozier

Syntypes, Belgian Congo: Lodja, iii.1928, (J. Ghesquière), ex: *Coffea robusta*, B.M. 1933.

Nigeria: Bida, 26.x.1960, (M.O. Ezeigwe), ex: indet.

Aldabra Island: Middle camp, 15.viii.1974, (L.F.H. Merton), ex: *Canthium* sp., 3/74 (5 slides).

A. fulakariensis Cohic

Syntype, Congo: Brazzaville, Fulakary, 28.iii.1965, (R. Paulian), ex: *Neosloetiopsis kamerunensis*, B.M. 1965-305.

A. hexalobi Bink-Moenen

Holotype+paratypes, Chad: Bentam, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*, B.M. 1977-471.

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 6.xi.1989, (L.D.C. Fishpool), ex: *Cryptolepis sanguinolenta*.

A. loganiaceae Dozier

Syntype, Belgian Congo: Sankuru, i.1928, (J. Ghesquière), ex: Loganiaceae, B.M. 1933.

A. ochraceae Dozier

Syntype, Belgian Congo: Kole, 8.i.1928, (J. Ghesquière), ex: *Ochna* sp., B.M. 1933.

Nigeria: Ibadan, viii.1966, (E.A. James), ex: *Cola nitida*, 22/66.

6. *Agrostaleyrodes* Ko

A. arcamus Ko

Paratype, Taiwan: Wulai, 13.viii.1990, (C.C. Ko), ex: *Miscanthus* sp.

7. *Aleurocanthus* Quaintance & Baker

A. banksiae (Maskell)

Australia: Aldingo Scrub, xii.1967, (L.A. Mound), ex: *Banksia marginata*, 39/71 (3 slides).

Australia: Kangaroo Island, 18.ii.1990, (J.H. Martin 5661), ex: *Banksia ornata*, B.M. 1998-50.

Australia: nr Victor Harbour, 20.iv.1977, (P. Borrett), ex: *Banksia marginata*, 25/80.

A. calophylli (Kotinsky)

Syntypes, Fiji: Levuca, 30.x.1899, (A. Koebele), ex: *Calophyllum inophyllum*.

A. ceracroceus Martin

Paratypes, Australia: Queensland, creek-bed, 3 km north of Cape Tribulation, 16.vii.1996, (J.H. Martin 6895), ex: *Neolitsea* spp. (2 slides).

A. citriperdus Quaintance & Baker

Hong Kong: 14.iv.1974, (D.S. Hill), ex: *Citrus* sp. (2 slides).

Hong Kong: Tai Pokau Orchard, 21.ii.1971, ex: Orange, C.I.E. 4663, 11/71 (2 slides).

A. cocois Corbett

Papua New Guinea: New Britain Prov., Gazelle Peninsula, Vunakanau, 18.v.1956, (J.L. Gressit), ex: *Cocos* sp., B.204 (2 slides).

Solomon Island: Kukum, 13.ii.1956, (E.S. Brown), ex: Coconut, C.I.E. 3898Aa/14609.
Solomon Island: Guadalcanal Kukum, 20.viii.1954, (E.S. Brown), ex: Coconut, C.I.E. 3177/13920.

A. delottoi Cohic

Paratype, Kenya: Nairobi, 27.viii.1958, (G. de Lotto), ex: *Chaetachme aristata*, 4/65, B.M. 1965-224.

Kenya: Langata, Nairobi, 7.v.1988, (J.H. Martin 5272), ex: *Pappea capensis*, B.M. 1988-188.

A. esakii Takahashi

Papua New Guinea: Morobe Prov. coast, Buso, 1.xi.1979, (J.H. Martin 2783), ex: *Pometia pinnata* (2 slides).

A. gateri Corbett

West Malaysia: Selangor, Sardang, 7.ii.1985, (J.H. Martin 4417), ex: *Cocos nucifera* (2 slides).

A. gordoniae Takahashi

Hong Kong: slopes of the Peak, above Pok Fu Lam Reservoir Road, 12.xi.1996, (J.H. Martin 6795), ex: *Gordonia axillaris* (2 slides).

A. inceratus Silvestri

Hong Kong: rec'd., v.1990, ex: *Michelia champaca*, C.I.E. A.20977.

A. leptadeniae Cohic

Sudan: Wad Medani, xii.1961, (L.A. Mound), ex: *Leptadenia heterophylla*, 161/162.

A. spiniferus (Quaintance)

Syntypes, Tanzania: ?Dar es Salaam, ?iv.1902, (?A. Zimmermann), ex: ?*Citrus* sp. (2 slides).

India: Coimbatore, 14.iv.1967, (B.V. David), ex: Rose, 10/67, B.M. 1967-280.

Uganda: E. Buganda, Bulemezi, Sabawali, Bugema, 5.vi.1970, (D.N. McNutt), ex: *Citrus sinensis* (2 slides).

Australia: Queensland, Stratford, Cairns, 8.viii.1983, (A.J. Wait), ex: Custard apple (3 slides).

Micronesia: Kosrae I., vi.2001, ex: *Citrus* sp.

8. *Aleurocerus* Bondar

A. cerifer (Sampson & Drews)

Panama: Canal Zone, Cerro Galera, 26.iii.1983, (J.H. Martin 4154), ex: *Tetracera* sp. (2 slides).

Panama: Canal Zone, Barro Colorado I., 13.iii.1983, (J.H. Martin 3986), ex: *Tetracera* sp. (2 slides).

Costa Rica: Guanacaste Province, 60 km south of Liberia, 26.i.1983, (J.H. Martin 3693), ex: *Mastichodendron ?capiri* (2 slides).

A. flavomarginatus Bondar

Costa Rica: N. Heredia Province, 15 km north of Puerto Viejo, 3.ii.1983, (J.H. Martin 3770), ex: *Heliconia* sp. (2 slides).

Ecuador: Mapo Coca, 16.v.1982, (G. Omore), ex: *Elaeis guineensis*, 80/82 (2 slides).

A. musae Russell

Mexico: San Diego, 31.vii.1995, (A. Endo), ex: *Zingiber* sp., 022090 CA, B.M. 1998-131.

Panama: Canal Zone, Pipeline Road, 19.iii.1983, (J.H. Martin 4050), ex: Bignoniaceae (2 slides).

Panama: Canal Zone, Gamboa Hill, 20.iii.1983, (J.H. Martin 4088 B), ex: *Warszewiczia coccinea* (2 slides).

Costa Rica: N. Heredia Province, 10 km north of Puerto Viejo, 2.ii.1983, (J.H. Martin 3740), ex: indet. (2 slides).

A. palmae Russell

Paratypes, Mexico: intercepted at Dallas, Texas, USA, 21.ii.1962, (L. Russell), ex: *Chamaedorea elegans*, ref. 62-15982 (2 slides).

Belize: Cayo, Chiquibul forest, Las Cuevas plots, 3.xi.1994, (J.H. Martin 6438), ex: *Pouteria* sp.

9. *Aleurochiton* Tullgren

A. acerinus Haupt

Czechoslovakia: Radotin, 2.iii.1969, (J. Zahradnik), ex: *Acer campestre* (2 slides).

England: Kent, Polhill Down, near Sevenoaks, 28.v.1983, (W.R. Dolling), ex: *Acer campestre*, 2/83 (2 slides).

Italy: Sicily, (CT), Bronte, 30.x.1981, (C. Rapisarda), ex: *Acer campestre*, 50/84.

Hungary: Szarvas, 21.vi.1990, (V.F. Eastop 19144), ex: *Acer campestre*.

A. aceris (Modeer)

England: Hants, Alice Holt forest, 29.vii.1984, (J.H. Martin 4363), ex: *Acer platanoides* (2 slides).

England: Hants, Alice Holt forest near Alton, 10.vii.1994, (J.H. Martin 6419), ex: *Acer platanoides*.

Netherlands: Bennekom, 7.x.1971, (V.F. Eastop), ex: *Acer platanoides*, 7/72 (3 slides).

A. forbesii (Ashmead)

USA: Illinois, Urbana, 10.x.1970, (L.A. Mound), ex: *Acer saccharinum*, 29/71 (4 slides).

USA: Maryland, Prince Georges Co Beaver Dam Rd, 2.vii.1998, (J.H. Martin 7103), ex: *Acer rubrum* (2 slides).

A. pseudoplatani Visnya

Sicily: (CT/ME), Monti Nebrodi, near Bronte, 10.x.1985, (J.H. Martin 5068), ex: *Acer pseudoplatanus* (2 slides).

Netherlands: Amerongen, 28.ix.1975, (R.M. Bink-Moenen), ex: *Acer pseudoplatanus*, B.M. 1976-56 (2 slides).

10. *Aleuroclava* Singh

A. aucubae (Kuwana)

Japan: Osaka, Taishi, 3.xi.1956, (R. Takahashi), ex: *Paederia* sp., 76/62, B.M. 1962-401 (2 slides).

Japan: Tokyo, 9.x.1963, (H. Takada), ex: *Celtis* sp.

A. bifurcata (Corbett)

Paratype, Malaya: Kajang, 4.v.1927, (G. H. Corbett), ex: *Nephelium lappaceum*, B.M. 1932.511.

West Malaysia: Pahang Prov., Tioman I., Kampung Tekek, 20.iii.1984, (J.H. Martin 4309), ex: indet.

West Malaysia: Pahang Prov., Taman Negara, Tembeling path, 11.iii.1984, (J.H. Martin 4250), ex: indet.

A. citrifolii (Corbett)

India: Punjab, 29.iii.1926, (M.A. Husain), ex: *Citrus* sp., I.B.E. 309.

Pakistan: Peshawar, 31.i.1963, (L.A. Mound), ex: *Murraya exotica*, 10/64.

A. euryae (Kuwana)

Japan: Odaiga - Hara, near Nara, 18.viii.1956, (R. Takahashi), ex: *Trochodendron* sp., 73/62, B.M. 1962-401.

A. ficicola (Takahashi)

Taiwan: Wufeng, 18.iv.1994, (K.C. Chou), ex: *Morus* sp., B.M. 1994.

Taiwan: Wufeng, 9.v.1994, (K.C. Chou), ex: *Morus* sp., B.M. 1994.

A. filamentosa (Corbett)

Brunei: Belait, Peat Swamp forest, East of Seria, 4.x.1992, (J.H. Martin 6122), ex: indet.

A. hikosanensis (Takahashi)

Japan: Mt. Ikoma, near Osaka, 18.ix.1955, (R. Takahashi), ex: Buxa, 78/62, B.M. 1962-401 (2 slides).

A. jasmmini (Takahashi)

Hong Kong: Kowloon, 29.viii.1979, (J.H. Martin 2478), ex: indet. (2 slides).

Singapore: Ayer Rajah, housing blocks, 10.iv.2000, (J.H. Martin 7338), ex: *Jasminum* sp.

Singapore: Ayer Rajah, housing blocks, 10.iv.2000, (J.H. Martin 7334), ex: *Citrus* sp.

Taiwan: Formosa, Tobacco Res. Inst., 12.iv.1967, (Y.C. Chang), ex: *Murraya paniculata*, 9/67, B.M. 1967-258.

A. magnoliae (Takahashi)

Japan: Odaiga - Hara near Nara, 16.viii.1956, (R. Takahashi), ex: Lauraceae, 75/62, B.M. 1962-401.

A. murrayae (Singh)

Pakistan: Peshawar, 31.i.1963, (L.A. Mound), ex: *Murraya exotica*, 9/64, C.I.E. 19214.

Pakistan: Peshawar, 31.i.1963, (L.A. Mound), ex: *Murraya exotica*, 9/64.

A. nigeriae (Mound)

Paratype, Nigeria: Ibadan, 28.ix.1956, ex: Guava, 54/59.

Paratype, Nigeria: Ibadan, Moor Plantation, xii.1960, (M.O. Ezeigwe), ex: *Psidium guajava* (2 slides).

Paratype, Nigeria: Kuseriki, ix.1960, (L.A. Mound), ex: indet. (2 slides).

A. piperis (Takahashi)

Japan: Enoshima, 31.viii.1949, (R. Takahashi), ex: *Piper* sp., 74/62, B.M. 1962-401 (2 slides).

A. psidii (Singh)

Taiwan: Formosa: Tobacco Res. Inst., 3.iv.1967, (Y.C. Chang), ex: *Litchi chinensis*, 4/67, B.M. 1967-186.

Taiwan: Taichung, 10.ix.1967, (Y.C. Chang), ex: *Prunus salicina*, 20/67, B.M. 1967-592.

Taiwan: Taichung, 18.xii.1968, (Y.C. Chang), ex: Grape, 2/69.

A. pulcherrimus (Corbett)

Java: Jakarta, Kemang, 16.xii.1979, (J.H. Martin 2953), ex: indet.

A. similis (Takahashi)

Japan: Asakawa, near Tokyo, 27.vii.1949, (R. Takahashi), ex: *Pieris japonica*, 77/62, B.M. 1962-401 (2 slides).

11. *Aleurocybotus* Quaintance & Baker

A. graminicolus (Quaintance)

Paralectotype, USA: Florida, Lake City, 25.vii.1898, (Prof. Rolfs), Q 6775, ex: Grass, B.M. 1998-131.

A. occiduus Russell

Paratype, USA: California, ix-x.1961, (L.D. Anderson), ex: Grass, 8/65, B.M. 1965-451 (2 slides).

USA: California, El Centro Imp. Co., 5.x.1961, (O.V. Wadgner), ex: *Sorghum* sp., B.M. 1968-368.

12. *Aleuroduplidens* Martin

A. eucalyptifolia Martin

Paratypes, Australia: ACT, Molonglo Gorge (near Queanbeyan, NSW), 8.xii.1996, (J.H. Martin 6852), ex: *Eucalyptus nortonii*, B.M. 1998-50 (2 slides).

Paratypes, Australia: S.A., Belair, National Park, 29.iv.1957, (D.C. Swan), ex: *Eucalyptus* sp. (3 slides).

Australia: WA, Bungle Bungles, Frog Hole Gorge, 8.v.2000, (J.H. Martin 7443), ex: *Eucalyptus saplings*.

A. santali Martin

Paratypes, Australia: Queensland, Mt. Isa, 3.x.1996, (P. De Barro 66), ex: *Santalum lanceolatum* (2 slides).

Paratypes, Australia: Queensland, 25 mi. W. of Dalby, ix.1985, (L.A. Mound), ex: indet., B.M. 1998-51 (2 slides).

A. triangularis Martin

Paratype, Australia: Northern Territory, King's Canyon, 5.ii.1990, (J.H. Martin 5643), ex: *Acacia ?mcdonnelliensis*, B.M. 1998-50 (2 slides).

A. wellsae Martin

Paratypes, Australia: Victoria, Merbein South, near Mildura, 27.i.1997, (A. Wells), ex: *Acacia melvillei*, B.M. 1998-50 (3 slides).

Paratypes, Australia: River Murray, 30 mls, N. of Overland Corner, 10.x.1965, (D.E. Symon), ex: *Acacia wilhelmiana*, 3/66, B.M. 1966-50 (3 slides).

13. *Aleuroglandulus* Bondar

A. magnus Russell

Costa Rica: Guanacaste Province, Santa Rosa Nat. park, 6.iii.1990, (J.M. Cox 63), ex: indet.

A. malangae Russell

Costa Rica: Turrialba, 14.xii.1981, (A. King), ex: *Colocasia* sp., 2/82.

Costa Rica: Turrialba, 640 m., Trop. Agric. Centre, C.A.T.I.E., 2.iii.1983, (J.H. Martin 3954), ex: *Colocasia* sp. (4 slides).

A. striatus Sampson & Drews

Costa Rica: N. Heredia Province, 10 km north of Puerto Viejo, 2.ii.1983, (J.H. Martin 3739), ex: indet. (2 slides).

A. subtilis Bondar

Panama: Darién, Ensenada del Guayabo, sea level - 250 m., 11.i.1983, (J.H. Martin 3588), ex: *Psidium guajava* (2 slides).

Panama: Canal Zone, Gamboa Hill, 20.iii.1983, (J.H. Martin 4095), ex: Bombacaceae.

Panama: Canal Zone, Barro Colorado I., 13.iii.1983, (J.H. Martin 3987), ex: *Inga* ?*pezizifera*.

Panama: Canal Zone, Barro Colorado I., 17.iii.1983, (J.H. Martin 4032), ex: ?*Palicourea guianensis* (2 slides).

14. *Aleuroinanis* Martin

A. myrtacei Martin

Paratypes, Australia: Queensland, strand 2 km north of Cape Tribulation, 15.xii.1996, (J.H. Martin 6883), ex: *Syzygium* sp., B.M. 1998-50.

15. *Aleurolobus* Quaintance & Baker

A. diastematus Bink-Moenen

Sudan: S. Darfur, Ed Da'ain, 3.v.1981, (J.H. Martin 3371), ex: *Guiera senegalensis*.

Sudan: S. Darfur, Jebel Marra, S. W. side, Wadi Golol, 28.iv.1981, (J.H. Martin 3332), ex: *Salix subserrata*.

A. graminicola Bink-Moenen

Paratypes, Chad: Baibokoum, 700 m., 8.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Cymbopogon giganteus*, B.M. 1977-471 (2 slides).

A. hargreavesi Dozier

Syntype, Uganda: Kampala, 16.ii.1928, (H. Hargreaves 0182), ex: *Hyparrhenia* sp.

Nigeria: Enugu, 15.i.1957, (V.F. Eastop 6558), ex: indet.

A. marlatti (Quaintance)

Paralectotypes, Japan: Hokata, 21.v.1901, (Quaintance 10227, via Marlatt), ex: Orange.

Australia: Queensland, rocky coast, 3 km north of Cape Tribulation, 16.xii.1996, (J.H. Martin 6897), ex: ?Euphorbiaceae, B.M. 1998-50.

Australia: Queensland, strand 2 km north of Cape Tribulation, 15.xii.1996, (J.H. Martin 6888), ex: *Clerodendron inerme*, B.M. 1998-50.

Australia: Queensland, Cape Tribulation, 18.xii. 1996, (J.H. Martin 6909), ex: *Aegiceras corniculatus*, B.M. 1998-50 (2 slides).

A. moundi David & Subramaniam

India: Kovilpatti, 31.i.1967, (B.V. David), ex: *Bassia latifolia*, 10/67, B.M. 1967-280/16 (3 slides).

India: Coimbatore, 7.xi.1967, (B.V. David), ex: *Bassia longifolia*, 10/67, B.M. 1967-280/17.

A. onitshae Mound

Paratypes, Nigeria: Onitsha, 13.i.1957, (V.F. Eastop 6553), ex: *Phyllanthus floribundus*, 198/61.

Paratype, Nigeria: Onitsha, 14.i.1957, (V.F. Eastop 6553), ex: *Phyllanthus floribundus*.

Paratypes, Nigeria: Moor Plantation area, i.1961, (L.A. Mound), ex: *Flueggea virosa*.

A. styraci Takahashi

Japan: Tokyo, 16.ix.1950, (R. Takahashi), ex: *Styrax* sp., 80/62, B.M. 1962-401 (2 slides).

A. teucrii Mifsud & Palmeri

Paratypes, Malta: Gozo Island, Xlendi, valley side, 16.iii.1994, (D. Mifsud & G.W. Watson 91), ex: *Teucrium fruticans* (3 slides).

16. *Aleurolonga* Mound

A. cassiae Mound

Paratypes, Nigeria: Ibadan, Moor Plantation, i.1960, (L.A. Mound), ex: *Cassia siamea* (2 slides).

Angola: 3 m N Santa Comba, 8.iii.1972, (D. Hollis), ex: *Brachystegia* sp., 2/74 (2 slides).

17. *Aleuromarginatus* Corbett

A. corbettiaformis Martin

Paratypes, Papua New Guinea: Morobe Province coast, Buso, 10.x.1979, (J.H. Martin 2680), ex: *Desmodium umbellatum* (3 slides).

A. dalbergiae Cohic

Chad: Bebedjia, 395 m, 22.xi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Pterocarpus lucens*, B.M. 1976-56.

A. kallarensis David & Subramaniam

Paratypes, India: Madras, Tambaram, 4.i.1970, (B.V. David), ex: Papilionaceous shrub, B.M. 1972-24.

A. littoralis Martin

Paratypes, Papua New Guinea: Morobe Province coast, Buso, 10.ix.1979, (J.H. Martin 2529), ex: ?*Derris trifoliata* (2 slides).

West Malaysia: Pahang Prov., Pulau Tioman, kg Penuba, beach-top, 19.xi.1992, (J.H. Martin 6219), ex: *Desmodium umbellatum* (2 slides).

A. marginiquus Martin

Paratypes, Australia: Western Australia, Perth, Forrestfield, 23.xi.1997, (P.J. Gullan), ex: *Daviesia horrida*.

A. moundi Martin

Paratypes, Australia: Queensland, 54 km W. of Mitchel, 30.iii.1998, (L.A. Mound 3420), ex: *Acacia ?aneura*, B.M. 1998-51.

A. nemciae Martin

Paratypes, Australia: Western Australia, Darling District, West Talbot Road, 27.ix.1993, (M.D. Crisp 8517), ex: *Nemcia* sp.

A. nigrus Martin

Paratypes, Australia: New South Wales, Braidwood - Nelligen Road, scarp descent, 30.xi.1996, (J.H. Martin 6840), ex: *Oxylobium ilicifolium*, B.M. 1998-50 (2 slides).

Paratypes, Australia: Australian Capital Territory, Molonglo Gorge, 8.xii.1996, (J.H. Martin 6854), ex: *Acacia* sp., B.M. 1998-50.

A. serdangensis Takahashi

Syntype, Malaya: Serdang, 20.i.1945, (R. Takahashi), ex: indet., B.M. 1955-799.

A. tephrosiae Corbett

Chad: Bebedjia, 395 m, 15.viii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Tephrosia linearis*, B.M. 1976-56.

18. *Aleuropapillatus* Regu & David

A. kumariensis Regu & David

Paratype, India: Tamil Nadu, Mondaikadu, 4.viii. 1989, (K. Regu), ex: *Gmelina asiatica*.

19. *Aleuroparadoxus* Quaintance & Baker

A. arctostaphyli Russell

Paratype, USA: Boulder Creek, Calif., 13.x.1891, (V. Bailey), ex: *Arctostaphylos nummularia*, B.M. 1998-131.

A. gardeniae Russell

Paratypes, Mexico: 29.xii.1937, ex: *Gardenia* sp., B.M. 1998-131.

Paratypes, Mexico: 10.xii.1936, (J.M. Singleton), ex: *Gardenia* sp., B.M. 1998-131.

A. ilicicola Russell

Paratype, USA: Alabama, Gulf Shores, 12.i.1944, (L.A. Mayer), ex: *Ilex* sp., B.M. 1998-131.

A. iridescens (Bemis)

?Paralectotype, USA: California, Santa Clara Valley, 1.iv.1901, (F.E. Bemis 7084), ex: *Heteromeles arbutifolia*, B.M. 1998-131.

USA: California, C. Costa Co., Russelmann Park, 21.v.1968, (R.F. Wilkey), ex: *Rhamnus crocea* var. *ilicifolia*, B.M. 1998-131.

A. punctatus Quaintance & Baker

Chile: Santiago, Univ. campus, ix.1995, (V.F. Eastop), ex: indet. (2 slides).

20. *Aleuroplatus* Quaintance & Baker

A. acaciae Bink-Moenen

Paratypes, Egypt: Aswan, Elephantine, 21.i.1977, (F.A. Bink & R.M. Bink-Moenen), ex: *Acacia tortilis*.

Sudan: Khartoum North, 11.iv.1981, (J.H. Martin 3265), ex: *Pithecollobium dulce* (2 slides).

A. berbericolus Quaintance & Baker

USA: California, Trial, Oregon in quarantine Hornbrook, 6.x.1966, (B. Palmer), ex: *Mahonia aquifolium*, 25/66 (4 slides).

A. bossi Takahashi

Kenya: Westermann's Safari Camp, 30 km S. of Voi, 16.v.1988, (J.H. Martin 5312), ex: ?Oleaceae, B.M. 1988-188.

Kenya: ~20 km S. W. of Malindi, inland, 19.v.1988, (J.H. Martin 2348), ex: *Ochna mossambicensis*, B.M. 1988-188.

Sudan: Kordofan, En Nahud - Wad, Banda route, 22.iv.1981, (J.H. Martin 3314), ex: *Boscia senegalensis*.

A. cadabae Priesner & Hosny

Sudan: Kassala Province, Jebel Kassala, 5.iv.1981, (J.H. Martin 3254), ex: *Cadaba rotundifolia* (2 slides).

A. cockerelli (Ihering)

Syntypes, Brazil: S. Paulo Prov., (?V. Ihering), ex: *Baccaris paucifloscula*, B.M. 1920-107.

A. cococolus Quaintance & Baker

Belize: Cayo, Chiquibul forest, San Pastor track, 14.ii.1996, (J.H. Martin 6668), ex: *pimenta dioica*.

Belize: Cayo, Chiquibul forest, Millionario - Grano de Oro, 12.ii.1996, (J.H. Martin 6645), ex: *Guettarda combesii*.

A. dorsipallidus Martin

Paratypes, Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Toraut forest, 11.iii.1985, (J.H. Martin 4639), ex: probably *Taxotrophis ilicifolius* (2 slides).

Paratypes, Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Toraut forest, 20.iii.1985, (J.H. Martin 4709), ex: *Taxotrophis ilicifolius*.

Paratypes, Indonesia: Sulawesi Utara, lakeside, Danau Mooat, Gunung Muajat, 22.iii.1985, (J.H. Martin 4716), ex: ?*Palaquium* sp.

A. epigaeae Russell

Paratypes, Canada: at Boston, 26.v.1939, (W.J. Ehinger), ex: *Epigaea repens*, B.M. 1998-131.

A. evodiae Takahashi

Syntypes, Réunion Island: Plaine des Affouches, v.1957, (J. Bosser), ex: *Evodia* sp., 111/62, B.M. 1962-401.

A. fici Takahashi

Taiwan: Taipei, 1.ii.1987, (C.C. Ker), ex: *Ficus vasculosa*, B.M. 1987-197.

A. ilicis Russell

USA: Texas, in quarantine at Sanders, Arizona, 13.xi.1965, (A.T. Guest), ex: *Ilex* sp., 25/66 (2 slides).

A. incisus Quaintance & Baker

Ceylon: Haputalo, xii.1904, ex: *Ostodes zeylanica*, B.M. 1927-451.

A. pectiniferus Quaintance & Baker

Paratype, India: Tamil Nadu, Burliar, 20.vi.1985, (R.W.A. Jesudasan), ex: *Cinnamomum* sp., B.M. 1987-185.

Paralectotypes, Ceylon: Peradeniya, i.1905, (E.E. Green), ex: *Ficus gibbosa*, B.M. 1927-451.

Australia: Northern Territory, Kakadu Nat. Park, Ubirr rocks, 35 km N. of Jabiru, 22.xii.1996, (J.H. Martin 6956), ex: *Strychnos* sp., B.M. 1998-50.

Hong Kong: Lantau Island, near Tung Chung, 16.x.1990, (J.H. Martin 5779), ex: indet.

Hong Kong: Lantau Island, near Tung Chung, 16.x.1990, (J.H. Martin 5781), ex: *Aguilaria sinensis*.

A. periplocae (Dozier)

Syntype, Belgian Congo: Barumbu, viii.1925, (J. Ghesquière 592), ex: *Periploca nigrescens*.

A. perseaphagus Martin *et al.*

Paratypes, Madeira: Funchal, Lombo da Boa Vista, 3.vii.1993, (via M. Pita C220), ex: *Persea americana* (2 slides).

Paratypes, Madeira: Santa Cruz village, 20.ii.1994, (A.F. Aguiar), ex: *Persea americana*.

A. quercusaquaticae (Quaintance)

USA: Virginia, Vienna, 18.viii.1912, (A.C. Baker 15), ex: Chestnut, B.M. 1924-137.

21. *Aleuropleurocelus* Drews & Sampson

A. nigrans (Bemis)

USA: California, Pasadena, 26.ii.1945, (R.H. Smith), ex: *Rhamnus californica*, B.M. 1998-131.

22. *Aleuropteridis* Mound

A. eastopi Mound

Holotype, Ghana: Tafo, 12.v.1957, (V.F. Eastop 6982), ex: Fern, 109/61.

Paratypes, Ghana: Tafo, 12.v.1957, (V.F. Eastop 6982), ex: Fern, 109/61 (3 slides).

A. filicicola (Newstead)

Syntypes, England: Surrey, Kew Gardens, ?1890, (J.W. Douglas 1205), ex: *Pteris togoensis* & *Cyclosorus dentatus*, 97/61, B.M. 1904-120 (2 slides).

Syntypes, Tanzania (German East Africa): Sigithal near Amani, 4.viii.1902, (A. Zimmermann 18), ex: Fern, 49/62 (4 slides).

A. hargreavesi Mound

Syntypes, Sierra Leone: Freetown, Mokinma, 9.ix.1924, (E. Hargreaves), ex: Bush fern, 108/61.

A. jamesi Mound

Syntypes, Nigeria: Ibadan, Moor Plantation, Ajibi V., 2.x.1959, (E.A. James), ex: *Pteris togoensis*, C.I.E. 5387, 21/60, B.M. 1960-101 (4 slides).

Syntypes, Nigeria: Ibadan, Moor Plantation, Ajibi V., 29.xii.1960, (E.A. James), ex: *Pteris togoensis*.

23. *Aleuroputeus* Corbett

A. perseae Corbett

West Malaysia: Pahang Prov., Taman Negara, Kuala Tahan, 11.iii.1984, (J.H. Martin 4260), ex: indet. small tree (2 slides).

Indonesia: Java, Jakarta, Kemang, 16.xii.1979, (J.H. Martin 2952), ex: *Psidium guajava*.

24. *Aleurothrixus* Quaintance & Baker

A. aepim (Göldi)

Brazil: 23.v.1978, (M. Samways "A"), ex: *Manihot* sp., 8/78.

Brazil: 23.v.1978, (M. Samways "B"), ex: *Manihot* sp., 8/78.

A. antidesmae Takahashi

Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Toraut forest, near base camp, 6.v.1985, (J.H. Martin 4926), ex: indet.

Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Toraut forest, near base camp, 30.iii.1985, (J.H. Martin 4767), ex: ?Myrtaceae.

Hawaiian Islands: Maui, Wailuku, 3.iv.1999, (J.H. Martin 7201), ex: *calophyllum inophyllum*.

A. bondari Costa Lima

Brazil: Vicoso, (H.L. Parker 953-16), ex: *Triumfetta semitriloba*, 44-27574, B.M. 1998-131.

A. floccosus (Maskell)

Singapore: Napier/Cluny Rds, Botanic Gardens, 11.iv.2000, (J.H. Martin 7364), ex: *Citrus retusa* (2 slides).

Singapore: Napier/Cluny Rds, Botanic Gardens, 11.iv.2000, (J.H. Martin 7363), ex: *Noringi crenulata*.

A. interrogationis (Bemis)

USA: California, Arcadia, 3.xi.1970, (H.G. Walker), ex: *Ceanothus cyaneus*, B.M. 1998-131.

USA: California, Riverside, 11.v.1961, (L.A. Mound), ex: *Ceanothus* sp., 91/61, B.M. 1961-369 (2 slides).

USA: California, Riverside, 13.ii.1957, (M. Johnson), ex: *Ceanothus* sp.

A. lucumai Costa Lima

Brazil: Vicoso, 25.v.1944, (H.L. Parker 953-26), ex: *Citrus* sp., 44-27574, B.M. 1998-131.

A. myrtacei Bondar

Belize: Cayo, Chiquibul forest, Las Cuevas plots, 17.xi.1994, (J.H. Martin 6491), ex: *Pimenta dioica* (2 slides).

A. porteri Quaintance & Baker

Syntypes, Chile: Villa del Mar, 1.iv.1899, (D.G. Fairchild) ex: Solanaceae, B.M. 1998-131.

A. proximans Bondar

?Syntypes, Brazil: xii.1922, (G. Bondar 653), ex: Lauraceae, B.M. 1998-131 (2 slides).

A. smilaceti Takahashi

Taiwan: Kenting, 4.iv.1986, (C.C. Ker.), ex: *Gardenia* sp., B.M. 1987-197.

25. *Aleurotithius* Quaintance & Baker

A. timberlakei Quaintance & Baker

USA: California, San Diego, Encanto, 10.iv.1968, (R. Thomas), ex: *Eriodictyon crassifolium*, B.M. 1968-368 (2 slides).

USA: California, Pala, 1.xi.1950, (R.A. Flock via L.A. Mound), ex: *Eriodictyon* sp., 87/61, B.M. 1961-369.

26. *Aleurotrachelus* Quaintance & Baker

A. alpinus Takahashi

Taiwan: 20.x.1989, (C.C. Ko), ex: *Polygonum* sp.

A. ampullatus Bink-Moenen

Paratypes, Chad: Boro, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Burkea africana*, B.M. 1977-471.

A. asparagi (Lewis)

Syntypes, South Africa: Natal, iv.1893, (R.T. Lewis), ex: *Asparagus* sp., B.M. 1893-75 (2 slides).

A. atratus Hempel

Topotype, Brazil: Bahia, 7.xii.1923, (G. Bondar), ex: *Cocos nucifera*, B.M. 1924-44.

Tenerife: Canary Islands, Santa Cruz de T., 22.v.1997, (J.H. Martin 7072 & E.H. Suarez), ex: *Arecastrum* sp. (3 slides).

A. camelliae (Kuwana)

Japan: Hyogo, 15.iv.1956, (R. Takahashi), ex: *Camellia japonica*, 83/62, B.M. 1962-401 (2 slides).

A. duplicatus Bink-Moenen

Paratypes, Chad: Bebedjia, 28.xi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Mitragyna inermis*, B.M. 1977-471.

A. fumipennis (Hempel)

Brazil: Mato Grosso, 25.iii.1968, (O.W. Richards), ex: Grass, B.M. 1968-260 (2 slides).

A. limbatus (Maskell)

Australia: Victoria, nr McKenzie Falls, 23.viii.1965, (D.E. Symon), ex: *Styphelia adscendens*.

Australia: Victoria, nr McKenzie Falls, 23.viii.1965, (D.E. Symon), ex: *Styphelia adscendens*, 3/65, B.M. 1966-50 (2 slides).

A. longispinus Corbett

Syntypes, Ceylon: Telloola, xi.1905, (E.E. Green), ex: indet. 86/61, B.M. 1927-451 (2 slides).

A. minutus Takahashi

Singapore: Botanic Garden, 15.ix.1980, (D.H. Murphy), ex: Annonaceae, 10/85.

A. multipapillus Singh

Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Toraut forest, near base camp, 14.iii.1985, (J.H. Martin 4660), ex: Bamboo.

A. obscurus Bink-Moenen

Paratypes, Chad: Boro, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Daniella oliveri*, B.M. 1977-471.

A. tarennae Bink-Moenen

Paratype, Chad: Timberi, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Tarenna* sp., B.M. 1977-471.

A. tracheifer (Quaintance)

Belize: Cayo, Chiquibul forest, Las Cuevas, Monkey Tail trail, 17.ii.1996, (J.H. Martin 6685), ex: *Strychnos* sp.

Belize: Cayo, Chiquibul forest, Las Cuevas, Monkey Tail trail, 5.iii.1996, (J.H. Martin 6752), ex: woody recumbent (2 slides).

Panama: Canal Zone, Barro Colorado I., 1.i.1983, (J.H. Martin 3478), ex: *Mouriri myrtilloides*.

A. trachoides (Back)

Tahiti; Papeete, 0-50 m., 8.iii.1979, (N.L.H. Krauss), ex: Vine, ?Convolvulaceae (2 slides).

Costa Rica: Guanacaste Province, 100 km S. E. of Liberia, 29.i.1983, (J.H. Martin 3734), ex: Simaroubaceae.

27. *Aleurotuba* Tremblay & Iaccarino

A. jelinekii (Frauenfeld)

England: Mddx., Staines station, 31.x.1976, (J.H. Martin 1660), ex: *Viburnum ?tinus*.

England: Notts., Radcliffe -on- Trent, 5.iv.1986, (J.H. Martin 5069), ex: *Viburnum tinus* (2 slides).

England: London, Battersea Park, 7.xi.1976, (J.H. Martin 1674), ex: *Viburnum ?tinus*.

Morocco: 45 km S. of Oujda, Col de Jerada, 23.iv.1992, (J.H. Martin 6060), ex: *Viburnum tinus*, B.M. 1992-73.

Morocco: Western Rif, 20 km S.W. of Chefchaouene, Ouezzane Road, 29.iv.1992, (J.H. Martin 6085), ex: *Arbutus unedo*, B.M. 1992-73.

France: Corsica, Gorges, de la Restonica, 23.v.1990, (J.H. Martin 5716), ex: *Arbutus unedo*.

28. *Aleurotulus* Quaintance & Baker

A. anthuricola Nakahara

Paratypes, Colombia: at Miami, 28.iii.1984, (J. Russo), ex: *Anthurium* sp.

A. arundinacea Singh

Papua New Guinea: Morobe Prov. coast, Buso, 21.ix.1979, (J.H. Martin 2598), ex: Bamboo.

A. nephrolepidis (Quaintance)

Type series, England: Surrey, Kew Gardens, 17.ix.1937, (C.B. Williams), ex: Ferns, 166/62, B.M. 1962-643 (2 slides).

Azores: São Miguel, road to Sete Cidades, 27.ix.1998, (A. Polaszek & E.M. Hernandez-Suares), ex: *Blechnum spicant* (2 slides).

Brazil: S.P., ?Campinas, rec'd., iii.1982, (A. Lourencao), ex: *Pteris* sp., 3/82.

A. pteridophytae Martin

Paratypes, Costa Rica: North Heredia Province, La Selva Biological Reserve, xi.1992, (L.A. Mound), ex: *Selaginella eurymota* (4 Slides).

Venezuela: Parque Nacional, El Cumbre, 2.xii.1938, (A.H.G. Alston), ex: *Anemia phyllitidis*, 162/60 (2 slides).

29. *Aleuroviggianus* Iaccarino

A. adanaensis Bink-Moenen

Paratypes, Turkey: Adana, 19.iv.1987, (H. Elekcioglu), ex: *Quercus coccifera*, 22/87 (4 slides).

Syria: Idleeb, 15.v.1995, (Badar Munir), ex: *Quercus calliprinos*, B.M. 1995-187.

A. adrianae Iaccarino

Italy: nr Naples, x.1981, (F.M. Iaccarino), ex: *Quercus ilex* group, 281/81.

Morocco: Middle Atlas, Ifrane, 25.iv.1992, (J.H. Martin 6067), ex: *Quercus* sp., B.M. 1992-73.

Morocco: Western Rif, 5 km S. of Chefchaouene, roadside, 19.iv.1992, (J.H. Martin 6043), ex: *Quercus ?suber*, B.M. 1992-73.

Costa Rica: Cargèse, 31.x.1998, (J.H. Martin 7189), ex: *Quercus ilex* (2 slides).

A. graecus Bink-Moenen

Holotype, Greece: Corfu, Kerkira Kariotikon, 13.v.1980, (F.A. Bink & R.M. Bink-Moenen), ex: *Quercus coccifera*.

A. halperini Bink-Moenen

Paratypes, Israel: Golan Heights, 5.xii.1967, (J. Halperin), ex: *Quercus calliprinos*.

Paratype, Israel: Tivon, 20.xii.1958, (M. Sternlicht 3811), ex: *Quercus calliprinos*, C.I.E. 5001/16428.

Turkey: Içel, 30.iii.1987, (H. Elekcioglu), ex: *Quercus coccifera* (2 slides).

A. polymorphus Bink-Moenen

Paratypes, Spain: Colmenar Viejo, 12.v.1978, (F.A. Bink & R.M. Bink-Moenen), ex: *Quercus rotundifolia* (2 slides).

Paratype, France: Pyrenées Or., N.W. of Ille-s-Têt, 11.v.1987, (F.A. Bink & R.M. Bink-Moenen), ex: *Quercus ilex*.

Morocco: Western Rif, 5 km S. of Chefchaouene, roadside, 19.iv.1992, (J.H. Martin 6043), ex: *Quercus ?suber*, B.M. 1992-73 (2 slides).

Morocco: 45 km S. of Oujda, Col de Jerada, 23.iv.1992, (J.H. Martin 6064), ex: *Quercus coccifera*, B.M. 1992-73.

A. zonalus Bink-Moenen

Paratype, Greece: Crete, Montes Psiloriti, 5.vii.?1942, (L.A. Mound & R.M. Bink), ex: *Quercus coccifera*.

Paratypes, Greece: Crete, Omalus, 8.v.1979, (F.A. Bink & R.M. Bink-Moenen), ex: *Quercus coccifera*.

30. *Aleyrodes* Latreille

A. amnicola Bemis

USA: California, Vista, San Diego, 14.xi.1966, (Waldrip & Kruz), ex: indet., 8/72 (3 slides).

A. elevatus Silvestri

Spain: Almeria Province, gorge N of Berja, Road A337, 9.x.1998, (J.H. Martin 7162), ex: *Ficus carica* (2 slides).

France: Corsica, Ile Rousse, 4.vii.1980, (J.H. Martin 3028), ex: *Ficus ?carica*, 47/80.

A. loniceræ Walker

England: London, New Cross, 21.x.1968, (J.M. Palmer), ex: *Lonicera serotina*, 5/69 (3 slides).

England: Notts., Radcliffe -on- Trent, 4.x.1980, (J.H. Martin 3108), ex: *Lonicera periclymenum*, 50/80.

England: Notts., Radcliffe -on- Trent, 18.ix.1982, (J.H. Martin 3448), ex: *Mentha* sp., 57/82.

Austria: Ossiach, 14.viii.1966, (V.F. Eastop), ex: *Aegopodium podograria*.

A. proletella (Linnaeus)

Angola: C.E. Cavaco, 7.xii.1970, (P. Carvalho), ex: *Brassica oleracea*, 17616/1571 (2 slides).

Austria: Ossiach, 6.viii.1966, (V.F. Eastop), ex: *Chelidonium* sp. (2 slides).

England: Kent, Dunton Green, 30.ix.1969, (W.R. Dolling), ex: *Aegopodium podograria*, 3/80 (2 slides).

Tenerife: Canary Islands, La Matanza de Acentejo, 20.v.1997, (J.H. Martin 7058), ex: *Lactuca serriola*.

Malta: Gozo Island, Xlendi, gariugue habitat, valley side, 16.iii.1994, (G.W. Watson 90), ex: *Reichardia picroides*.

A. pruinosus Bemis

Type material: USA: Colorado, Boulder, pre 1912, (T.D.A. Kockerell), ex: indet., 96/61, B.M. 1912-49.

USA: California, San Diego, 24.x.1966, (J. Kenyon), ex: *Heteromeles arbutifolia*, 8/12 (2 slides).

USA: California, Riverside, 11.ii.1957, (M. Frost), ex: *Photinia arbutifolia*.

A. singularis Danzig

Paratype, USSR: Georgia, Marneul District, 1962, (T. Chavchanidze), ex: *Euphorbia* sp., 4/66, B.M. 1966-67 (4 slides).

31. *Aleyrodiella* Danzig

A. lamellifera Danzig

Paratypes, USSR: Vladivostock, Okeanskaya, 31.viii.1961, (E.M. Danzig), ex: *Ulmus laciniata*.

32. *Apobemisia* Takahashi

A. kuwanai (Takahashi)

Japan: Amani, Oshima, 6.iv.1960, (R. Takahashi), ex: *Ficus pumila*, 87/62, B.M. 1962-401 (3 slides).

33. *Arachnaleyrodes* Bink-Moenen

A. insignis Bink-Moenen

Paratypes, Chad: Bebedjia, 395 m, 27.vi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Combretum hypopilinum*, B.M. 1976-56 (2 slides).

Paratypes, Chad: Baïbokoum, 700 m, 9.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Combretum glutinosum*, B.M. 1976-56.

34. *Asialeyrides* Corbett

A. lushanensis Ko

Paratypes, Taiwan: Lushanwenchuan, 6.ix.1986, (C.C. Ko), ex: *Syzygium buxifolium*, B.M. 1987-197 (2 slides).

A. maesae (Takahashi)

Taiwan: Hapan, 11.i.1987, (C.C. Ko), ex: indet., B.M. 1987-197.

35. *Asterobemisia* Trehan

A. carpini (Koch)

England: Herts., near Barford, 18.iv.1936, (C.B. Williams), ex: Hornbeam, 133/60, B.M. 1960-483 (2 slides).

England: Herts., Harpenden, Bluebell Wood, 16.x.1935, (C.B. Williams), ex: Hornbeam, 94/61, B.M. 1969-483 (2 slides).

A. dentata Danzig

Turkmenistan: Kapem, Dag, 10.x.1988, ex: *Rhamnus* sp.

A. obenbergeri (Zahradnik)

Paratype, Czechoslovakia: Mohelno, Moravia Occid., 29.ix.1954, ex: *Thymus* sp.

Paratypes, Czechoslovakia: Mohelno, 30.viii.1957, (J. Zahradnik), ex: *Thymus* sp., B.M. 1969-370.

36. *Asterochiton* Maskell

A. cordiae David & Subramaniam

Paratype, India: Madras, 19.vii.1971, (B.V. David), ex: *Cordia myxa*, B.M. 1972-24.

A. pittospori Dumbleton

New Zealand: Auckland, i.1979, (L.A. Mound 1346), ex: *Pittosporum eugenioides* (2 slides).

37. *Bellitudo* Russell

B. jamaicae Russell

Paratype, Jamaica: Holly Mount, 25-27.v.1904, (W.R. Maxon), ex: *Coccoloba longiflora*, B.M. 1998-131.

West Indies: Jamaica, La Rozelle, Morant Bay, 16.iv.1969, (K. Heinze 12), ex: *Coccoloba unifera*.

Jamaica: La Rozelle, 23.x.1969, (K. Heinze), ex: *Coccoloba unifera*, B.M. 1998-131.

B. hispaniolae Russell

Haiti: Port-au-Prince, 28.v.1930, (H.L. Dozier), ex: *Coccoloba unifera*, B.M. 1998-131.

38. *Bemisaleyrodes* Cohic

B. brideliae Bink-Moenen

Paratypes, Chad: Boro, 450 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Bridelia ferruginea*, B.M. 1976-56 (3 slides).

B. pauliani Cohic

Paratype, Nigeria: Ibadan, 29.x.1960, (F.A. Squire), ex: *Ficus* sp.

Paratype, Cameroon: Bamenda, 2.xi.1957, (V.F. Eastop), ex: *Ficus* sp.

39. *Bemisia* Quaintance & Baker

B. afer (Priesner & Hosny)

Israel: Slopes of Wadi Keren, after Zuriel, Upper Galilee, 15.x.1968, (D. Gerling 132), ex: *Smilax aspersa*, B.M. 1969-220.

Israel: Yoknean Junction, nr Zikhron Yaakov., 15.x.1968, (D. Gerling 013), ex: *Morus* sp., B.M. 1969-220.

Pakistan: Lyallpur, 24.viii.1966, (J.I. Qureshi 102), ex: *Morus alba*, 21/66 (2 slides).

Pakistan: Lyallpur, 25.viii.1966, (J.I. Qureshi 104), ex: *Dalbergia sissoo*, 21/66.

Iraq: Mosul University, ?1977, ex: *Bauhinia* sp., No. 35, C.I.E. A.10124.

Iran: Fars, Shiraz, 28.x.1990, (M. Zarrabi 8), ex: *Rosa canina*.

B. antennata Gameel

Chad: Bebedjia, 395 m, 4.vii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Cordia africana*, B.M. 1976-56.

Sudan: Wad Medani, x.1961, (L.A. Mound), ex: *Ficus sycamora*.

Sudan: Wad Medani, xi.1961, (L.A. Mound), ex: *Ficus sycamora*.

Sudan: Wad Medani, i.1962, (L.A. Mound), ex: *Ficus sycamora*, B.M. 1963-777.

Sudan: Wad Medani, v.1962, (L.A. Mound), ex: *Ficus sycamora*.

Sudan: Wad Medani, 1962, (L.A. Mound), ex: *Ficus sycamora*.

B. berbericola (Cockerell)

USA: California, Riverside, 11.v.1961, (L.A. Mound), ex: *Photinia arbutifolia*, 86/61, B.M. 1961-369.

USA: California, Riverside, 11.v.1961, (L.A. Mound), ex: *Ceanothus arbutifolia*, 93/61, B.M. 1961-369.

USA: California, Riverside, 11.v.1961, (L.A. Mound), ex: *Ceanothus* sp., 91/61, B.M. 1961-369 (2 slides).

B. combreticula Bink-Moenen

Holotype, Chad: Bebedjia, 395 m, 27.vii.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Anogeissus leiocarpus*.

B. formosana Takahashi

Taiwan: Chihpenwenchuan, 26.ix.1989, (I.C. Hsu), ex: *Setaria palmifolia*, B.M. 1994 (3 slides).

B. giffardi (Kotinsky)

Australia: Queensland, Koah, 19.iii.1959, (G.W. Saunders), ex: *Citrus* sp., C.I.E. 5435 1686, 40/60, B.M. 1960-240.

Australia: Adelaide, Highgate, iv.1958, (H.M. Brookes), ex: Orange, 88/58, B.M. 1965-517.

Hong Kong: NT, Sek Kong, 1.iii.1990, (J.H. Martin 5681), ex: *Citrus* sp.

Nepal: Kathmandu, 15.iv.1965, (K.C. Sharma), ex: *Citrus* sp., C.I.E. 8855.A266, B.M. 196.

Tahiti: Fada, ix.1976, (M. Monnot), ex: *Citrus aurantiifolia*, C.I.E. A.10644, 10/82.

B. gigantea Martin

Australia: Queensland, Happy Valley, 3 km W. of Babinda, 3.v.1998, (C. Burwell & C. Rodriguez), ex: *Elaeocarpus angustifolius*, B.M. 1998-51 (3 slides).

B. guieriae Bink-Moenen

Sudan: Kordofan, 100 km W. of Umm, Ruwaba, EL Obeid Road, 18.iv.1981, (J.H. Martin 3296), ex: *Guiera senegalensis* (2 slides).

B. hirta Bink-Moenen

Sudan: N. Darfur, EL Fasher, 25.iv.1981, (J.H. Martin 3329), ex: *Boscia senegalensis*.

Sudan: Kassala Prov., Jebel Kassala, 6.iv.1981, (J.H. Martin 3258), ex: *Boscia senegalensis*.

Sudan: Kordofan, 20 km W. of EL Obeid, 18.iv.1981, (J.H. Martin 3300), ex: *Boscia senegalensis*.

B. lauracea Martin et al.

Paratypes, Madeira: Faja da Nogueira, 28.iii.1995, (J.H. Martin & A.F. Aguiar, JHM 6553), ex: *Ocotea foetens* (2 slides).

Paratypes, Madeira: cliffs above Seixal, 200–300 m, 29.iii.1995, (J.H. Martin & A.F. Aguiar, JHM 6556), ex: *Laurus azorica* (2 slides).

Paratype, Madeira: Fanal, 1150 m. N. E. of Rabacal, 30.iii.1995, (J.H. Martin & A.F. Aguiar, JHM 6574), ex: *Laurus azorica*.

Paratypes, Madeira: Fanal, 1150 m. N. E. of Rabacal, 30.iii.1995, (J.H. Martin & A.F. Aguiar, JHM 6575), ex: *Ocotea foetens*.

B. mesasiatica Danzig

Paratype, Tadjikistan, 15.vi.1964, (E. Danzig), ex: *Exochorda alberti*.

B. pongamiae Takahashi

West Malaysia: Pahang Prov., Tioman I., Kampung Tekek, 16.iii.1984, (J.H. Martin 4272), ex: ?Leguminosae (2 slides).

Philippine Islands: Palawan, st. paul N. P., Sabang H. Q., 24.i.1988, (J.H. Martin 5163), ex: strand legume vine, B.M. 1988-2.

Philippine Islands: Palawan, Puerto Princesa, White Beach, 27.i.1988, (J.H. Martin 5177), ex: strand Leguminosae shrub, B.M. 1988-2.

Taiwan: Taipei, 6.ix.1985, (C.C. Ko), ex: *Pongamia pinnata*.

B. silvatica Danzig

Paratypes, USSR: Alekseyevka, Lazarevskaya, District, 11.viii.1960, (E.M. Danzig), ex: *Rhododendron flavum*.

B. subdecipiens Martin

Paratypes, Australia: Kangaroo Island, Flinders Chase N. P., 17.ii.1990, (J.H. Martin 5652), ex: *Melaleuca* sp., B.M. 1998-50 (2 slides).

B. tabaci (Gennadius)

Jordan: 1977, (K. Makkouk 21), ex: *Solanum lycopersicon*, 105/84 (2 slides).

Kuwait: Al-Jabriyah, 10.i.1977, (W. Al-Houty), ex: *Solanum melongena*, 6/77 (2 slides).

Greece: N. Evia, Isteia, viii.1982, (A. Kourmadas), ex: *Hibiscus esculentus*, 34/82.

Iran: Gorgan, Ghalandar Mahaleh, 1.xi.1986, ex: Cotton, C.I.E. A18482 (2 slides).

Iran: Gorgan, ?.xi.1986, ex: Cotton, C.I.E. A18482.

Iran: Khuzistan, Ramin, 2.iii.1978, (V.F. Eastop 15893), ex: Euphorbiaceae, 129/81.

Iran: Khuzistan, Ahvaz, 6.ii.1978, (V.F. Eastop 15787), ex: *Malva* sp. or ?*Hibiscus* sp., 120/81.

Iraq: Haidei, 21.vi.1961, ex: Cotton, C.I.E. 6384-18046, B.M. 1962-3 (2 slides).

B. tuberculata Bondar

Paraguay: iv.1995, (Romero), ex: *Manihot esculenta*, P95-042 (2 slides).

Colombia: Ciat, Valle Palmira, viii.1997, (M. Pilar Hernandez), ex: *Yuca* sp.

40. *Bemisiella* Danzig

B. artemisiae Danzig

Paratypes, Russia: Voronezh, 3.viii.1999, (J. Yavrilov) ex: *Artemisia vulgaris*, BMNH (E) 2000-64.

41. *Brazzaleyrodes* Cohic

B. eriococciformis Cohic

Belgian Congo: Yang-Steui, 13.xii.1936, (J. Ghesquière 3472), ex: *Cynometra hankei*.

42. *Bulgarialeurodes* Corbett

B. cotesii (Maskell)

Syntypes, Bulgaria: 30.viii.1930, (Dr P. Tchorbadjiev), ex: *Rosa damascena*, B.M. 1935-284.

Iran: Fars, Shiraz, 28.x.1990, (M. Zarrabi 8), ex: *Rosa canina*.

43. *Calluneyrodes* Zahradnik

C. callunae (Ossiannilsson)

?Syntype, Sweden: Södermanland, Nacka Erstavik, 27.x.1946, (Ossiannilsson).

44. *Chitonaleyrodes* Martin

C. canberrensis Martin

Paratype, Australia: Canberra, A.C.T., viii.1968, (L.A. Mound), ex: *Callistemon* sp., 34/71 (4 slides).

Paratypes, Australia: Canberra, A.C.T., i.1968, (L.A. Mound), ex: *Callistemon* sp., 33/71 (2 slides).

45. *Cockerelliella* Sundararaj & David

C. curcuma (Corbett)

West Malaysia: Pahang Prov., Taman Negara, Kuala Tahan, 4.iii.1984, (J.H. Martin 4220), ex: Musaceae (2 slides).

West Malaysia: Pahang Prov., Taman Negara, Kuala Tahan, 2.iii.1984, (J.H. Martin 4191), ex: Zingiberaceae.

West Malaysia: Cameron Highlands, nr Ringlet, ~1000 m, 14.ii.1985, (J.H. Martin 4489), ex: Zingiberaceae.

West Malaysia: Genting Highlands, ~3000', 10.ii.1985, (J.H. Martin 4445), ex: Zingiberaceae.

C. psidii (Corbett)

Singapore: Bukit Timah, Forest Reserve, 13.v.2000, (J.H. Martin 7465), ex: indet.

Papua New Guinea: Morobe Prov. coast, Buso, 9.x.1979, (J.H. Martin 2681), ex: *Premna* sp. (2 slides).

Papua New Guinea: Morobe Prov. coast, Buso, 3.x.1979, (J.H. Martin 2655), ex: *Myrtella* sp.

Papua New Guinea: Morobe Prov. coast, Buso, 12.x.1979, (J.H. Martin 2687), ex: *Anisoptera thurifera* (2 slides).

Papua New Guinea: Morobe Prov. coast, Lasanga Island, 7.xi.1979, (J.H. Martin 2817), ex: indet. tree (2 slides).

46. *Cohicaleyrodes* Bink-Moenen

C. alternans (Cohic)

?Paratype, Congo: Brazzaville, 17.iv.1964, (F. Cohic), ex: Papilionaceae, B.M. 1965-305.
Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 13.x.1989, (L.D.C. Fishpool 17), ex: *Terminalia ivorensis*.

C. crossopterygis Bink-Moenen

Paratypes, Chad: Boro, 453 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Crossopteryx febrifuga*.

C. obscura Bink-Moenen

Holotype, Chad: Timberi, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Tarenna* sp.

C. platysepali (Cohic)

Paratype, Congo: Brazzaville, 16.iv.1965, (F. Cohic), ex: *Platysepalum vanderystii*, B.M. 1965-305.

C. quadrilongispinae Bink-Moenen

Paratypes, Chad: Bebedjia, 395 m, 23.vii.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Cissus* sp.

Paratype, Chad: Bebedjia, 395 m, 25.x.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Cissus* sp.

Paratypes, Chad: Doba, 21.vi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Paullinia pinnata*.

C. recurvispina (Cohic)

Syntype, Congo: Brazzaville, 21.iii.1964, (F. Cohic), ex: *Tetracera alnifolia*, B.M. 1965-305.

Chad: Doba, 27.xii.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*, B.M. 1973-226.

47. *Combesaleyrodes* Cohic

C. bouqueti Cohic

Syntype, Congo: Brazzaville, 25.ii.1965, (F. Cohic), ex: *Collettoecema dewevrei*, B.M. 1965-305.

C. tauffliebi Cohic

Syntype, Congo: Brazzaville, 25.ii.1965, (F. Cohic), ex: *Urophyllum hirtellum*, B.M. 1965-305.

48. *Corbettia* Dozier

C. baphiae Russell

Kenya: Matuga, 4.ix.1982, (I. Robertson 3361), ex: *Cassia* sp., 54/82 (2 slides).

Malawi: Bunda, Lilongwe, 17.xi.1995, (Y. Tembo), ex: *Sesbania sesban*.

C. bauhiniiae Cohic

Sudan: Bahr el Ghazal Prov., N. of Aweil, 9.v.1981, (J.H. Martin 3387), ex: *Piliostigma* sp. (2 slides).

Sudan: S. Darfur, Kas-Nyala Road, 1.v.1981, (J.H. Martin 3368), ex: *Bauhinia* | *Piliostigma* sp. (2 slides).

C. graminis Mound

Paratypes, Nigeria: Onitsha, 13.i.1957, (V.F. Eastop 6544), ex: Gramineae (3 slides).

C. isoberliniae Bink-Moenen

Paratypes, Chad: Boro, 453 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Isoberlinia doka*, B.M. 1976-56 (2 slides).

C. lonchocarpi Bink-Moenen

Paratypes, Chad: Boro, 450 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Lonchocarpus laxiflorus*, B.M. 1976-56 (2 slides).

C. milletticola Dozier

Paralectotype, Belgian Congo: Sankuru, Kole, 22.i.1928, (J. Ghesquière), ex: *Millettia versicolor*, B.M. 1933.

Nigeria: Ibadan, Moor Plantation, 10.v.1956, (V.F. Eastop 5929), ex: *Lonchocarpus sericeus*.

Nigeria: Olokomeji, (V.F. Eastop 5942), ex: *Mucuna* sp.

Chad: Bebedjia, 395 m, 30.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Swartzia madagascariensis*, B.M. 1976-65 (2 slides).

C. tamarindi Takahashi

Chad: Bebedjia, 395 m, 19.ix.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Tamarindus indicus*, B.M. 1976-56 (2 slides).

49. *Crenidorsum* Russell

C. aroidephagus Martin & Aguiar

Paratypes, Costa Rica: San José, 22.i.1983, (J.H. Martin 3648), ex: ?*Philodendron* (2 slides).

Paratypes, Madeira: Funchal, 16.iii.1999, (A. Aguiar C873), ex: *Epipremnum aureus* (2 slides).

Paratypes, Germany: Berlin, Botanic Gardens, iii.2001, (P. Baufeld), ex: *Philodendron gloriosum* (2 slides).

Paratype, USA: Hawaii, Foster Botanical Gardens, 10.iv.2001, (B. Kumashiro & J. Dooley), ex: Aroid. B.M.N.H. (E) 2001-140.

C. celebes Martin

Paratypes, Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Clark's camp area, 18.iv.1985, (J.H. Martin 4881), ex: ?Proteaceae (cf. *Helicia* sp.) (4 slides).

Paratype, Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Clark's camp area, 12.v.1985, (J.H. Martin 4940), ex: ?Sapotaceae.

Paratypes, Indonesia: Sulawesi Utara, lakeside, Danau Mooat, Gunung Muajat, 22.iii.1985, (J.H. Martin 4720), ex: *Persea americana*.

Paratypes, Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Clark's camp area, 15.iv.1985, (J.H. Martin 4846), ex: *Weinmannia* sp.

C. commune Russell

Paratype, Bahamas: Harbour Island, 18.ii.-4.iii.1907, (E.G. Britton), ex: *Coccoloba laurifolia*, B.M. 1998-131.

Paratypes, Navassa Island: 6.i.1930, (H. Rehder), ex: *Coccoloba laurifolia*, B.M. 1998-131.

C. lasangensis Martin

Paratypes, Papua New Guinea: Morobe Province coast, Lasanga Island, 18.ix.1979, (J.H. Martin 2586), ex: *Musa* sp. (2 slides).

C. magnisetae Russell

Paratype, Haiti: Massif, Port Margot, (E.L. Ekman), ex: *Coccoloba retusa*, B.M. 1998-131.
Haiti: Morne-d-Cabrits, 19.xii.1930, (H.L. Dozier), ex: *Bunchosia* sp., B.M. 1998-131.

C. malpighiae Russell

Puerto Rico: Guajataca, Gorge, 13.iii.1952, (Martorell & L.M. Russell), ex: *Torrubia fragans*, B.M. 1998-131.

C. micheliae (Takahashi)

Hong Kong: NT, Tai Po Kau, headland forest, 21.xi.1999, (J.H. Martin 7247), ex: *Embelia laeta* (2 slides).

C. millennium Martin

Holotype, Australia: Northern Territory, Kakadu Nat. Park, Gubara Walk, Burdulba Creek, 20 km S. of Jabiru, 22.xii.1996, (J.H. Martin 6954), ex: *Xanthostemon eucalyptoides*.

C. morobensis Martin

Paratypes, Papua New Guinea: Morobe Province coast, Buso, 3.x.1979, (J.H. Martin 2655), ex: *Myrtella* sp. (3 slides).

Paratype, Papua New Guinea: Morobe Province coast, Buso riverbank, 11.ix.1979, (J.H. Martin 2531), ex: ?*Decaspermum* sp.

Paratype, Papua New Guinea: Morobe Province coast, Buso riverbank, 12.ix.1979, (J.H. Martin 2547), ex: indet.

C. stigmaphylli Russell

Paratypes, Cuba: Hanabanilla Falls, near Cumanayagua, Santa Clara, 7.iv.1925, (J.G. Myers Q.Z. 3388), ex: *Stigmaphyllon sagraeanum*, B.M. 1998-131.

Puerto Rico: El Asomante Rd, 21.iii.1952, (Martorell & L.M. Russell), ex: *Stigmaphyllon tomentosum*, B.M. 1998-131.

C. tuberculatum Russell

Paratype, Puerto Rico: West of Ponce, 26.x.1902, (A.A. Heller), ex: *Coccoloba obtusifolia*, B.M. 1998-131.

Puerto Rico: Central Aguirre, 25.vi.1925, (H.L. Dozier), ex: *Cordia* sp.

50. *Crescentaleyrodes* David & Jesudasan

C. paulianae (Cohic)

Ivory Coast: Lamto, 25.xi.1988, (J.S. Noyes), ex: *Sorghastrum bipennatum* (3 slides).

C. semilunaris (Corbett)

Syntypes, Ceylon: Colombo, xi.1903, (E.E. Green), ex: *Cymbopogon* sp., 232/61, B.M. 1927-451 (4 slides).

India: Vellayani, (via New Delhi), 5.viii.1985, ex: *Cymbopogon flexuosus*, C.I.E. A.17357, 25/85.

India: Burliar, 20.vi.1985, (B.V. David), ex: *Cymbopogon* sp. (2 slides).

Brunei: Borneo, Penanjong, 3.iii.1989, (J.H. Martin 5432), ex: Coastal grass, B.M. 1989-89.

51. *Cryptolingula* Martin & Carver

C. perplexa Martin & Carver

Paratypes, Australia: S.A., Black Hill, 14.ix.1952, (D.E. Symon), ex: indet., (4 slides).

52. *Dialeurodes* Cockerell*D. agalmae* TakahashiHong Kong: Lantau Island, Po Lin, Monastery – Tung Chung path, 26.xi.1999, (J.H. Martin 7300), ex: *Schefflera octophylla* (2 slides).*D. ara* CorbettSingapore: Napier, Cluny Rds, 29.xii.1996, Botanic Gardens, (J.H. Martin 6991), ex: *Calophyllum* sp. (2 slides).*D. armatus* David & SubramaniamParatypes, India: Coimbatore, 5.xi.1966, (B.V. David), ex: *Azadirachta indica*, B.M. 1972-24.*D. bicornicauda* MartinParatypes, Australia: Queensland, 2 km north of Cape Tribulation, 14.xii.1996, (J.H. Martin 6881), ex: *Dysoxylum parasiticum* (2 slides).*D. cerifera* Quaintance & Baker

?Syntypes, South Africa: Am. Bur. Ent. Coll., B.M. 1921-35, (6 slides).

D. cinnamomi Takahashi

Taiwan: Shihmen, 23.i.1987, (C.C. Ker), ex: indet., B.M. 1987-197.

D. citri (Ashmead)USSR: Truziya, Batumi, 23.viii.1960, (E.M. Danzig), ex: *Citrus* sp., B.M. 1963-708.Turkey: Adana, 1976, ex: *Citrus* sp., C.I.E. A.9030, 8/77.USA: Florida, Nursery at Enfield, Middlesex, 19.xi.1974, ex: *Citrus mitis*.

Greece: Corfu, (Olive Inst.), vii.1979, (S. Pappas), ex: indet., 13/80.

Algeria: Boufarik, 14.xi.1978, (F.D. Bennett), ex: *Citrus* sp., C.I.E. A.10842.Italy: Vicenza, Po Valley, x.1993, ex: *Ligustrum* sp., I.I.E. 22957.Taiwan: Meifeng, 9.vi.1986, (C.C. Ker), ex: *Prunus phaeosticta*, B.M. 1987-197.Taiwan: Wuling, 10.xi.1986, (C.C. Ker), ex: *Eriobotrya deflexa*, B.M. 1987-197.*D. cyathispinifera* Corbett

?Syntype, Malaya: Kuala Lumpur, 11.iii.1929, (G.H. Corbett), ex: indet., B.M. 1932-511.

D. decaspermi MartinParatypes, Papua New Guinea: Morobe Province coast, Buso, 10.ix.1979, (J.H. Martin 2528), ex: *Decaspermum* sp., (4 slides).Paratypes, Papua New Guinea: Morobe Province coast, Buso riverbank, 16.x.1979, (J.H. Martin 2713), ex: *Decaspermum* sp.Paratypes, Papua New Guinea: Morobe Province coast, Buso riverbank, 13.x.1979, (J.H. Martin 2690), ex: ?*Decaspermum* sp.Paratypes, Papua New Guinea: Morobe Province coast, Buso riverbank, 23.x.1979, (J.H. Martin 2755), ex: *Decaspermum* sp.*D. dubia* Corbett

West Malaysia: Pahang Prov., Taman Negara, Tembeling path, 12.iii.1984, (J.H. Martin 4263), ex: indet. (3 slides).

D. emarginata (Mound)Holotype, Nigeria: Olokomeji, near Ibadan, iii.1961, (E.A. James), ex: *Cola cordifolia*.Paratypes, Nigeria: Olokomeji, near Ibadan, iii.1961, (E.A. James), ex: *Cola cordifolia* (2 slides).*D. erythrinae* Corbett

Indonesia: Java, Jogjakarta, Hotel Santika, 14.i.2001. (P. De Barro), ex: indet. (2 slides).

D. ixorae Singh?Syntypes, India: Madras, (Y. Ram Chandra Rao), ex: *Ixora coccinea* (2 slides).

?Syntypes, India: Chepauk, Madras, 10.viii.1907, (E. Ballard), ex: *Ixora coccinea*, B.M. 1916-75.

Denmark: import fm. Sri Lanka, 1994, Ministry of Agriculture, Sample no. 1057/94, ex: *Ixora odorata*, I.I.E. 23154.

D. joholensis Corbett

West Malaysia: Pahang Prov., Taman Negara, Tembeling path, 12.iii.1984, (J.H. Martin 4261), ex: indet. (3 slides).

D. kirkaldyi (Kotinsky)

Paralectotypes, Hawaii: Honolulu, Hawaiian Sugar Planters Experimental Station, 22.viii.1906, (J. Kotinsky/Quaintance collection Q.4107), ex: Jessamine, B.M. 1998-131 (2 slides).

Malaya: Kuala Lumpur, 16.x.1973, (L.A. Mound), ex: *Pittosporum ferrugineum*, 68/84.

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 20.ix.1989, (L.D.C. Fishpool), ex: *Clerodendron* sp.

Cyperus: ex: *Jasminum sambac*.

D. lanceolata Takahashi

Singapore: Botanic Gardens, 15.ix.1980, (D.H. Murphy), ex: *Bhesa panicula*, 14/85 (2 slides).

D. minahassai Martin

Paratypes, Indonesia: Sulawesi Utara, Gunung Mogogonipa, 13.vi.1985, (J.H. Martin 5025), ex: *Eugenia caryophyllata* (2 slides).

Paratypes, Indonesia: Sulawesi Utara, Gunung Mogogonipa, 8.iv.1985, (J.H. Martin 4806), ex: *Eugenia caryophyllata*.

D. mirabilis Takahashi

Hong Kong: NT, Tai Lung Farm, Sheung Shui, 13.xii.2001, (J.H. Martin 7539A), ex: *Aporusa dioica*.

D. platicus Bondar

Brazil: Bahia, 7.xii.1923, (G. Bondar), ex: Myrtaceae, B.M. 1924-44.

53. *Dialeurolobus* Danzig

D. rhamni Bink-Moenen

Paratypes, Israel: 1 km S. Solomon's pools, Judaeen Hills, 18.ix.1968, (D. Gerling), ex: *Rhamnus* sp. (2 slides).

Iran: Fars, Farashband, 22.x.1990, (M. Zarrabi 19), ex: *Punica granatum* (2 slides).

Turkey: Iskenderun, 15.xii.1986, (via N. Hygun 3), ex: *Punica granatum*.

54. *Dialeurolonga* Dozier

D. africana (Newstead)

Syntypes, Nigeria: Ibadan, Moor Plantation, v.1917, (C.O. Farquharson), ex: *Salacia* sp. (4 slides).

D. communis Bink-Moenen

Holotype, Chad: Bebedjia, 395 m, 26.xii.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Allophylus africanus*, B.M. 1976-56.

Paratype, Chad: Bebedjia, 395 m, 26.xii.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Allophylus africanus*, B.M. 1976-56.

Paratype, Chad: Baïbokoum, 500 m, 19.xi.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Citrus* sp., B.M. 1976-56.

Sudan: Wad Medani, xi.1961, (L.A. Mound), ex: *Citrus* sp., B.M. 1963-777 (2 slides).

D. elliptica Takahashi

Madagascar: Ihosy, v.1951, (R. Takahashi), ex: *Plumeria* sp., 104/62, B.M. 1962-401 (3 slides).

D. elongata Dozier

Syntypes, India: Punjab, 29.iii.1926, (M.A. Husain), ex: *Citrus* sp., I.B.E. 309 (2 slides).

Syntypes, India: Punjab, ?1928, (M.A. Husain), ex: *Citrus* sp., I.B.E. 309, 46/61, cf. B.M. 1928-181 (2 slides).

Syntypes, India: Punjab, (M.A. Husain), ex: *Citrus* sp., I.B.E. 309, B.M. 1928-181.

India: Maharashtra, Akola, 10.i.1985, ex: *Citrus* sp., C.I.E. A.17113 (2 slides).

D. hoyti Mound

Holotype+paratype, Nigeria: Agege, near Lagos, xi.1959, (C.P. Hoyt), ex: Coffee.

Paratypes, Nigeria: Agege, near Lagos, xi.1959, (C.P. Hoyt), ex: ?Coffee, (4 slides).

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 22.xi.1989, (L.D.C. Fishpool 81), ex: *Trichilia heudelotii*.

D. maculata (Singh)

Patatypes, India: Madras, 3.viii.1971, (B.V. David), ex: *Ficus religiosa*, B.M. 1972-24.

D. nemoralis Bink-Moenen

Holotype, Chad: Bentam (Donia), 451 m, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*, B.M. 1976-56.

Paratype, Chad: Bentam (Donia), 451 m, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*, B.M. 1976-56.

D. phyllarthronis Takahashi

Madagascar: Antsirabé, vii.1955, (R. Paulian), ex: Cerisier du Brésil, 106/62, B.M. 1962-401 (3 slides).

D. ravensarae Takahashi & Mamet

Syntypes, Madagascar: Besanatrihely, Haut Sambirano, 1000 m, x. 1949, (R. Paulian), ex: *Ravensara* sp.

D. rusostigmoides Martin

Paratypes, Australia: Queensland, Cooper's Creek, 20 km S. of Cape Tribulation, 18.xii.1996, (J.H. Martin 6915), ex: indet., B.M. 1998-50 (2 slides).

D. swaini Martin

Paratypes, Australia: Queensland, Broadbeach, viii-ix.1970, (G. Swaine), ex: *Ficus* sp., 16/71, (6 slides).

D. tenella Takahashi

Syntypes, Madagascar: Ambilobé, iv.1951, (R. Paulian), ex: Barabanja, 103/62, B.M. 1962-401 (2 slides).

Syntype, Madagascar: Ambilobé, iv.1951, (R. Paulian), ex: Barabanja.

D. trialeuroides Takahashi & Mamet

Syntypes, Madagascar: Antsohihy, x.1949, (R. Paulian), ex: indet.

55. *Dialeuropora* Quaintance & Baker

D. brideliae (Takahashi)

Taiwan: Keelung, 28.vi.1990, (C.C. Ko), ex: indet., B.M. 1994 (2 slides).

Taiwan: Hungyeh, 26.ii.1990, (C.C. Ko), ex: indet.

D. decempuncta (Quaintance & Baker)

Paralectotypes, Sri Lanka: Royal Botanic Gardens, 7.x.1910, (R.S. Woglum/Quaintance collection Q. 6729A), ex: *Cinnamomum* sp., B.M. 1998-131 (3 slides).

Hong Kong: NT, East Ping Chau I, 5.xii.2001, (J.H. Martin 7528), ex: *Celtis* sp.

Hong Kong: Lantau Island, hillside E. of Shek Pik Reservoir, 15.xi.1996, (J.H. Martin 6820), ex: *Glochidion eriocarpum*.

Hong Kong: Victoria Peak, Harlech Road, 22.x.1990, (J.H. Martin 5826), ex: *Litsea glutinosa*.

West Malaysia: Pahang Prov., Taman Negara, Kuala Tahan, 11.iii.1984, (J.H. Martin 4259), ex: *Flemingia macrophylla* (2 slides).

Australia: WA, Kununurra, 5.v.2000, (J.H. Martin 7430), ex: *Eucalyptus grandifolia*.

D. mangiferae (Corbett)

West Malaysia: Pahang Prov., Genting Highlands at 2000', 24.iii.1984, (J.H. Martin 4314), ex: *Mangifera indica*.

D. papillata Cohic

Syntype, Congo: Brazzaville, 5.iii.1965, (F. Cohic), ex: *Platysepalum vanderystii*, B.M. 1965-305.

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 5.x.1989, (L.D.C. Fishpool 8), ex: *Microdesmis puberula*.

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 17.xi.1989, (L.D.C. Fishpool 79), ex: *Smeathmannia pubescens*.

Sudan: S. Darfur, Jebel Marra, S. W. side, Wadi Golol, 28.iv.1981, (J.H. Martin 3348), ex: *Ficus ?capensis*.

Sudan: S. Darfur, Jebel Marra, S. W. side, Wadi Golol, 30.iv.1981, (J.H. Martin 3358), ex: *Syzygium guineense* (2 slides).

Cameroon: Mbalmayo, viii.2000, (A. Polaszek), ex: *Musa* sp. (2 slides).

D. silvarum (Corbett)

West Malaysia: Pahang Prov., Tioman I., Kampung Tekek, 16.iii.1984, (J.H. Martin 4276), ex: ?Leguminosae (3 slides).

D. viburni (Takahashi)

Taiwan, Nanjenshan, 24.ii.1990, (C.C. Ko), ex: indet.

Taiwan, Nanjenshan, 24.ii.1990, (C.C. Ko), ex: indet., B.M. 1994.

Taiwan, Loloshan, 13.ii.1987, (C.C. Ko), ex: *Turpinia formosana*, B.M. 1994.

56. *Disiphon* Russell*Disiphon* sp.

Belize: Cayo, Chiquibul forest, Las Cuevas plots, 6.xi.1994, (J.H. Martin 6462), ex: *Piper* sp. (2 slides).

Belize: Cayo, Chiquibul forest, Grano de Oro track, 22.xi.1994, (J.H. Martin 6506), ex: ?Laurel.

57. *Dumbletoniella* Jesudasan & David*D. callistemoni* Martin

Paratypes, Australia: A.C.T., Canberra, Black Mountain, Nat. Botanic Gardens, 27.xi.1996, (J.H. Martin 6833), ex: *Callistemon shiresii*, B.M. 1998-50 (2 slides).

Paratypes, Australia: A.C.T., Canberra, Black Mountain, Nat. Botanic Gardens, 27.xi.1996, (J.H. Martin 6833), ex: *Callistemon* sp., B.M. 1998-50.

Australia: New South Wales, Univ. W. Sydney, Hawkesbury – Richmond campus, 15.iv.2000, (J.H. Martin 7387), ex: *Callistemon* sp. (2 slides).

D. eucalypti (Dumbleton)

Paratypes, New Zealand: Waikakaho, Blenheim, 3.xi.1950, (L.J. Dumbleton), ex: *Eucalyptus globulus* (3 slides).

New Zealand: ?Nelson, ?Waskaho, ex: *Eucalyptus globulus*, 9/65, B.M. 1964-721 (2 slides).

New Zealand: ?Nelson, ?Waskaho, ex: *Eucalyptus globulus*, B.M. 1964-721.

New Zealand: ?Nelson, ex: *Eucalyptus globulus*, 9/65, B.M. 1964-721.

D. pittospori Martin & Carver

Paratypes, Australia: Victoria, Mallee, 1925, (J.E. Dixon), ex: *Pittosporum phylliraeoides*, 32/61, B.M. 1961-4 (3 slides).

Paratypes, Australia: Adelaide, Waite Hill, 23.xi.1967, (L.A. Mound), ex: *Pittosporum phylliraeoides*, 36/71 (3 slides).

D. rotunda Martin & Carver

Paratypes, Australia: Vic, Indigo, 17.ix.1914, (E.E. Pescott), ex: *Eucalyptus leucoxydon*.

Paratypes, Australia: Queensland, Barakula, 5.x.1939, ex: *Eucalyptus crebra* (4 slides).

Paratypes, Australia: Western Australia, Perth, King's Park, 4.iii.1959, (V.F. Eastop 7362), ex: *Banksia* sp. (?*Dryandra* sp.), 149/61, B.M. 1960-144. (In the original description ex: *Eucalyptus* sp.).

D. xanthorrhoeae Martin

Paratypes, Australia: S.A., Kangaroo Island, Flinders Chase N. P., 17.ii.1990, (J.H. Martin 5657), ex: *Xanthorrhoea australis*, B.M. 1998-50, (5 slides).

58. *Extensaleyrodes* Bink-Moenen

E. akurensis (Mound)

Holotype, Nigeria: Akure, 8.i.1957, (V.F. Eastop 6511), ex: indet., 65/61.

Paratypes, Nigeria: Akure, 8.i.1957, (V.F. Eastop 6511), ex: indet., 65/61 (2 slides).

E. falcata Bink-Moenen

Holotype, Chad: Boro, 453 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Isobertinia doka*, B.M. 1976-56.

Paratypes, Chad: Boro, 453 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Isobertinia doka*, B.M. 1976-56 (2 slides).

59. *Fascaleyrodes* Bink-Moenen

F. palmae (Gameel)

Nigeria: Moor Plantation area, iii.1961, (?L.A. Mound), ex: Palm (2 slides).

Nigeria: Moor Plantation area, iv.1961, (?L.A. Mound), ex: Palm.

F. rara Bink-Moenen

Holotype, Chad: Donia (Bentam, in the original description), 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*.

Paratypes, Chad: Donia (Bentam, in the original description), 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*.

60. *Filicaleyrodes* Takahashi

F. bosseri Takahashi

Syntypes, Madagascar: Tsimbazaza, Tananarive, viii.1956, (J. Bosser), ex: Fern, 114/61, B.M. 1961-455.

F. williamsi (Trehan)

Lectotype, England: Surrey, Kew Gardens, 17.ix.1937, (K.N. Trehan and/or C.B. Williams), ex: *Oleandra africana*, 207/62, B.M. 1962-643.

Type series, England: Surrey, Kew Gardens, 20.i.1937, (K.N. Trehan and/or C.B. Williams), ex: *Oleandra* sp., 207/62, B.M. 1962-643.

Type series, England: Surrey, Kew Gardens, ii.1938, ex: Ferns.

61. *Fippataleyrodes* Sundararaj & David

F. indica Sundararaj & David

Paratype, India: Cumbum, 10.i.1988, (S. Selvakumaran), ex: *Ficus racemosa*.

F. litseae ?Sundararaj & David

Paratype, India: Mahableshwar, 28.iii.1987, (B.V. David), ex: *Litsea stocksii*.

62. *Gagudjuia* Martin

G. allosyncarpiae Martin

Paratypes, Australia: Northern Territory, Kakadu Nat. Park, Gubara Walk, Burdulba Creek, 20 km S. of Jabiru, 22.xii.1996, (J.H. Martin 6955), ex: *Allosyncarpia ternata* (6 slides).

63. *Gomenella* Dumbleton

G. dryandrae (Takahashi)

Lectotype, Australia: Western Australia, Nedlands, 20.v.1940, (K.R. Norris), ex: *Dryandra floribunda*, 7/40, B.M. 1955-799.

Australia: Western Australia: Perth, King's Park, 4.iii.1959, (V.F. Eastop 7362 × 1), ex: *Banksia* sp. (or ?*Dryandra* sp.), B.M. 1960-144.

Australia: New South Wales, Cowan, 24.x.1978, (F. Wheelhouse 10), ex: *Banksia spinulosa*, C.I.E. A.12322.

USA ex-Australia: New York, 30.iii.1974, (R. Scott), ex: *Banksia hookeriana*, B.M. 1998-131.

G. multipora Dumbleton

Paratype, New Caledonia: Tinip, 4.xi.1954, (L.J. Dumbleton), ex: indet., B.M. 1964-221.

64. *Harpaleyrodes* Bink-Moenen

H. tuberculata Bink-Moenen

Paratypes, Chad: Bebedjia, 390 m, 17.vi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Combretum hypopilinum*, B.M. 1977-471 (3 slides).

65. *Hempelia* Sampson & Drews

H. chivelensis Sampson & Drews

Syntype, Mexico: Chivela, Oaxaca, iv.1926, (G.F. Ferris), ex: indet., B.M. 1998-131.

Belize: Cayo, Chiquibul forest, Las Cuevas Res. Stn, 4.xi.1994, (J.H. Martin 6451), ex: *Persea ?americana* (3 slides).

Belize: Cayo, Chiquibul forest, Las Cuevas clearing, 10.ii.1996, (J.H. Martin 6633), ex: *Persea* sp.

66. *Heteraleyrodes* Takahashi

H. bambusicola Takahashi

Holotype, Malaya: Kuala Lumpur, 13.iii.1944, (R. Takahashi), ex: Bamboo, B.M. 1955-799.

West Malaysia: Kuala Lumpur, Lake Gardens, 11.ii.1985, (J.H. Martin 4454), ex: Bamboo.

67. *Heterobemisia* Takahashi

H. alba Takahashi

Syntype, Japan: Odaig-ahara, Nara Pref., 15.viii.1956, (R. Takahashi), ex: *Itea japonica*.

68. *Indoaleyrodes* David & Subramaniam

I. glochidioni Martin & Carver

Paratypes, Australia: Queensland, Great Keppel Island, 2.vii.1991, (J.J. Bruhl & G.N. Batianoff), ex: *Glochidion lobocarpum*, B.M. 1998-51 (2 slides).

Australia: Queensland, 50 km S. of Bowen, 20.iv.2001, (P. De Barro 76), ex: *Glochidion* sp. *I. laos* (Takahashi)

Paratypes: India: Coimbatore, 25.iii.1967, (B.V. David), ex: *Morinda tinctoria*, B.M. 1972-24 (2 slides).

India: Madras, 6.i.1972, (B.V. David), ex: *Morinda tinctoria*, 3/72, (4 slides).

I. pseudoculatus Martin

Paratypes, Papua New Guinea: Morobe Province coast, Buso, 8.x.1979, (J.H. Martin 2674), ex: *Syzygium* sp. (4 slides).

69. *Kanakarajiella* David & Sundararaj

K. vulgaris (Singh)

India: Chikmugalur, Coffee Research Stn, i-ii.1969, ex: Coffee, C.I.E. A2803, 1/69 (4 slides).

Burma: Rangoon, 28.viii.1984, (T.J. Crowe), ex: *Jasminum* sp., C.I.E. A16401, 147/84 (2 slides).

70. *Leucopogonella* Dumbleton

L. pallida Dumbleton

Paratypes, New Caledonia: Plum, (L.J. Dumbleton), ex: *Leucopogon* sp., B.M. 1964- 721.

L. simila Dumbleton

Paratypes, New Caledonia: Plum, (L.J. Dumbleton), ex: *Leucopogon* sp., B.M. 1964- 721.

71. *Lipaleyrodes* Takahashi*L. atriplex* (Froggatt)

Paralectotype, Australia: N.S.W. Broken hill, 25.v.1911, (W.W. Froggatt), ex: Saltbush (2 slides).

L. breyniae (Singh)Maldives: Addu Atoll, Feydhoo, 12.xi.1993, (G.W. Watson 100), ex: *Phyllanthus distichus*, I.I.E. 22974, (4 slides).*L. crossandrae* David & SubramaniamParatypes, India: Coimbatore, 18.xi.1966, (B.V. David), ex: *Achyranthes aspera*, B.M. 1972-24 (2 slides).*L. euphorbiae* David & SubramaniamParatypes, India: Madurai, 28.i.1967, (B.V. David), ex: *Euphorbia prostrata*, B.M. 1972-24.India: Hebbal, Bangalore, 20.vi.1997, (via A. Polaszek), ex: *Phyllanthus fratteranae*.Kenya: Ganze, 30.xi.1983, (I. Robertson 9/9), ex: *Phyllanthus amarus*, 21/84 (3 slides).*L. phyllanthi* TakahashiSyntypes, Madagascar: Massif de l'Tremo, 1700 m, (J. Bosser), ex: *Phyllanthus* sp., 115/61, B.M. 1961-455.Hong Kong: HK, Tai Tam, Interm. Reservoir, 6.xii.1999, (J.H. Martin 7311), ex: *Blumea* sp.**72. *Malayaleyrodes* Corbett***M. ?lumpurensis* corbettWest Malaysia: Pahang Prov., Pulau Tioman I., Kampung Tekek, 17.ii.1985, (J.H. Martin 4509), ex: ?*Tetracera* sp.**73. *Marginalleyrodes* Takahashi***M. ixorae* TakahashiSyntypes, Madagascar: Between Mahattsinjo and Andriba, ex: *Ixora* sp., 102/62, B.M. 1962-401 (3 slides).**74. *Massilieurodes* Goux***M. americanus* JensenUSA: Maryland, Prince George's Co., Beltsville, Beaver Dam Road area, 2.vii.1998, (J.H. Martin 7108), ex: *Ilex* sp.USA: Maryland, Prince George's Co., Beltsville, USDA campus, 3.vii.1998, (J.H. Martin 7112), ex: *Ilex* sp. (2 slides).*M. chittendeni* (Laing)Lectotype, England: Ascot, 23.v.1928, (G. Fox-Wilson), ex: *Rhododendron* sp.Paralectotypes, England: Ascot, vi.1928, (G. Fox-Wilson), ex: *Rhododendron* sp., (4 slides).England: Surrey, Richmond Park, 19.vi.1984, (J.H. Martin 4352), ex: *Rhododendron* sp.England: Surrey, Richmond Park, 27.vi.1984, (J.H. Martin 4353), ex: *Rhododendron* sp.Italy: Cermenake (Co.), vi.2000, (M. Colombo 2), I.I.E. 24199, ex: *Rhododendron* ifs, B.M. (E) 2001-110.*M. euryae* (Takahashi)

Syntypes, Taiwan: Neng Kao, 6.viii.1939, (R. Takahashi), ex: *Eurya glaberrima*, B.M. 1994 (3 slides).

M. fici (Takahashi)

Taiwan: Sanhsing, 23.ix.1993, (K.C. Chou), ex: *Ficus virgata*, B.M. 1994 (2 slides).

Taiwan: Taipei, 4.x.1986, (C.C. Ker), ex: *Bischofia javanica*.

Taiwan: Taichung, 5.ix.1967, (Y.C. Chang 8), ex: *Bischofia javanica*, 20/67, B.M. 1967-592 (3 slides).

M. setiger (Goux)

France: Corsica, Solenzara, Col de Larone road, D268, 27.x.1999, (J.H. Martin 7237), ex: *Viburnum tinus*.

Italy: Campania, Portici grounds, 10.vi.1987, (A. Polaszek & S. Laudonia), ex: *Viburnum* sp., C.I.E. A19249, 4/88 (2 slides).

Italy: Portici, 9.iv.1984, (F.M. Laccarino), ex: *Viburnum tinus*, 59/84.

Morocco: Middle Atlas, Ifrane, ± 1600 m, 26.iv.1992, (J.H. Martin 6076), ex: *Viburnum tinus*, B.M. 1992-73 (2 slides).

75. *Metabemisia* Takahashi

M. filicis Mound

Paratypes, Scotland: Edinburgh, Royal Botanic Gardens, glasshouse, 27.v.1966, (L.A. Mound & B.R. Pitkin), ex: *Dryopteris* sp. (4 slides).

Paratypes, Scotland: Edinburgh, Royal Botanic Gardens, glasshouse, 31.i.1966, (E.C. Pelham-Clinton), ex: Ferns, B.M. 1966-73 (4 slides).

M. palawana Martin

Paratypes, Philippine Islands: Palawan, Irawan/Iwahig, forest, 2000', 11.ii.1988, (J.H. Martin 5265), ex: *Lastreopsis* ?*subsparsa*, B.M. 1988-2 (2 slides).

76. *Minutaleyrodes* Jesudasan & David

M. minuta (Singh)

India: Madras, 11.ii.1972, (B.V. David), ex: *Ixora coccinea*, B.M. 1972-24.

Singapore: Botanic Gardens, 15.ix.1980, (D.H. Murphy), ex: *Bhesa paniculata*, 14/58.

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 1.xi.1989, (L.D.C. Fishpool 31), ex: *Ixora* sp.

Taiwan: Fengshan, 22.xi.1993, (K.C. Chou), ex: *Ixora* sp.

M. suishanus (Takahashi)

Philippine Islands: Palawan, Honda Bay, 7.ii.1988, (J.H. Martin 5251), ex: *indet.*, B.M. 1988-2 (2 slides).

77. *Mixaleyrodes* Takahashi

M. ?polystichi Takahashi

Taiwan: Lienhuachih, 2.v.1994, (K.C. Chou), ex: *Angiopteris lygodiiifolia*, B.M. 1994 (2 slides).

78. *Neoaleurotrachelus* Takahashi & Mamet

N. bertilloni (Cohic)

Syntype, Congo: Brazzaville, 21.iii.1964, (F. Cohic), ex: *Tetracera alnifolia*, B.M. 1965-305.

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 6.xi.1989, (L.D.C. Fishpool 40), ex: *Cryptolepis sanguinolenta*.

N. graberi (Cohic)

Sudan: S. Darfur, Jebel Marra, S. W. side, Wadi Golol, 28.iv.1981, (J.H. Martin 3348), ex: *Ficus ?capensis*.

Sudan: Khartoum, Botanic Garden, 13.iv.1981, (J.H. Martin 3280), ex: indet. (2 slides).

Sudan: Wad Medani, xii.1962, (L.A. Mound), ex: *Zizyphus spinachrish*, B.M. 1963-777 (2 slides).

Sudan: Wad Medani, i.1962, (L.A. Mound), ex: *Lawsonia alba*, B.M. 1963-777.

Chad: Timberi, 407 m, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Gardenia triacantha*, 74/81.

79. *Neomaskellia* Quaintance & Baker

N. andropogonis Corbett

Syntypes, Ceylon: Bandarawalla, v.1906, (E.E. Green), ex: *Andropogon* sp., B.M. 1927-451.

Hong Kong: NT, west Tai Mo Shan, Country Park, Ho Pui reservoir, 13.xii.2001, (J.H. Martin 7567), ex: *Saccharum ?spontaneum*, (2 slides).

India: Lucknols, Inst. Sugar Cane Res., iv.1966, ex: Sugar Cane, C.I.E. A1284, 26/66 (2 slides).

West Malaysia: Pahang Prov., Taman Negara, Kuala Tahan, 10.iii.1984, (J.H. Martin 4247), ex: *Saccharum* sp.

West Malaysia: Selangor Prov., Genting Highlands, 1.v.1994, (D.J. Williams 1507), ex: indet.

Pakistan, Rawalpindi, 1.iv.1960, ex: *Bauhinia variegata*, C.I.E. 18414, 16/63.

N. bergii (Signoret)

S. Nigeria: Ibadan, 6.i.1964, (W.A. Lambom), ex: Grass, 40/62 (2 slides).

E. Nigeria: Ikom, 11.ii.1957, (Eastop 6766), ex: Elephant grass.

Brunei: 1971, (Kassim bin Ahmad), ex: Sugar Cane, C.I.E. A4463/18, 1/71 (2 slides).

Seychelles: Mahé, v.1970, (P.L. Mathias), ex: Gramineae, 1/72 (3 slides).

Papua New Guinea: Morobe Prov., Lasanga Island, 20.x.1979, (J.H. Martin 2735), ex: *Saccharum ?officinarum* (2 slides).

Papua New Guinea: Buba, Dept. of Agriculture, 15.vii.1968, ex: Sugar Cane, C.I.E. A3299/12, 11/69.

Pakistan: Dacca, Locust Warning & Plant Quar. Dept., 27.vi.1967, ex: Sugar Cane, C.I.E. A1797/10, 16/67.

Topotypic, Fiji Islands: (R. Veitch), ex: Gramineae, B.M. 1915-240 (3 slides).

80. *Neopealius* Takahashi

N. rubi Takahashi

Taiwan: 24.xi.1993, (Warren), ex: *Rubus* sp., 22176/7, 93-1369, I.I.E. 22997.

81. *Nigrasialeyrodes* Martin & Carver

N. convexus Martin & Carver

Paratypes: Australia: Queensland, Townsville, James Cook University campus, 2.vi.1992, (D.W. Burrows), ex: *Melaleuca viridiflora*, B.M. 1998-51 (3 slides).

Paratypes: Australia: Queensland, Townsville, James Cook University campus, 19.xii.1991, ex: *Melaleuca quinquenervia*, B.M. 1998-51.

82. *Orchamoplatus* Russell

O. caledonicus (Dumbleton)

Paratype, New Caledonia: Noumea, 24.v.1955, (L.J. Dumbleton), ex: *Citrus* sp., B.M. 1964-721.

New Caledonia: 28.ii.1956, (J.F. Schoen 56-2755), ex: *Ochrosia* sp., B.M. 1998-131.

O. citri (Takahashi)

Australia: Adelaide, xi.1967, (L.A. Mound), ex: *Citrus* sp., 35/71 (4 slides).

Lord Howe Island: xi.2001, (L.A. Mound), ex: *Psidium cattelianum* (2 slides).

O. dentatus (Dumbleton)

Paratype, New Caledonia: Carenage, 8.iv.1955, (L.J. Dumbleton), ex: Myrtaceae, B.M. 1964-721.

New Caledonia: near Nouméa, 1912, (Franc), ex: *Microsemma salicifolia* (2 slides).

O. louiserussellae Martin

Paratypes, Australia: Queensland, strand 2 km north of Cape Tribulation, 15.vii.1996, (J.H. Martin 6883), ex: *Syzygium* sp., B.M. 1998-50 (5 slides).

O. mammaeferus (Quaintance & Baker)

Lectotype, Samoa: Upolu, Apia, iv.1925, (P.A. Buxton & G.H. Hopkins), ex: Croton.

Paralectotypes, Samoa: Upolu, Apia, iv.1925, (P.A. Buxton & G.H. Hopkins), ex: Croton (3 slides).

Cook Islands: Aitutaki, ii.1960, (N.L.K. Krauss 2), ex: indet. (3 slides).

Fiji: Viti Levu, Rakiraki, iv.1982, (L.A. Mound), ex: indet., 14/82.

Fiji: Nausori, 30.iii.1967, (G. Swaine), ex: Croton, C.I.E. A1757, 15/67.

Fiji: Viti Levu, Suva, 23.iii.1976, (P. Maddison), ex: *Codiaeum variegatum*, C.I.E. A9356.

French Polynesia: Tahiti, 9.vii.1976, (P. Maddison), ex: *Citrus reticulata*, C.I.E. A9023.

French Polynesia: Tahiti, 8.vii.1976, (P. Maddison), ex: *Citrus grandis*, C.I.E. A9023.

O. montanus (Dumbleton)

Paratypes, New Caledonia: Plaine des Lacs, 8.iv.1955, (L.J. Dumbleton), ex: indet., B.M. 1964-721 (2 slides).

New Caledonia: near Nouméa, 1912, (Franc), ex: *Microsemma salicifolia* (2 slides).

O. niuginii Martin

Paratypes, Papua New Guinea: Morobe Prov. coast, Lasanga Island, 18.ix.1979, (J.H. Martin 2581), ex: *Calophyllum inophyllum* (4 slides).

Paratypes, Papua New Guinea: Morobe Prov. coast, Buso, 30.ix.1979, (J.H. Martin 2643), ex: *Calophyllum inophyllum* (2 slides).

Paratypes, Papua New Guinea: Morobe Prov. coast, Buso, 8.ix.1979, (J.H. Martin 2676), ex: *Durandea* sp. (2 slides).

O. noumeae Russell

Paratype, New Caledonia: Noumea, 31.x.1940, (F.X. Williams), ex: *Citrus* sp., B.M. 1998-131.

O. plumensis (Dumbleton)

Paratype, New Caledonia: Plum, (L.J. Dumbleton), ex: indet., B.M. 1964-721.

O. porosus (Dumbleton)

Paratype, New Caledonia: Carenage, 8.iv.1955, (L.J. Dumbleton), ex: Myrtaceae, B.M. 1964-721.

83. *Orientalleyrodes* David

O. zeylanicus (Corbett)

Syntypes, Sri Lanka: Trincomalee, v.1906, (T.B. Fletcher), ex: indet., B.M. 1927-451.

Syntypes, Sri Lanka: Trincomalee, vi.1906, (T.B. Fletcher), ex: indet., B.M. 1927-451 (3 slides).

84. *Orstomaleyrodes* Cohic

O. fimbriata (Mound)

Paratypes, Nigeria: Ibadan, Moor Plantation, ii.1961, (L.A. Mound), ex: *Cassia siamea* (5 slides).

Paratypes, Nigeria: Ibadan, Moor Plantation, i.1961, (L.A. Mound), ex: *Cassia siamea*.

Paratypes, Nigeria: Ibadan, Moor Plantation, iii.1961, (E.A. James), ex: *Cassia siamea*.

Congo: Brazzaville, 29.i.1964, ex: *Caloncoba welwitschii*, Al 14-65, B.M. 1965-305.

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 3.xi.1989, (L.D.C. Fishpool 33), ex: *Milletia zechiana*.

85. *Papillipes* Bink-Moenen

P. spinifer Bink-Moenen

Holotype, Chad: Bebedjia, 395 m, 4.i.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Daniellia olivera*, B.M. 1977-471.

86. *Parabemisia* Takahashi

P. aceris (Takahashi)

Taiwan: Taipei, 15.i.1986, (C.C. Ker), ex: *Acer* sp., B.M. 1987-197.

P. javani Martin

Paratypes, Papua New Guinea: Morobe Prov. coast, Jawani Island, 2.xi.1979, (J.H. Martin 2789), ex: ?*Timonius* sp. (5 slides).

P. lushanensis Ko & Luo

Paratypes, Taiwan: Nengkao, 24.x.1989, (C.C. Ko), ex: indet.

P. myricae (Kuwana)

Australia: Queensland, Carnarvon Gorge, 24.x.2001, (P. De Barro 107), ex: *Glochidion* sp.

Israel: Hefar, Placcab, 26.ix.1978, (A. Sternlicht), ex: *Citrus* sp., 15/78.

Israel: Akko, iii.1979, (A. Sternlicht), ex: *Citrus* or *Persea*, 1/69.

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 20.xii.1989, (L.D.C. Fishpool 115), ex: *Persea americana*.

Turkey: Adana, 15.ii.1986, (E. Sekeroglu), ex: *Citrus* sp., 2/86.

P. myrmecophila Martin

Paratypes, Papua New Guinea: Morobe Prov. coast, Buso, 27.ix.1979, (J.H. Martin 2629), ex: *Cryptocarya* sp. (3 slides).

Paratypes, Papua New Guinea: Morobe Prov. coast, Buso, 27.ix.1979, (J.H. Martin 2626), ex: *Anisoptera thurifera* (2 slides).

Paratypes, Papua New Guinea: Morobe Prov. coast, Buso, 27.ix.1979, (J.H. Martin 2627), ex: *Macaranga* sp.

Paratypes, Papua New Guinea: Morobe Prov. coast, Buso, 26.ix.1979, (J.H. Martin 2622), ex: *Prunus* sp.

Paratypes, Papua New Guinea: Morobe Prov. coast, Buso, 5.xi.1979, (J.H. Martin 2799), ex: *Prunus* sp.

87. *Paraleurolobus* Sampson & Drews

P. chamaedoreae Russell

Paratypes, Mexico: intercepted at San Antonio, USA, 4.i.1970, (C. Parker), ex: *Chamaedorea* sp., B.M. 1998-131.

Paratype, Mexico: intercepted at San Antonio, USA, ix.1974, (C. Parker), ex: *Chamaedorea* sp.

Paratypes, Mexico: intercepted at San Antonio, USA, 2.v.1970, (D. Johnston), ex: *Chamaedorea* sp., B.M. 1998-131.

Costa Rica: N. Heredia Province, ~10 km north of Puerto Viejo, 2.ii.1983, (J.H. Martin 3746), ex: indet.

Costa Rica: N. Heredia Province, ~10 km north of Puerto Viejo, 2.ii.1983, (J.H. Martin 3747), ex: ?*Flacoutiae* sp.

88. *Paulianaleyrodes* Cohic

P. splendens Cohic

Syntype, Congo: Brazzaville, 10.xii.1964, ex: *Dichapetalum brazzae*, B.M. 1965-305.

P. tetracerae Cohic

Syntype, Congo: Brazzaville, 8.i.1964, ex: *Tetracera alnifolia*, B.M. 1965-305.

89. *Pealius* Quaintance & Baker

P. akebiae (Kuwana)

Japan: Hikawa, Tokyo, 9.viii.1950, (R. Takahashi), ex: *Akebia lobata*, 97/62, B.M. 1962-401 (3 slides).

P. azaleae (Baker & Moles)

Paratypes, India: Ootacamund (Nilgiris), 7000', 3.vii.1969, (B.V. David), ex: *Azalea indica*, B.M. 1972-24.

Australia: A.C.T., Canberra, Weston, 19.xi.1996, (J.H. Martin 6826), ex: *Azalea* sp., B.M. 1998-50 (3 slides).

England: Surrey, Wisley, R.H.S. gdns., xi.1968, (K.M. Harns), ex: *Azalea* sp.

England: Wisley, R.H.S. gdns., 8.vii.1965, (L.A. Mound 76), ex: *Rhododendron mucronatum*.

Taiwan: Fenchifu, 11.iii.1990, (C.C. Ko), ex: *Rhododendron* sp.

P. cryptus Martin

Paratypes, Australia: Northern Territory, Kakadu Nat. Park, Nawurlandja lookout, 25 km S. of Jabiru, 21.xii.1996, (J.H. Martin 6948), ex: *Ficus platypodia*, B.M. 1998-50 (3 slides).

P. euryae (Takahashi)

Syntypes, Japan: Ome, nr Tokyo, 17.viii.1949, (R. Takahashi), ex: *Eurya* sp., 96/62, B.M. 1962-401 (3 slides).

P. fici Mound

Paratypes, Nigeria: Samaru, Zaria, 26.x.1960, (M.O. Ezeigwe), ex: *Ficus asperifolia* (3 slides).

Paratypes, Nigeria: Agege nr Lagos, 10.viii.1960, (M.O. Ezeigwe), ex: *Ficus asperifolia*.

Paratype, Nigeria: Ibadan, Oyo Road, vi.1961, ex: *Ficus* sp.

Paratypes, Nigeria: Ibadan, 16.v.1956, (V.F. Eastop 5950), ex: *Ficus* sp.

Sudan: S. Darfur, Jebel Marra, S. W. side, Wadi Golol, 28.iv.1981, (J.H. Martin 3348), ex: *Ficus ?capensis*.

Sudan: S. Darfur, Jebel Marra, S. W. side, Wadi Golol, 29.iv.1981, (J.H. Martin 3353), ex: *Ficus ?glumosa*.

P. kelloggi (Bemis)

USA: California, Sacramento, 11.vii.1966, (R.F. Wilkey), ex: *Prunus lyoni*, 25/66 (8 slides).

P. liquidambari (Takahashi)

Taiwan: Taipei, 28.xii.1985, (C.C. Ko), ex: *Liquidambar formosana*.

Taiwan: Taipei, 22.xi.1985, (C.C. Ko), ex: *Liquidambar formosana*, B.M. 1987-197.

P. madeirensis Martin *et al.*

Paratypes, Madeira: Faja da Nogueira, hydro dam, 28.iii.1995, (J.H. Martin & A.F. Aguiar, JHM 6552), ex: *Picconia excelsa* (2 slides).

Paratypes, Madeira: Ribeiro Frio, 800 m, 28.iii.1995, (J.H. Martin & A.F. Aguiar, JHM 6545), ex: *Picconia excelsa* (2 slides).

Paratypes, Madeira: Santana, Caldeirao do Inferno, 985 m, 27.vi.1993, (M. Pita C214), ex: *Picconia excelsa*.

P. mori Takahashi

Taiwan: Taipei, 18.vi.1968, (Y.C. Chang), ex: *Morus australis*, 4/68 (4 slides).

Taiwan: Taipei, 26.iv.1992, (C.C. Ko), ex: *Ficus* sp.

Thailand: Bangkok, iii.1973, (K. Sombatsiri), ex: Mulberry, 6/73 (3 slides).

P. rhododendri (Takahashi)

Hawaiian Islands: Hawai'i, Akaka Falls, 10 mi. N. of Hilo, 5.iv.1999, (J.H. Martin 7207), ex: *Rhododendron* sp. (2 slides).

Hong Kong: H K Island west, University campus, 12.xi.1996, (J.H. Martin 6784), ex: *Rhododendron pulchrum* (4 slides).

Japan: Minoh nr Osaka, 29.viii.1955, (R. Takahashi), ex: *Rhododendron* sp., 95/62, B.M. 1962-401 (3 slides).

90. *Pectinaleyrodes* Bink-Moenen

P. culcasiae (Cohic)

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 20.x.1989, (L.D.C. Fishpool 20), ex: *Culcasia liberica* (2 slides).

P. trichlisiae (Cohic)

Syntype, Belgian Congo: Brazzaville, 7.ix.1965, ex: *Trichlisia gilletii*, B.M. 1965-305.

Chad: Donia, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: Annonaceae, B.M. 1976-56 (2 slides).

91. *Pentaleyrodes* Takahashi

P. cinnamomi (Takahashi)

Taiwan, Wulai, 27.xii.1931, (R. Takahashi), ex: *Machilus* sp., B.M. 1994 (3 slides).

P. hongkongensis Takahashi

Hong Kong: Pok Fu Lam, Country Park, below High West, 12.xii.2001, (J.H. Martin 7557), ex: *Litsea rotundifolia* var. *oblongifolia* (2 slides).

Hong Kong: Victoria Peak, 3.iii.1990, (J.H. Martin 5693), ex: *Machilus chinensis* (2 slides).

P. yasumatsui Takahashi

Japan: Nara, 10.v.1959, (R. Takahashi), ex: *Lindera* sp., 99/62, B.M. 1962-401, (5 slides).

92. *Pogonaleyrodes* Takahashi

P. fastuosa Takahashi

Syntype, Madagascar: Manjakatempo, 2000 m, 24.v.1950, (R. Mamet), ex: indet., B.M. 1998-131.

P. zimmermanni (Newstead)

Syntypes, German East Africa (Tanzania): Amani, ix.1902, (A. Zimmermann 19), ex: Acanthaceae, 48/62 (2 slides).

Nigeria: Ibadan, Moor Plantation, xi.1959, (E.A. James), ex: indet.

Nigeria: Ibadan, Moor Plantation, vii.1960, (M.O. Ezeigwe), ex: indet.

Tanganyika: Tukuyu, South Highlands Prov., xii.1957, (R.G. Tapley), ex: *Coffea arabica*, C.I.E. 4678/15780, 51/62, B.M. 1962-3 (2 slides).

Cote d'Ivoire (Ivory Coast): 20 km W. Abidjan, Adiopodoume, IIRSDA, 17.xi.1989, (L.D.C. Fishpool 74), ex: *Aidia genipiflora*.

Kenya: Gedi forest, south of Malindi, 17.v.1988, (J.H. Martin 5318), ex: *Tricalysia ovalifolia*.

93. *Pseudaleuroplatus* Martin

P. kiensis Martin

Paratypes, Australia: S.A., Kangaroo Island, 17.ii.1990, (J.H. Martin 5652), ex: *Melaleuca* sp., B.M. 1998-50 (2 slides).

94. *Ramsesseus* Zahradnik

R. follioti Zahradnik

Paratype, Egypt: Ramsesseum, 10.i.1963, (J. Zahradnik), ex: *Acacia* sp.

95. *Rhachisphora* Quaintance & Baker

R. alishanensis Ko et al.

Paratype, Taiwan: Alishan, 26.iv.1990, (C.C. Ko), ex: *Eurya* sp.

Paratypes, Taiwan: Alishan, 26.iv.1990, (C.C. Ko), ex: *Eurya strigillosa*.

Paratype, Taiwan: Tatachiaanpu, 27.iv.1990, (C.C. Ko), ex: *Eurya* sp.

R. elongatus Regu & David

Paratype, India: Kunnathoor, 24.i.1989, (K. Regu), ex: *Mimusops elengi*.

R. franksae Martin

Paratypes, Australia: Queensland, 2 km north of Cape Tribulation, 13.xii.1996, (J.H. Martin 6870), ex: *Fagraea gracilipes*, B.M. 1998-50 (3 slides).

R. koshunensis (Takahashi)

Taiwan: Kentingkunyaen, 22.i.1990, (W.J. Wu), ex: *Cinnamomum reticulatum* (2 slides).

R. maesae (Takahashi)

Taiwan: Nanshanchi, 28.x.1989, (C.C. Ko), ex: *Maesa* sp.

Taiwan: Jihyuetan, 9.vi.1986, (C.C. Ker), ex: *Maesa japonica*, B.M. 1987-197.

Taiwan: Chiaochiwenchuan, 14.ii.1990, ex: *Maesa* sp.

Malaya: Cameron Highlands, Tanah Rata, 12.x.1973, (L.A. Mound), ex: indet. shrub, 66/84.

R. rutherfordi (Quaintance & Baker)

India: Kunnathoor, 24.i.1989, (K. Regu), ex: *Loranthus elasticus*.

R. trilobitoides (Quaintance & Baker)

Ceylon: Sigiriya, 23.viii.1909, (E.E. Green), ex: indet., B.M. 1927-451.

Ceylon: Trincomalee, ix.1910, (E.E. Green), ex: ?Tree, B.M. 1927-451 (2 slides).

Ceylon: Maha, ix.1909, (E.E. Green), ex: *Memecylon* sp., B.M. 1927-451.

Ceylon: Kesbewa, iv.1903, (E.E. Green), ex: *Memecylon* sp., B.M. 1927-451.

Pakistan: Lahore, 24.iii.1981, ex: *Syzygium cumini*, C.I.E. A12961 (2 slides).

Philippine Islands: Palawan, Irawan forest, 1000', 19.i.1988, (J.H. Martin 5133), ex: indet., B.M. 1988-2.

96. *Rositaleyrodes* Meganathan & David

R. oplismeni (Takahashi)

India: Kunnathoor, 25.i.1989, (K. Regu), ex: Grass.

97. *Rugaleyrodes* Bink-Moenen

R. angolensis (Cohic)

Chad: Doba, 389 m, 21.vi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Syzygium guineense*, 73/81.

R. bidentata Bink-Moenen

Holotype+paratypes, Chad: Baibokoum, 19.xi.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*.

R. tetracerae (Cohic)

Syntype, Congo: Brazzaville, 18.iii.1964, (F. Cohic), ex: *Tetracera alnifolia*, B.M. 1965-305.

98. *Rusostigma* Quaintance & Baker

R. eugeniae (Maskell)

Syntypes, India: Maharashtra, Poona, (W.M. Maskell), ex: *Eugenia jambolana*, 13/85 (3 slides).

India: Gujarat, Navsari, ix.1985, ex: *Syzygium* sp., C.I.E. 18160, 27/86 (2 slides).

R. radiirugosa (Quaintance & Baker)

Syntypes, Indonesia: Billiton Isle, 5.ii.1911, (R.S. Woglum), ex: Mango, Q.6727-A, B.M. 1998-131.

Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Clark's camp area, 10.v.1985, (J.H. Martin 4934), ex: [*Eugenia* sp.] *Syzygium* sp.

West Malaysia: Pahang Prov., Taman Negara, Bukit Teresek, 2.iii.1984, (J.H. Martin 4202), ex: indet.

Malaysia: Sarawak, Gn. Mulu Nat. Park, Kerangas plot, 14.x.1977, (D. Hollis M9/13), ex: indet., 53/82 (2 slides).

Singapore: Bukit Timah Forest, 31.iii.1989, (J.H. Martin 5543), ex: *Eugenia* sp., B.M. 1989-89.

Brunei: Temburong District, Sungai, Temburong, Kuala Baki, 16.x.1992, (J.H. Martin 6145), ex: indet.

R. tokyonis (Kuwana)

Japan: Tokyo, 9.viii.1949, (R. Takahashi), ex: *Sakakia* sp., 94/62, B.M. 1962-401 (3 slides).

99. *Russellaleyrodes* David

R. cumiugum (Singh)

Philippine Islands: Palawan, Irawan forest, 1000', 19.i.1988, (J.H. Martin 5127), ex: indet., B.M. 1988-2.

Philippine Islands: Palawan, Irawan forest, 1000', 21.i.1988, (J.H. Martin 5148), ex: indet., B.M. 1988-2.

100. *Setaleyrodes* Takahashi

S. mirabilis Takahashi

Taiwan: Wufeng, 24.v.1994, (K.C. Chou), ex: *Stephania japonica*, B.M. 1994 (3 slides).

Taiwan: Wufeng, Taiwan Agric. Res. Institute, 18.v.1994, (K.C. Chou), ex: *Stephania japonica*, B.M. 1994 (2 slides).

S. thretaonai David

Paratypes, India: Kerala, Walayar, 24.iii.1972, (B.V. David), ex: *Homonoia riparia*, 18/85.

Australia: Northern Territory, Humpty Doo, Solar Village, 45 km S. E. of Darwin, 24.xii.1996, (J.H. Martin 6963), ex: *Ficus* sp.

S. vigintiseta Martin

Paratypes, Australia: Northern Territory, Humpty Doo, Solar Village, 45 km S. E. of Darwin, 24.xii.1996, (J.H. Martin 6965), ex: *Acacia ?aulacocarpa*, B.M. 1998-50 (3 slides).

101. *Simplaleurodes* Goux

S. hemisphaerica Goux

Morocco: 45 km S. of Oujda, Col de Jerada, 22.iv.1992, (J.H. Martin 6054), ex: Oleaceae, B.M. 1992-73.

102. *Singhiella* Sampson

S. bicolor (Singh)

India: New Delhi, iv.1964, (T.S. Muthukrishnan), ex: *Eugenia jambolana*, B.M. 1967-280/19 (2 slides).

Pakistan: Rawalpindi, 23.iv.1979, ex: *Eugenia jambolana*, C.I.E. A11232.

S. cardamomi (David & Subramaniam)

West Malaysia: Pahang Prov., Taman Negara, Kuala Tahan, 2.iii.1984, (J.H. Martin 4191), ex: Zingiberaceae (2 slides).

Brunei: Borneo, nr Bukit Bedawan, 30.iii.1989, (J.H. Martin 5516), ex: Zingiberaceae, B.M. 1989-89.

S. chinensis (Takahashi)

Hong Kong: Lantau Island, Shek Pik – Taio Road, 18.x.1990, (J.H. Martin 5798), ex: *Machilus chinensis*.

S. citrifolii (Morgan)

Hong Kong: NT, Tai Lung Farm, Sheung Shui, 22.xi.1999, (J.H. Martin 7250), ex: *Citrus grandis*.

Hong Kong: NT, Tai Lung Farm, Sheung Shui, 22.xi.1999, (J.H. Martin 7252), ex: *Citrus paradisi*.

Hong Kong: HK, Botanical & Zoological Gardens, 7.xii.1999, (J.H. Martin 7319), ex: *Randia spinosa*.

Costa Rica: Turrialba, 13.iii.1990, (J.M. Cox 81), ex: *Citrus* sp.

Costa Rica: San Gerardo, nr San Isidro, 10.ii.1983, (J.H. Martin 3818), ex: *Citrus* sp.

Jamaica: Kingston, 26.x.1970, (L.A. Mound), ex: *Citrus* sp., 31/71.

Lebanon: Byblos, ix.2000, (M.R. Wilson), ex: *Citrus* sp.

USA: Florida, Nursery at Broxbourne, Herts., 19.xi.1974, ex: *Citrus mitis*.

S. delamarei (Cohic)

Chad: Doba, 389 m, 21.vi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*, B.M. 1976-56.

Kenya: Gedi forest, south of Malindi, 17.v.1988, (J.H. Martin 5321), ex: *Monanthotaxis fornicata*, B.M. 1988-188.

S. kuraruensis (Takahashi)

Taiwan: Sanhsian, 22.vi.1990, (C.C. Ko), ex: *Machilus kusanoi*.

S. premmae Martin

Paratypes, Australia: Queensland, strand 2 km north of Cape Tribulation, 13.xii.1996, (J.H. Martin 6862), ex: *Premna serratifolia*, B.M. 1998-50.

S. subrotunda (Takahashi)

Taiwan: Wuling, 11.x.1986, (C.C. Ker), ex: *Litsea akoensis*, B.M. 1987-197.

Taiwan: Fenchifu, 13.ii.1986, (C.C. Ker), ex: *Phoebe formosana*, B.M. 1987-197.

103. *Singhius* Takahashi

S. hibisci (Kotinsky)

Syntypes, Hawaii: Hilo, xi.1905, (B.M. Newell), ex: *Hibiscus roseorinctus*, B.M. 1923-278 (3 slides).

Hong Kong: Victoria Peak, 2.iii.1990, (J.H. Martin 5684), ex: ?*Bridelia monoica*.

Hong Kong: Wanchai, 25.ii.1990, (J.H. Martin 5667), ex: indet. shrub.

Taiwan: Wufeng, 8.xii.1985, (C.C. Ker), ex: *Mallotus japonicus*, B.M. 1987-197.

S. russellae (David & Subramaniam)

Paratypes, India: Valparai, 6000 ft, 16.iv.1967, (B.V. David), ex: indet., B.M. 1972-24.

104. *Siphoninus* Silvestri

S. immaculatus (Heeger)

- England: Colchester, Essex, ii.1973, (E. Milne-Redhead), ex: Ivy, 11/73 (5 slides).
 England: Camberley, Surrey, 25.viii.1920, (E.E. Green), ex: Ivy, B.M. 1930-488 (2 slides).
S. phillyreae (Haliday)
 France: Corsica, Bussaglia, near Porto, 2.vii.1980, (J.H. Martin 3014), ex: *Crataegus* sp.
 France: Corsica, Bussaglia, near Porto, 2.vii.1980, (J.H. Martin 3015), ex: *Fraxinus* sp.
 Hungary: Budapest, Plant Protection Inst., 19.viii.1983, (L.A. Mound), ex: indet.
 Spain: Cadiz Prov., E. of Grazalema, A372, 11.x.1998, (J.H. Martin 7170), ex: *Crataegus* sp.
 Spain: Burgos Prov., Santuario de Santa Casilda, near Briviesca, 950 m, 6.viii.1993, (J.H. Martin 6294), ex: *Crataegus* sp.
 England: Camberley, Surrey, 1.x.1927, (E.E. Green), ex: Hawthorn, B.M. 1930-488.
 USA: California, U. Cal. Riverside campus, 3.vii.1990, (J. LaSalle), ex: *Citrus* sp.

105. *Taiwanaleyrodes* Takahashi

- T. carpini* Takahashi
 Taiwan: Hapan, 17.vi.1994, (K.C. Chou), ex: *Maesa* sp., B.M. 1994 (2 slides).
T. meliosmae Takahashi
 Japan: Kushimoto, Wakayama Prefecture, vi.1955, (R. Takahashi), ex: *Daphniphyllum* sp., 101/62, B.M. 1962-401, (5 slides).

106. *Tetraleyrodes* Cockerell

- T. acaciae* (Quaintance)
 Mexico: Tamaulipas state, Miquihuana, 1.iv.2001, (E. Danzig), ex: *Sophora secundaria* (2 slides).
 Hong Kong: Pok Fu Lam, Country Park, 10.xii.2001, (J.H. Martin 7545), ex: ?Mimosaceae (2 slides).
 Jamaica: Negril, 17.vi.1969, (M. Schuiling 39), ex: *Piscidia piscipula*, 10/69 (2 slides).
T. andropogoni (Dozier)
 Syntypes, Belgian Congo: Lodja, x.1929, (J. Ghesquière), ex: *Andropogon* sp., B.M. 1933.
 Uganda: W. Buganda, Kyadondo, 3900', Kawanda Res. Stn, 25.iii.1971, (D.N. McNutt), ex: *Phragmanthera rufescens* var. *sigensis*, 9/72 (3 slides).
 Belgian Congo: Brazzaville, 23.ii.1966, ex: *Loranthus* sp., Al 263-66.
 Burundi: Rumonge, 3.ix.1983, (G. Remaudière), ex: *Elaeis guineensis*, 46/83.
T. banksiae Martin
 Paratypes, Australia: S.A., Kangaroo Island, 17.ii.1990, (J.H. Martin 5654), ex: *Banksia marginata*, B.M. 1998-50 (3 slides).
T. bicolor Bink-Moenen
 Turkey: Adana, 16.ii.1987, (via N. Uygun), ex: *Myrtus communis*, 13/87, (4 slides).
T. caulicola Nakahara
 Paratypes, USA: California, Palm Canyon, 31.iii.1963, (L.D. Anderson), ex: *Dalea* sp., B.M. 1998-131.
T. confusa Nakahara
 Paratypes, USA: Florida, Ormond Beach, 7.viii.1959, (Smith & Roberts), ex: *Magnolia virginiana*, B.M. 1998-131.
T. granulata Bink-Moenen

Paratypes, Chad: Doba, 21.vi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *gardenia triacantha*, B.M. 1977-471 (2 slides).

Paratypes, Chad: Bebedjia, 11.vii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hymenocardia acida*, B.M. 1977-471.

Paratypes, Chad: Donia, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hexalobus monopetalus*, B.M. 1977-471 (2 slides).

T. pluto Dumbleton

Australia: W.A. Ravensthorpe, 25.ix.1980, (H.J. Banks), ex: *Hakea* sp. (3 slides).

T. quercicola Nakahara

Paratypes, USA: Utah, iv.1966, (G.F. Knowlton), ex: Evergreen oak, B.M. 1998-131.

T. selachidentata Bink-Moenen

Paratypes, Chad: Bebedjia, 26.xi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Stereospermum kunthianum*.

T. stirlingiae Martin

Paratypes, Australia: intercepted at SFKIA, 26.x.1987, (T. Moreno 88 0550), ex: *Stirlingia* sp., B.M. 1998-131.

Paratypes, Australia: intercepted at San Francisco, 25.vii.1988, (A. Mendoza), ex: *Stirlingia latifolia*, B.M. 1998-131.

T. sulcistriatus Martin

Paratypes, Australia: Queensland, strand 2 km north of Cape Tribulation, 15.xii.1996, (J.H. Martin 6887), ex: *Ficus microcarpa*, B.M. 1998-50.

Paratypes, Australia: Queensland, strand 2 km north of Cape Tribulation, 13.xii.1996, (J.H. Martin 6866), ex: *Calophyllum inophyllum*, B.M. 1998-50 (2 slides).

Paratype, Australia: Northern Territory, Kakadu Nat. Park, Gubara Walk, Burdulba Creek, 20 km S. of Jabiru, 22.xii.1996, (J.H. Martin 6954), ex: ?*Xanthostemon eucalyptoides*, B.M. 1998-50.

107. *Tetralicia* Harrison

T. ericae Harrison

Topotypes, England: Durham, Waldrige Fell, (B.J. Selman), ex: *Erica tetralix*, 10/65, (4 slides).

England: Surrey, Camberley, xi.1929, (E.E. Green), ex: *Erica tetralix*, B.M. 1929-545 (3 slides).

T. graminicola Bink-Moenen

Paratypes, Chad: Baibokoum, 8/9.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Hyparrhenia rufa*.

T. iberiaca Bink-Moenen

Paratypes, Portugal: Monchique, ca. 300 m, 29.iv.1978, (F.A. Bink & R.M. Bink-Moenen), ex: *Erica arborea*.

T. tuberculata Bink-Moenen

Holotype+paratypes, Chad: Bebedjia, 395 m, 11.vii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Vitex simplicifolia*.

108. *Trialeurodes* Cockerell

T. abutiloneus (Haldeman)

USA: Arizona, East of Roll, 22.ix.1954, (R.D. Dickson), ex: *Euphorbia* sp., B.M. 1998-131.

USA: Florida, Haven, 2.xi.1959, (V. Norton & F. Weems), ex: *Hibiscus mutabilis*, B.M. 1998-131.

USA: California, Three Rivers Tulore Co., 4.xi.1978, (L. Vet), ex: *Euphorbia supina*.
T. bruneiensis Martin

Paratypes, Brunei: Borneo, near Bukit Bedawan, 30.iii.1989, (J.H. Martin 5523), ex: Fern, B.M. 1989-89, (5 slides).

T. darwiniensis Martin

Paratypes, Australia: Northern Territory, Darwin, Nightcliff, foreshore, 20.xii.1996, (J.H. Martin 6931 & A. Wells), ex: *Flueggea ?virosa*, B.M. 1998-50 (5 slides).

T. dicksoniae Martin

Paratypes, Australia: Victoria, Tall Trees, nr Mansfield, 22.i.1990, (J.H. Martin 5621), ex: *Dicksonia antarctica*, B.M. 1998-50 (4 slides).

T. drewsi Sampson

USA: California, Sacramento, Sac. Co., 27.ii.1968, (P.W. Hiatt), ex: *Quercus ilex*, 9/68, B.M. 1968-368.

T. ericae Bink-Moenen

Paratypes, Netherlands: Leersum, reserve Leersumse Veld, 6.vi.1975, (R.M. Bink-Moenen), ex: *Erica tetralix*, B.M. 1976-56 (2 slides).

Netherlands: Ilpendam, reserve Ilpenveld, 29.v.1975, (F.A. Bink & R.M. Bink-Moenen), ex: *Erica tetralix*, B.M. 1976-56.

Netherlands: Amerongen, 27.iv.1975, (F.A. Bink & R.M. Bink-Moenen), ex: *Erica tetralix*, B.M. 1976-56.

T. floridensis (Quaintance)

USA: Florida, Tampa, 11.ii.1976, (C.W. Hale), ex: *Persea americana*, B.M. 1998-131.

Panama: Canal Zone, Barro Colorado I., 13.iii.1983, (J.H. Martin 3988), ex: *Inga* nr *pezizifera*.

Costa Rica: Guanacaste Province, 60 km S. of Liberia, 26.i.1983, (J.H. Martin 3693), ex: *Mastichodendron ?capiri* (2 slides).

T. pergandei (Quaintance)

USA: Florida, nr Brooksville, 11.ii.1922, (A.L. Quaintance 20590), ex: *Bignonia* sp., B.M. 1998-131.

USA: Maryland, Silver Spring, 1.ix.1941, (L.M. Russell), ex: *Rubus* sp., B.M. 1998-131.

?USA: pres. T.D.A. Cockerell, det. Quaintance, ex: Rose, B.M. 1923-178 (2 slides).

USA: Maryland, Allegany Co., Green Ridge, State Forest, 29.vi.1998, (J.H. Martin 7101), ex: *Crataegus* sp.

T. rex Martin

Paratypes, Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Clark's camp area, 18.iv.1985, (J.H. Martin 4869), ex: Fren.

Paratypes, Indonesia: Sulawesi Utara, Dumoga-Bone N.P., Clark's camp area, 18.iv.1985, (J.H. Martin 4879), ex: Fren (3 slides).

T. vaporariorum (Westwood)

?Type series, England: "on leaf in Westwood's Taxonomic Series", Hope Department, Oxford, no data (3 slides).

Syntypes, South Africa: Durban, Pretoria, Chief Entomologist, vi.1923, ex: *Nicotiana tabacum*, B.M. 1935-248 (2 slides).

England: Berkshire, Ascot, Silwood Park, 5.xi.1978, (J.H. Martin 2323), ex: *Solanum ducamara*, 9/78.

South Africa: GAU, Delmas, v.2000, (R. Tukker HA/60), ex: *Cucumis sativus*, B.M. 2001-198.

- Azores: Pico, hotel grounds, 26.vi.2000, (A. Polaszek), ex: *Oenothera* sp.
 Madeira: Portela, 550 m above Porto da Cruz, 28.iii.1995, (J.H. Martin 6560 & A.F. Aguiar), ex: *Ageratina* sp.
 Iran: Isfahan, University of Technology, viii.1986, ex: *Nicotiana tabacum*, C.I.E. A18545, 2/87.
 Iran: Fars, Badjgah, 13.x.1990, (M. Zarrabi 17), ex: *Cichorium intybus*.
 Italy: Po River Valley, 1998, ex: Strawberry, C.I.E. A20129.
 USA: California, Riverside Co., UCR campus, (L. Vet), ex: *Nicotiana glauca*.
T. variabilis (Quaintance)
 USA: Florida, Miami, 1.xii.1921, (G.F. Moznette), ex: *Carica papaya*, B.M 1998-131.
 Jamaica: Kingston, vii.1965, ex: *Carica papaya*, B.M 1965-433.
 Belize: Cayo, Chiquibul forest, Las Cuevas, 6.xi.1994, (J.H. Martin 6463), ex: *Carica* sp. (2 slides).
 Trinidad: Tableland, 3.xii.1991, I.I.E. 22137, ex: *Carica papaya*.

109. *Tuberaleyrodes* Takahashi

- T. machili* Takahashi
 Hong Kong: Victoria Peak, 3.iii.1990, (J.H. Martin 5693), ex: *Machilus chinensis* (4 slides).

110. *Vasdauidius* Russell

- V. cobarensis* (Martin)
 Paratypes, Australia: New South Wales, Cobar, 1979, ex: Pasture grass, C.I.E. 11292 (2 slides).
V. concursus (Ko)
 Paratypes, Taiwan: Taipei City, 10.xii.1989, (T.C. Hsu), ex: Gramineae.
 Paratypes, Taiwan: Taipei City, 30.i.1994, (K.C. Chou), ex: Gramineae.
 Hong Kong: NT, west Tai Mo Shan, Country Park, Ho Pui area, 13.xii.2001, (J.H. Martin 7566), ex: *Saccharum ?spontaneum*, (2 slides).
 Hong Kong: NT, E. side of Ma On Shan, Country Park, near Hebe Haven, 14.xi.1996, (J.H. Martin 6817), ex: Grass.
V. indicus (David & Subramaniam)
 Paratypes, India: Coimbatore, 20.v.1966, (B.V. David), ex: *Chloris barbata*, B.M. 1972-24 (2 slides).
 Nigeria: Ibadan, I.I.T.A., 20.ii.1983, (M.S. Alam), ex: Rice, C.I.E. A15041.
 Nigeria: Badeggi, 25.x.1977, (S.A. Apeji), ex: Rice, C.I.E. A9972.
 Burkina Faso (Upper Volta): Bo-Bo rice area, 31.vii.1973, (C.L. Tao), ex: *Oryza sativa*, C.I.E. A10811, 64/82 (2 slides).
V. setiferus (Quaintance & Baker)
 Hong Kong: Tai Lung Farm, NT, 14.vii.1976, (R. Winney), ex: Grass, C.I.E. A9278 (2 slides).
 Papua New Guinea: Morobe Prov. coast, Buso, 15.x.1979, (J.H. Martin 2702), ex: *Imperata cylindrica* (3 slides).

111. *Venezaleurodes* Russell

- V. pisoniae* Russell
 Paratypes, Venezuela: Ipore, near Altigracia de Orituco, xi.1966, (F.A. Lee), ex: *Pisonia macranthocarpa*, B.M. 1998-131 (2 slides).

Paratypes, Venezuela: Guarcio, near Altagracia de Orituco, Guatopo, xi.1966, ex: *Pisonia macranthocarpa* (2 slides).

112. *Viennotaleyrodes* Cohic

V. bicolorata Martin

Paratypes, Australia: Queensland, 150 km S. of Mackay, 19.vii.1968, (L.A. Mound), ex: *Acacia harpophylla*, 22/71, (9 slides).

V. bosciae Bink-Moenen

Holotype: Cameroun: 22 km S. of Waza, 23.i.1966, (A.J.M. Leeuwenberg), ex: *Boscia senegalensis*.

Sudan: Bahr el Ghazal Prov., ~20 km N. of Aweil, 9.v.1981, (J.H. Martin 3386), ex: *Boscia senegalensis* (2 slides).

V. curvisetosus Martin

Paratypes, Australia: New South Wales, Tamworth, 29.iii.1920, (W.W. Froggatt), ex: *Acacia pendula* (4 slides).

V. fallax Bink-Moenen

Paratypes, Chad: Boro, 450 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Burkea africana*, B.M. 1976-56 (2 slides).

Paratypes, Chad: Boro, 443 m, 30.vii.1972, (F.A. Bink & R.M. Bink-Moenen), ex: *Burkea africana*, B.M. 1976-56.

V. lacunae Martin

Paratypes, Australia: ACT, Molonglo Gorge (near Queanbeyan, NSW), 8.xii.1996, (J.H. Martin 6849), ex: *Acacia rubida*, B.M. 1998-50 (4 slides).

V. megapapillae (Singh)

Hong Kong: Pok Fu Lam, Country Park, 10.xii.2001, (J.H. Martin 7547), ex: *Millettia* sp., (4 slides).

Taiwan: Hsini, 23.xii.1995, (K.C. Chou), ex: *Bauhinia championii*.

Taiwan: 22.i.1990, (Y.C. Shiau), ex: indet.

V. platysepali (Cohic)

Syntype, Congo: Brazzaville, 5.iii.1965, ex: *Platysepalum vanderystii*, B.M. 1965-305.

Chad: Galaba, 407 m, 2.ix.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Entada africana*, B.M. 1976-56 (2 slides).

Chad: Timberi, 435 m, 29.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Dialium guineense*, B.M. 1976-56.

113. *Xenaleyrodes* Takahashi

X. artocarpi Takahashi

Australia: Queensland, strand 2 km north of Cape Tribulation, 13.xii.1996, (J.H. Martin 6863), ex: *Ficus microcarpa*, B.M. 1998-50 (2 slides).

Australia: Queensland, strand 2 km north of Cape Tribulation, 15.xii.1996, (J.H. Martin 6887), ex: *Ficus microcarpa*, B.M. 1998-50.

Australia: Queensland, creek mouth 2 km north of Cape Tribulation, 15.xii.1996, (J.H. Martin 6885), ex: *Pouteria obovoidea*, B.M. 1998-50.

Australia: Queensland, 2 km north of Cape Tribulation, garden/orchard, 17.xii.1996, (J.H. Martin 6903), ex: *Pouteria caimito*, B.M. 1998-50.

Papua New Guinea: Morobe Prov. coast, Buso, 9.x.1979, (J.H. Martin 2681), ex: *Premna* sp. (2 slides).

X. broughae Martin

Paratypes, Papua New Guinea: S. Highlands Province, Eraue, 21.vii.1983, (E.J. Brough C630), ex: *Citrus* sp., C.I.E. A15579, 37/84, (7 slides).

X. fauceregius Martin

Paratypes, Australia: Northern Territory, King's Canyon, 5.ii.1990, (J.H. Martin 5643), ex: *Acacia ?mcdonnelliensis*, B.M. 1998-50.

Paratypes, Australia: New South Wales, Native Dog Bore, 1890, (W.W. Froggatt), ex: indet. (2 slides).

Australia: South Australia, 2 km N. of Roxby Downs, 25.iv.2000, (J.H. Martin 7397), ex: *Acacia ?ramulosa*.

Australia: Queensland, 40 km E. of St. George, 12.iv.2000, (L.A. Mound 3860), ex: *Acacia aneura*.

114. *Yleyrodes* Bink-Moenen

Y. isoberliniae Bink-Moenen

Paratypes, Chad: Boro, 440 m, 6.xii.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Isoberlinia doka*, B.M. 1977-471.

115. *Zaphanera* Corbett

Z. capparis Bink-Moenen

Paratypes, Chad: Bebedjia, 12.vi.1973, (F.A. Bink & R.M. Bink-Moenen), ex: *Capparis corymbosa*, B.M. 1977-471 (2 slides).

Sudan: Bahr el Ghazal Prov., 160 km S.W. of El Muglad, 7.v.1981, (J.H. Martin 3385), ex: indet. woody plant.

Z. cyanotis Corbett

Lectotype, Ceylon: Maskeliya, v.1911, (J. Pole), ex: *Cyanotis* sp., B.M. 1927-451.

Paralectotype, Ceylon: Maskeliya, v.1911, (J. Pole), ex: *Cyanotis* sp., B.M. 1927-451.

Paralectotypes, Ceylon: Pundaluoya, (E.E. Green), ex: *Cyanotis* sp., B.M. 1927-451 (2 slides).

Pakistan: Balakot, 12.iv.1963, ex: *Dicliptera roxburghii*, C.I.E. 19214, 2/64.

Z. niger (Maskell)

Australia: S.A., Mt Lofty Ranges, Basket Range, 1.i.2000, (J. Hardy), ex: *Acacia* sp. (2 slides).

Australia: S.A., Carey Gully, nr Adelaide, i.2000, (J. Hardy), ex: *Acacia* sp.

Australia: S.A., Adelaide, Crafers, 25.xii.1997, (L.A. Mound & A. Wells), ex: *Acacia melanoxydon* (2 slides).

Australia: A.C.T., Canberra, Nat. Bot. Gdns, Black Mtn, 21.x.1994, (L.A. Mound), ex: *Acacia longifolia*.

Z. papyrocarpae Martin

Paratypes, Australia: S.A., Billakilina Station, 26.iv.2000, (J.H. Martin 7406), ex: *Acacia papyrocarpa* (3 slides).

Paratypes, Australia: S.A., Roxby Downs, 27.iv.2000, (J.H. Martin 7408), ex: *Acacia papyrocarpa* (3 slides).

Paratypes, Australia: S.A., Roxby Downs, 11.i.2000, (J. Hardy), ex: *Acacia papyrocarpa*.

Paratypes, Australia: S.A., Roxby Downs, 20.x.1999, (P. Bailey), ex: *Acacia papyrocarpa*.

Z. rhachisreticulata Martin

Paratypes, Australia: W.A., Wubin, iv.1997, (L.A. Mound), ex: *Acacia mackeyana*, B.M. 1998-51 (4 slides).

116. New genus 1

New genus 1 sp. 1

Belize: Cayo, Chiquibul Forest, Las Cuevas plots, 20.xi.1994, (J.H. Martin 6502), ex: Piperaceae (2 slides).

Belize: Cayo, Chiquibul Forest, New Maria track, 12.ii.1996, (J.H. Martin 6647), ex: indet. (2 slides).

Belize: Cayo, Chiquibul Forest, San Pastor track, 28.xi.1994, (J.H. Martin 6523), ex: *Piper* sp. (3 slides).

Belize: Cayo, Chiquibul FR., Monkey Tail trail, 22.vi.2002, (J.H. Martin 7703), ex: *Piper* sp. (2 slides).

New genus 1 sp. 2

Belize: Cayo, Chiquibul Forest, Las Cuevas plots, 5.xi.1994, (J.H. Martin 6458), ex: *Piper* sp.

Belize: Cayo, Chiquibul Forest, Las Cuevas plots, 20.xi.1994, (J.H. Martin 6502), ex: Piperaceae.

Belize: Cayo, Chiquibul Forest, San Pastor Hill path, 26.ii.1996, (J.H. Martin 6710), ex: *Piper* sp.

Belize: Cayo, Chiquibul FR., Monkey Tail trail, 22.vi.2002, (J.H. Martin 7703), ex: *Piper* sp.

117. New genus 2

New genus 2 sp. 1

Costa Rica: Turrialba, 640 m, Trop. Agric. Centre, C.A.T.I.E., 3.iii.1983, (J.H. Martin 3964), ex: ?Myrtaceae (2 slides).

Costa Rica: Turrialba, 640 m, Trop. Agric. Centre, C.A.T.I.E., 1.iii.1983, (J.H. Martin 3923), ex: *Hoffmannia* sp.

New genus 2 sp. 2

Costa Rica: Cerro, Chirripó Nat. Park, ~2600 m, 18.ii.1983, (J.H. Martin 3882), ex: indet. (2 slides).

Aleurodicinae

1. *Aleurodicus* Douglas

A. cocois (Curtis)

Paralectotypes, Barbados: (Sir R. Schomburgk), ex: Coconut tree (3 slides).

A. dispersus Russell

Paratypes, USA: Florida, Key West, 12.vi.1964, (H.V. Weems), ex: Coconut Palm (3 slides).

Hawaiian Islands: Oahu, Bishop, Museum grounds, 30.iii.1999, (J.H. Martin 7194), ex: *Colocasia esculenta*.

Hawaiian Islands: Hawai'i, coast, Kilauea Volcano, Pu'u O'o lava edge, 6.iv.1999, (J.H. Martin 7209), ex: indet. shrub.

Hawaiian Islands: Maui, Kailua, Hana route, Waianapanapa, 1.iv.1999, (J.H. Martin 7197), ex: *Hibiscus tiliaceus*.

Panama: Canal Zone, Cerro Galera, 26.iii.1983, (J.H. Martin 4144), ex: *Acalypha diversifolia*.

Panama: Darién, Ensenada del Guayabo, sea level - 250 m., 11.i.1983, (J.H. Martin 3582), ex: Leguminosae.

2. *Azuraleurodicus* Martin

A. pentarthrus Martin

Paratype, Belize: Cayo, Chiquibul forest, Grano de Oro track, 22.xi.1994, (J.H. Martin 6486), ex: *Guettarda ?combesii*.

Paratypes, Belize: Cayo, Chiquibul forest, Grano de Oro track, 16.xi.1994, (J.H. Martin 6486), ex: *Guettarda ?combesii*.

Paratypes, Belize: Cayo, Chiquibul For. Res., near Caracol Jct., 16.ii.1996, (J.H. Martin 6630A), ex: *Guettarda combesii* (2 slides).

3. *Dialeurodicus* Cockerell

D. cockerellii (Quaintance)

Syntypes, Brazil: Campinas, São Paulo, 30.iii.1898, (F. Noak, via Cockerell), ex: Myrtaceae, B.M. 1998-131.

Brazil: São Paulo, Embu-Guaco, 29.v.-2.vi.1998, (A.E. Campos-Farinha), ex: *Psidium* sp., Lot #9805640, B.M. 1998-168.

4. *Eudialeurodicus* Quaintance & Baker

E. bodkini Quaintance & Baker

Syntypes, Guyana: Berbice, 2.iii.1915, (G.E. Bodkin), ex: *Erythrina glauca*, B.M. 1998-131 (2 slides).

5. *Octaleurodicus* Hempel

O. nitidus Hempel

?Syntypes, Brazil: Bahia, 7.xii.1923, (G. Bondar), ex: *Cocos nucifera*, B.M. 1924-44.

6. *Stenaleyrodes* Takahashi

S. vinsoni Takahashi

Paratype, New Caledonia: Noumea, 20.v.1955, (L.J. Dumbleton), ex: *Cocos nucifera*, B.M. 1964-721.

Réunion Island: Saint Denis, 14.x.1963, (J.G. Pointel), ex: Palm, B.M. 1998-131.

7. *Synaleurodicus* Solomon

S. hakeae Solomon

Lectotype, Australia: Western Australia, Perth, iii.1934, (M.E. Solomon), ex: *Hakea prostrata*, B.M. 1936-84.

Australia: Western Australia, Perth, King's Park, 26.ix.1967, (L.A. Mound 215), ex: *Hakea ?pritzelli* (4 slides).

Appendix D: data matrix used in the cladistic analysis of the subfamily Aleoerodinae

Note: polymorphisms are coded as follows: A = 0&1, B = 0&2, C = 0&3, D = 0&6, E = 0&7, F = 1&2, G = 1&3, H = 1&8, K = 2&3, M = 2&5, N = 2&7, P = 0&1&2.

In-groups	1	2	3	4	5	6	7	8	9
<i>Acanthaleyrodes calliicarpa</i>	1234567890112345678901234567890123456789012345678901234567890123456789012345678901234								
<i>Acanthaleyrodes spinifera</i>	1?0111010101010?0?0?120000?10005?2?111231021?000?5?1101010010000000?01010010000000000000101000								
<i>Acanthaleyrodes styraci</i>	1?0111001010102013110?0?0000?00005?2?1110233021?000?5010111001000000?0101001000000000000101000								
<i>Acanthobemisia</i>	2001101110110012120?0?0?1110?000020?000111031?201011100?0?110000000?01010000000000000000101000								
<i>Acaudaleyrodes africana</i>	2021110000?11121100?0?0210?01004?2?110002031?101001000?000B110000?0?111001000000000000101000								
<i>Acaudaleyrodes raphipora</i>	2201110000?1111100?0?0210?01004?2?110002031?101001000?00021100000?0?1110010A001000000000101100								
<i>Acaudaleyrodes tuberculata</i>	2021110000?1111100?0?0210?01004?2?110002031?101001000?0?1100000?00101100000000000000101100								
<i>Acutaleyrodes</i>	2001110110?101110?00?00001?1012116031?2A104100?0010000000?010100000000000000000000101000								
<i>Africaleurodes coffeacola</i>	1?01110000?1010011A0?0?01013000025?010211031?20104110101100000000?01010000000000000000101000								
<i>Africaleurodes fulakariensis</i>	1?01110000?10100100?0?01013000020?010211031?20104110101100000000?01010000000000000000101000								
<i>Africaleurodes hexalobi</i>	1?01110000?10100100?0?01013000020?010211031?20104110101100000000?01010000000000000000101000								
<i>Africaleurodes loganiaceae</i>	1?01110000?10100100?0?01013000021?0102115031?200?6110101100000000?01010000000000000000101000								
<i>Africaleurodes ochnaceae</i>	1?01110000?10100100?0?01013000021?010211031?20104110101100000000?01010000000000000000101000								
<i>Agrostaleyrodes</i>	1?01110010?101020?0?0?0010?00105?000211021?000?00001110101000000?01010000000000000000101001								
<i>Aleurocanthus banksiae</i>	202111000111102100?0?101310?00005?2?11023300?0?00?01100?0000110000?01001000010000000000101000								
<i>Aleurocanthus calophylli</i>	230111010111102100?0?110200?001021?10023100?0?00?00100?0000100000?10100000000000000000101000								
<i>Aleurocanthus cerarocoeus</i>	20011000110112100?0?130210?00005?2?10023300?0?00?0000?0000000000?10010000000000000000101000								
<i>Aleurocanthus citripedus</i>	222110101110102100?1011210?00005?2?110031021?000?01100?0000100000?00000001000000000000101000								
<i>Aleurocanthus coccis</i>	22011010111112100?10110210?00105?2?110231021?011001100?0000100000?0000000000010100A0								
<i>Aleurocanthus delottoi</i>	22011010111112100?0?110310?00001?2?11013300?0?00?00100?0000110000?010000100000000000101000								
<i>Aleurocanthus esakii</i>	2021100011112110?0?11310?00005?2?11013300?0?00?0000?0000100000?10000000000000000000101000								
<i>Aleurocanthus gateri</i>	23011000110112100?0?11310?00105?2?100131021?001000100?0000100000?10000000000000000000101100								
<i>Aleurocanthus gordoniae</i>	2301100110112100?0?11210?00105?2?11003300?0?00?0000?0000100000?00000000000100000000101000								
<i>Aleurocanthus incertus</i>	2301101111112100?0?11110?00101?2?110231021?000?01000?00001100000?000000000100100000000101000								
<i>Aleurocanthus leptadeniae</i>	22011010111112100?0?11310?00005?2?100231021?000?01100?0000100000?0000?010000000010000000101100								
<i>Aleurocanthus spiniferus</i>	220110111110102100?1010310?00005?2?110133021?00100000?0000100000?0100100000100100000000101000								
<i>Aleurocerus cerifer</i>	210110?010?11122110?0?0211200000?2?10023400?0?00?01100?00101010000?10100000100000000000101000								
<i>Aleurocerus flavomarginatus</i>	220110?010?11122110?0?0211200000?2?10023400?0?00?01100?00101010000?10100000100000000000101000								
<i>Aleurocerus musae</i>	220110?010?11122110?0?0211200000?2?10023400?0?00?01100?00101010000?1010000010010000000101000								
<i>Aleurocerus palmae</i>	220110?010?11122110?0?0211200000?2?10023400?0?00?01100?00101010000?1010000010010000000101000								
<i>Aleurochiton acerinus</i>	202111000101010B1A100?0?1410?00003?1000235031?11100000111010000000?01010000000000000000101000								
<i>Aleurochiton aceris</i>	2021110000?10100100?0?1410?00003?1000235031?11100000111010000000?0101000000000000000000101000								
<i>Aleurochiton forbesii</i>	2121110000?10100100?0?1110?000020?010211031?20100000?001000000?0100000010?0101000000000000101000								
<i>Aleurochiton pseudoplatani</i>	2021110001010100100?0?1110?00003?1000235031?111000001110101000000?01010000000000000000101000								

Appendix D. (Continued).

	1	2	3	4	5	6	7	8	9
	12345678901234567890123456789012345678901234567890123456789012345678901234								
<i>Aleuromarginatus corbettiaformis</i>	2100110111011000?	100?0?1110?000021?	7010215031?	22010011013111?	20000000?	70101101000001000000000101001			
<i>Aleuromarginatus dalbergiae</i>	2000110111011000?	140?0?1110?001021?	7010215031?	22010010010110010000000?	7010110000000?	70101100000010000000000101001			
<i>Aleuromarginatus kallarensis</i>	2000110111011000?	140?0?1110?001021?	7010215031?	2200?01101010010000000?	70101100000010000000000101001				
<i>Aleuromarginatus littoralis</i>	20001101110101000?	140?0?1110?000024?	7010215031?	220101110101100000000?	701010000001000000000101001				
<i>Aleuromarginatus marginiquus</i>	2000110111011200?	100?0?1110?000021?	7010215031?	7000?00001010100000000?	701010000000000000000001001				
<i>Aleuromarginatus moundi</i>	2000110111011100?	140?0?1110?000021?	7011215031?	701001100?01020000000?	70101110000100000000000101001				
<i>Aleuromarginatus nemciae</i>	1?001101110101200?	100?0?1110?000021?	7000225031?	22010000000?	70110000000000000000000001001				
<i>Aleuromarginatus nigrus</i>	2000110111011100?	140?0?1110?001021?	7010215031?	22010000000?	7011010000000?	70101101000000000000000101001			
<i>Aleuromarginatus serdangensis</i>	2001110010101000?	100?0?0000?	7000121?	70023300???	210000?00100000000?	701010000001000000000101000			
<i>Aleuromarginatus tephrosiae</i>	2100110111011000?	140?0?1110?000021?	7010215031?	22010000013A100000000?	70111001000010000000000001001				
<i>Aleuropapillatus</i>	1?0111011011001110?	0001110?0?113000020?	7010127031?	2201000001010100000100?	701010000100000000000001010A0				
<i>Aleuoparadoxus arctostaphyli</i>	21071110010101001?	0?0?0?0110?00005?	7011037011?	401041001010100000100?	70101000010000000000000101000				
<i>Aleuoparadoxus gardeniae</i>	2101110001011011?	0?0?0?0110?00005?	7011037011?	401061101010100000100?	7010101110000100000000000101000				
<i>Aleuoparadoxus ilicicola</i>	200111001011001?	0?0?0?0110?00005?	7011037011?	401061001010100000100?	701011100010000000000101000				
<i>Aleuoparadoxus iridescens</i>	200111001011001?	0?0?0?0110?00005?	7010A037011?	4010611010101010000100?	70101010100010000000000101000				
<i>Aleuoparadoxus punctatus</i>	2101110001011001?	0?0?0?0110?00105?	7010137011?	40104110101010000100?	70101011100000000000000101000				
<i>Aleuoplatus acaciae</i>	2021110000?121100?	100?0?0110?000021?	7100023031?	2211001000?00100000000?	70001000010000000000000101000				
<i>Aleuoplatus berbericolus</i>	2201110000?111000?	0?0?0?0210?00105?	711002300???	700?41000?00001100000?	7010100001001000000000000101000				
<i>Aleuoplatus bossi</i>	2021110000?111010?	100?0?1110?00005?	71000330B1?	001001100?000000000?	700100001001000000000101100				
<i>Aleuoplatus cadabae</i>	2001110000?101020?	0?0?0?0110?000021?	710023700???	700?0?1100?00000000000?	70010000010010000000000101100				
<i>Aleuoplatus cockerelli</i>	210110?000?101000?	0?0?0?0100?0?0?0?00005?	710013300???	700?0?1100?00001100000?	701100000000000000000101000				
<i>Aleuoplatus coccolus</i>	2201110110?11010?	0?0?0?0210?00005?	711013300???	700?0?1100?00000000000?	701100001001000000000101000				
<i>Aleuoplatus dorsipallidus</i>	2201110110?101020?	0?0?0?0510?00005?	710023300???	700?0?1100?00011000000?	70011001000001000000000101000				
<i>Aleuoplatus epigaeae</i>	2101110000?1112100?	0?0?0?0210?00005?	710112300???	700?0?31100?00101000000?	701010000100100000000101000				
<i>Aleuoplatus evodiae</i>	230110000?1111?100?	0?0?0?0210?00005?	710013300???	700?0?1100?00001000000?	70111001000010000000000101000				
<i>Aleuoplatus fici</i>	1?011100010101020?	100?0?0110?000021?	700023502?	0010111010110000000?	701010000000000000000101000				
<i>Aleuoplatus ilicis</i>	2201110110?11211?	0?0?0?0210?00005?	710011300???	700?0?11000?0000100000000?	701010000100100000000101000				
<i>Aleuoplatus incisus</i>	220?110000?101070?	0?0?0?0510?00005?	7110113300???	700?61000?00?010000000?	70111000000100000000010?700				
<i>Aleuoplatus pectiniferus</i>	2201110000?101000?	100?0?0110?000021?	701121300???	700?0?41100?000000000?	701110000010000000000101000				
<i>Aleuoplatus periplocae</i>	220110010?111000?	0?0?0?010?00005?	7100231021?	010?00000?00000000000?	7001000000000000000000010?7??				
<i>Aleuoplatus perseaphagus</i>	200110?000?11020?	0?0?0?0210?00005?	710013300???	700?0?1100?00001000000?	70011000000000000000000101000				
<i>Aleuoplatus quercusagmaticae</i>	200110?000?11100?	0?0?0?0210?00006?	711013300???	700?0?10000?00100000000?	70111000010000000000000101000				
<i>Aleuoplueroceus</i>	202?10?110?111100?	0?0?0?0110?00005?	710023300???	700?0?00000?00101000000?	70101000100000000000010?000				
<i>Aleuopteridis eastopi</i>	2021110010?10102100?	0?0?0?0100?00005?	7110133011?	7010140000?00100000000?	701010000001000000000101000				
<i>Aleuopteridis filicicola</i>	2021110010?1010F10100?	0?0?0?0100?00005?	7110233011?	7010141100?00100000000?	701010000001000000000101000				

Appendix D. (Continued).

	1	2	3	4	5	6	7	8	9
	12345678901234567890123456789012345678901234567890123456789012345678901234								
<i>Aleuroviggianus adrianae</i>	2021110001011010?	100?0?1110?	0000024?010123021?	000?00001110011000000?00100010000000000000000101000					
<i>Aleuroviggianus graecus</i>	21011101101000?0?0?	21110?000020?000111031?2A0?	0000111031?2A0?	000011110000000?0010000000000000000101000					
<i>Aleuroviggianus halperini</i>	20211100010100100?0?0?	1110?000024?010111031?200?	400010111000000?00010010000000000000000101000						
<i>Aleuroviggianus polymorphus</i>	2021110001010AA0A10?0?1110?	000024?010111031?200?	01010111A000000?000100A000000000000000010100A						
<i>Aleuroviggianus zonalis</i>	20B1110000?101000?0?0?	1110?000020?010211031?200?	111010111000000?0010000000000000000101000						
<i>Aleyrodes amnicola</i>	1?0111000101100100?0?0?	0000?000020?010111031?201001100?00101000000?010100000000000000000101000							
<i>Aleyrodes elevatus</i>	2001110001010210110?0?0?	0000?000020?010111031?20100000101001100000?010100100000000000010100A							
<i>Aleyrodes loniceræ</i>	1?0111000101A0A10A00?0?0?	0000?000020?010115031?201000101010111000000?001001000000000000000101000							
<i>Aleyrodes proleteria</i>	1?011100010100100?0?0?	0000?000021?0000115031?201000101010100000?001000000000000000000101000							
<i>Aleyrodes prunosus</i>	1?011100010100100?0?0?	0000?000021?010111031?201000101010100000?001000000000000000000101000							
<i>Aleyrodes singularis</i>	1?0111000101010100?0?0?	0000?00005?1110111031?201000100?0010100000?00100000000000000000010100A							
<i>Aleyrodiella</i>	2101110000?101000?0?0?	007013000021?00023?011?200?	00001010101000100?0101000100?0101001000000000000101000						
<i>Apobemisia</i>	2021110000?101010?1110?	0?0?1110?000024?000211031?21110000011111000000?000100100000000000000101000							
<i>Arachnaleyrodes</i>	2001110000?1112010100?0?0?	0410?00001?211013300?0?00?41100?0010000001001010000000000000000101100							
<i>Asialeyrodes lushanensis</i>	0?0111000?111000?100?	0?01013000021?00023?00?0?00?511010100000000?001010000000000000000101000							
<i>Astaleyrodes maesae</i>	0?0111000101100100?0?0?	01011000021?111023?00?0?00?61100?0010100000?011000100000000000000101000							
<i>Asterobemisia carpini</i>	1?011100010101000?100?	0?0700?000020?010111031?201011100?111A000000?00101001000000000000101000							
<i>Asterobemisia dentata</i>	1?011100010100100?0?0?	000023?01011103300?010111100?111000000?001010010000000000000101000							
<i>Asterobemisia obenbergeri</i>	2101100010101000?100?	0?0700?000020?01011103300?010111010101000000?0010010000000000000101000							
<i>Asterochiton cordiae</i>	1?01110001010010110?0?0?	01113000020?010117021?00101010010110000000?00100000000000000001010A0							
<i>Asterochiton pittospori</i>	1?0110001010101110?0?0?	0110?000020?010211031?70105110100000000?001010000000000000000101000							
<i>Bellitudo hispaniolae</i>	2101110001011020?0?0?	01113000021?011123?011?201061101010000000?00100000000110000010100?							
<i>Bellitudo jamaicae</i>	2101110001011010?0?0?	01113000021?01123?011?401061101010000000?0010000000001100000101000							
<i>Bemisaaleyrodes brideliae</i>	1?0110001010014140?0?0?	0081021103300?010410101010000000?00100100000000000000000101010							
<i>Bemisaaleyrodes pauliani</i>	1?01110001010014140?0?0?	00900?001022?010211031?201104110101000000?0010000000000000000101010							
<i>Bemisia afer</i>	1?01110001010000?1A0?	0?0010?000025?01011103300?0101110111A000000?00100A000000000000000101000							
<i>Bemisia antennata</i>	1?01110001010000?1A0?	0?0010?000025?01011103300?0101110111000000?001001000000000000000101000							
<i>Bemisia berbericola</i>	1?0111000101010?110?	0?010?000020?0101110331?201000101011000000?0010000000000000000101000							
<i>Bemisia combreticula</i>	1?0111000101020?1A0?	0?0100?000020?010117031?20100110111000000?001001000000000000000101000							
<i>Bemisia formosana</i>	1?0111000101010?110?	0?0110?001023?01021703300?0101110111000000?0010010000000000000101001							
<i>Bemisia giffardi</i>	1?0111000101000?0?0?	010?000025?01021703300?01011101011000000?001001000000000000000101000							
<i>Bemisia gigantea</i>	1?0111000101000?0?0?	010?000025?01021703300?01011101011000000?001001000000000000000101000							
<i>Bemisia guieriae</i>	1?01110111011000?110?	0?010?000025?01001703300?01001001001111000000?001001000000000000000101000							
<i>Bemisia hirta</i>	0?21110001011010?110?	0?010?000020?0101110331?20100000101111000000?001001000000000000000101000							
<i>Bemisia lauracea</i>	2001110001010010100?0?0?	0110?0000125?010217031?20101110101110000000?01111001000000000000000101000							

Appendix D. (Continued).

	1	2	3	4	5	6	7	8	9																										
<i>Orchamoplatus porosus</i>	12345678901234567890123456789012345678901234567890123456789012345678901234	1?0111000110112100?0?0?200?000015??1?0127?31?200?80000?001000000011100100000000000000100101000	1?01110000?1112110?0?04011000021?10013700??00?4110100110000000?1001000010000000000001010A0	220111000A1101100?0?070210?00005??1A023300??00?11100?00010000000?10010010000000000000101000	0?2010?000?10102110?0?1110?00005??11023300??00?0A00?01100000100?0101010000000000000001001	2001110000?10100100?0?0?0110?000022?0002110300?0101110101110000000?010100000000000000010100?	200010?001010101100?0?0?1110?000021?000221031??2A0?2110101100000000?10100000000000000000101001	2001110000?10100100?0?0?1110?00002?000211031?20100A1010110000000?0101000000000000000101000	2101110000?10100100?0?0?1110?000020?0111031?201010A101110000000?0101000000000000000101000	210111000010101100?0?0?1110?000024?01011031?20101010101110000000?0101000000000000000101000	2301110110?11100?0?0?01011000021?10023700??00?41100?001000000?10010000100100000000101000	2121110110?111020?0?0?0110?00005??10022302?1000?00000?0110000010?01010011000000000000101000	1?2111110?111020?0?0?0110?00005??10022302?100100000?0110000010?01010110000000000000101000	2201110000?10101100?0?0?0110?00005??000111031?7A1001101111000000?0101000000000000000101000	200111000010102100?0?0?0110?00005??010111031?70101000011110000000?0101000000000000000101000	2001110000?101020?110?0?110?00003?2000211031?70100000110000000?01010010000000000000101000	2201110000?10101100?0?0?0110?00002?00011031??20100000121110000000?0101000000000000000101000	1?B111000A?101011A10?0?1110?00003?200225031?7110110110110110000000?010100A0000000000000101000	2101110001012100100?0?070110?00005??000102031?101011100?000000000?0101000000000000000101000	F0B11100010110210110?0?0110?00003?2000225031?7A0?1000110110000000?010100000000000000010100?	1?0111000101102110?0?0?0110?00005??100111031?101001100?010000000?0101000000000000000101000	2001110000?1010F1A0?0?1110?00003?2000225031?7A100110111000000100?0101100000000000000101000	2201110000?1010F1A0?0?1110?00002?01011031?2010000011110000000?0101000000000000000101000	20011001010102100?0?070210?00005??10023300??00?41100?000000000?10100000000000000101000	2001110000?12100100?0?0210?00001??11023300??00?41100?00000000?101000000000000000101000	200111000010102110?0?0110?00005??1023300??00?01100?0010000000?0101000000100000000101000	200111000010102110?0?0110?00005??1023300??00?01100?0010000000?0101000000100000000101000	1?2111000101001110?0?0110?00005??010233021?000?5000121010000010?01010010000000000000?????	1?211100010100140?0?0100?000021?0102213031?20105000101010000010?011100100000000000000101000	200110?001110100100?0?0410?00005??11003300??00?01000?0010000000?01010000001000000010?01	1?2110?000?1000?110?0?1410?000021?110121031?201000101010000000?01010000000000000010?01	1?01110000?11001110?0?0210?010024?01142300??00?5110110001100000?0101000000000000001000101000	1?011101110101121K0?0?0010?00105??10123300??00?5110101000100000?00010000000000000101010	1?01110000?11000K130?0?0210?010021?100231021?010?5110110001100000?010100000000000000101010	0?011101110101001K130?0?0210?01001??11123300??00?51101110001100000?010100000000000000101010

Appendix D. (Continued).

	1	2	3	4	5	6	7	8	9
	123456789012345678901234567890123456789012345678901234567890123456789012345678901234								
<i>Xenaleyrodes broughae</i>	2421110001110112100?120?0600?00015??11023300??00?00000?00100000000?01010000100100000000101000								
<i>Xenaleyrodes fauceregus</i>	2020110001111102111112140600?00015??110221021?000?00000?00101000000?10110000100000000000101000								
<i>Yleyrodes</i>	2001110001110100100?0?0?11111000020?000211031?201011101000100000000?01010000000000000000101000								
<i>Zaphanera capparidis</i>	2101110000?10102110?0?0?01A0?00005??10023300??00?01100?00101000000?01010000100100000000101000								
<i>Zaphanera cyanotis</i>	2101110000?10102110?0?0?0210?00005??10023300??00?01100?00101000000?01010000100100000000101100								
<i>Zaphanera niger</i>	200110?000?11100130?0?0?1410?00005??110233011?200?01100?00100000000?01010010000000000000101010								
<i>Zaphanera papyrocarpae</i>	200110?000?1110013130?0?1410?00005??110233011?200?01101310101000000?01010010000000000000101011								
<i>Zaphanera rhachisreticulata</i>	250110?000?11100130?0?0?1410?00005??110223011?200?01100?00101000000?010100001001000000100101010								
New genus 1 sp 1	2621110000?1110B0?0?0?0610?00005??10023300??01001100?00101000000?01010000000000000000101000								
New genus 1 sp 2	2121110000?111020?0?0?0610?00005??10023300??01011100?00101000000?01010010000000000000101000								
New genus 2 sp 1	1?011100011101021G130?0?0110?000024?11123300??01051101010010000000?01010010000000000000101010								
New genus 2 sp 2	1?211100011101021G130?0?0110?00005??10023300??00?051101010010000010?0101001000000000000010??10								
Out-groups									
<i>Stenaleyrodes</i>	2001110110?1211010100?0?0100?000020?0000181300?01140000?00100000100?10100000000100000000001011								
<i>Dialeurodicus</i>	0?01110110?1110010100?0?0010?000021?000215131?501140000?00100000000?1010000000000000000011011								
<i>Aleurodicus cocois</i>	0?0010?110?1210110100?0?0110?00005??0002151302?11100000?01100000000?10100010001000000000011010								
<i>Aleurodicus dispersus</i>	0?0010?110?12101100?0?0?0100?000021?0002151302?11100000?01000000000?10100000001000000000011010								
<i>Azuraleurodicus</i>	0?0110?000?1210A100?0?0?1110?00003?0000216131?511100000?01100000000?00100000001000000000011010								
<i>Eudialeurodicus</i>	1?01110110?1010010100?0?0000?00006??100228031?801000000?01000000000?01110010001000000000011011								
<i>Octaleurodicus</i>	1?0110?110?1110010100?0?0000?000024?010215131?201100000?00100000000?0111000000110000000001101?								
<i>Synaleurodicus</i>	1?0110?000?12100120?0?0?0?0?00005??110218131?2A0?00000?00100000100?1000000000100000000001100?								

For monobasic genera and/or those genera, for which only one species were available for study, the names of species were deleted in the data matrix in order to be indicated in the obtained trees from the other genera with multiple representative species. See Appendix C for the species names of these genera.

Appendix E: host plant association of the whitefly species selected for the phylogenetic study

Note: an asterisk indicates families unplaced to order but included in higher groups. All botanical names follow the system of the Angiosperm Phylogeny Group (Bremer et al. 1998, 2003; Soltis et al. 2000). Pteridophyta (ferns) are unplaced to higher group (§ shows orders of Pteridophyta). ‡ shows the only gymnosperm record (unplaced to higher group). Bold taxa are out-groups.

Species	Host plant family	Host plant order	Host plant higher group
<i>Acanthaleyrodes callicarpae</i>	Euphorbiaceae, Rosaceae, Verbenaceae, Vitaceae*	Lamiales, Malpighiales, Rosales	euasterids I, eurosids I, rosids
<i>Acanthaleyrodes spiniferosa</i>	Ebenaceae	Ericales	asterids
<i>Acanthaleyrodes styraci</i>	Styracaceae	Ericales	asterids
<i>Acanthobemisia distylli</i>	Hamamelidaceae	Saxifragales	core eudicots
<i>Acaudaleyrodes africana</i>	Fabaceae	Fabales	eurosids I
<i>Acaudaleyrodes rachipora</i>	Euphorbiaceae, Fabaceae	Fabales, Malpighiales	eurosids I
<i>Acaudaleyrodes tuberculata</i>	Combretaceae, Euphorbiaceae	Malpighiales, Myrtales	eurosids I, rosids
<i>Acutaleyrodes palmae</i>	Areceaceae	Arecales	commelinids
<i>Africaleurodes coffeacola</i>	Combretaceae, Euphorbiaceae, Malvaceae, Rhamnaceae, Rubiaceae, Sapindaceae	Gentianales, Malpighiales, Malvales, Myrtales, Rosales, Sapindales	euasterids I, eurosids I, eurosids II, rosids
<i>Africaleurodes fulakariensis</i>	Moraceae	Rosales	eurosids I
<i>Africaleurodes hexalobi</i>	Annonaceae, Apocynaceae	Gentianales, Magnoliales	euasterids I, magnoliids
<i>Africaleurodes loganiaceae</i>	Icacinaceae*, Loganiaceae, Malvaceae, Pandaceae, Rubiaceae, Salicaceae	Gentianales, Malpighiales, Malvales	euasterids I, eurosids I, eurosids II
<i>Africaleurodes ochmaceae</i>	Malvaceae, Ochnaceae	Malpighiales, Malvales	eurosids I, eurosids II
<i>Agrostaleyrodes arcanus</i>	Poaceae	Poales	commelinids
<i>Aleurocanthus banksiae</i>	Myrtaceae, Proteaceae	Myrtales, Proteales	eudicots, rosids
<i>Aleurocanthus calophylli</i>	Clusiaceae	Malpighiales	eurosids I
<i>Aleurocanthus ceracroceus</i>	Lauraceae, Monimiaceae, Myrtaceae	Laurales, Myrtales	magnoliids, rosids
<i>Aleurocanthus citriperdus</i>	Rutaceae	Sapindales	eurosids II
<i>Aleurocanthus cocois</i>	Areceaceae, Lauraceae, Myrtaceae	Arecales, Laurales, Myrtales	commelinids, magnoliids, rosids
<i>Aleurocanthus delottoi</i>	Rutaceae, Ulmaceae	Sapindales, Rosales	eurosids I, eurosids II
<i>Aleurocanthus esakii</i>	Rosaceae	Rosales	eurosids I
<i>Aleurocanthus gateri</i>	Areceaceae	Arecales	commelinids
<i>Aleurocanthus gordoniae</i>	Theaceae	Ericales	asterids
<i>Aleurocanthus inceratus</i>	Magnoliaceae, Rutaceae	Magnoliales, Sapindales	eurosids II, magnoliids
<i>Aleurocanthus leptadeniae</i>	Apocynaceae, Brassicaceae, Zygophyllaceae*	Brassicales, Gentianales	euasterids I, eurosids I, eurosids II
<i>Aleurocanthus spiniferus</i>	Annonaceae, Convolvulaceae, Ebenaceae, Elaeocarpaceae, Euphorbiaceae, Hamamelidaceae, Lardizabalaceae, Rosaceae, Rutaceae, Sabiaceae*, Salicaceae, Vitaceae*	Ericales, Magnoliales, Malpighiales, Oxalidales, Ranunculales, Rosales, Sapindales, Saxifragales, Solanales	asterids, core eudicots, euasterids I, eudicots, eurosids I, eurosids II, magnoliids, rosids
<i>Aleurocerus cerifer</i>	Dilleniaceae*, Sapotaceae	Ericales	asterids, core eudicots
<i>Aleurocerus flavomarginatus</i>	Areceaceae, Musaceae	Arecales, Zingiberales	commelinids
<i>Aleurocerus musae</i>	Bignoniaceae, Rubiaceae, Zingiberaceae	Gentianales, Lamiales, Zingiberales	euasterids I, commelinids
<i>Aleurocerus palmae</i>	Areceaceae, Sapotaceae	Arecales, Ericales	asterids, commelinids
<i>Aleurochiton acerinus</i>	Sapindaceae	Sapindales	eurosids II
<i>Aleurochiton aceris</i>	Sapindaceae	Sapindales	eurosids II
<i>Aleurochiton forbesii</i>	Sapindaceae	Sapindales	eurosids II
<i>Aleurochiton pseudoplatani</i>	Sapindaceae	Sapindales	eurosids II
<i>Aleuroclava aucubae</i>	Aquifoliaceae, Araliaceae, Caprifoliaceae, Cornaceae, Juglandaceae, Lauraceae, Moraceae, Oleaceae, Pittosporaceae, Rosaceae, Rubiaceae, Rutaceae, Salicaceae, Theaceae, Ulmaceae	Apiales, Aquifoliales, Cornales, Dipsacales, Ericales, Fagales, Gentianales, Lamiales, Laurales, Malpighiales, Rosales, Sapindales	asterids, euasterids I, euasterids II, eurosids I, eurosids II, magnoliids
<i>Aleuroclava bifurcata</i>	Sapindaceae	Sapindales	eurosids II
<i>Aleuroclava citrifolii</i>	Rutaceae	Sapindales	eurosids II

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Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Aleuroclava euryae</i>	Aquifoliaceae, Ericaceae, Lauraceae, Magnoliaceae, Theaceae, Trochodendraceae	Aquifoliales, Ericales, Laurales, Magnoliales	asterids, euasterids II, magnoliids
<i>Aleuroclava ficicola</i>	Moraceae	Rosales	eurosid I
<i>Aleuroclava filamentosa</i>	Euphorbiaceae, Rubiaceae	Gentianales, Malpighiales	euasterids I, eurosid I
<i>Aleuroclava hikosanensis</i>	Aquifoliaceae, Buxaceae, Lauraceae, Pittosporaceae, Theaceae	Apiales, Aquifoliales, Ericales, Laurales	asterids, euasterids II, magnoliids
<i>Aleuroclava jasmini</i>	Combretaceae, Euphorbiaceae, Myrsinaceae, Oleaceae, Rubiaceae, Rutaceae	Ericales, Gentianales, Lamiales, Malpighiales, Myrtales, Sapindales	asterids, euasterids I, eurosid I, eurosid II, rosids
<i>Aleuroclava magnoliae</i>	Actinidiaceae, Betulaceae, Clethraceae, Cornaceae, Ericaceae, Fabaceae, Hamamelidaceae, Lauraceae, Magnoliaceae, Oleaceae, Rosaceae, Sapindaceae, Saxifragaceae	Cornales, Ericales, Fabales, Fagales, Lamiales, Laurales, Magnoliales, Rosales, Sapindales, Saxifragales	asterids, core eudicots, euasterids I, eurosid I, eurosid II, magnoliids
<i>Aleuroclava murrayae</i>	Lauraceae, Proteaceae, Rutaceae	Laurales, Proteales, Sapindales	eudicots, eurosid II, magnoliids
<i>Aleuroclava nigeriae</i>	Anacardiaceae, Bignoniaceae, Combretaceae, Ebenaceae, Loganiaceae, Moraceae, Myrtaceae, Rubiaceae, Salicaceae, Sapindaceae, Sapotaceae, Simaroubaceae, Ulmaceae, Verbenaceae	Ericales, Gentianales, Lamiales, Malpighiales, Myrtales, Rosales, Sapindales,	asterids, euasterids I, eurosid I, eurosid II, rosids
<i>Aleuroclava piperis</i>	Piperaceae	Piperales	magnoliids
<i>Aleuroclava psidii</i>	Caprifoliaceae, Euphorbiaceae, Lauraceae, Moraceae, Myrsinaceae, Myrtaceae, Rosaceae, Salicaceae, Sapindaceae, Ulmaceae	Dipsacales, Ericales, Laurales, Malpighiales, Myrtales, Rosales, Sapindales	asterids, euasterids II, eurosid I, eurosid II, magnoliids, rosids
<i>Aleuroclava pulcherrimus</i>	Fabaceae	Fabales	eurosid I
<i>Aleuroclava similis</i>	Aquifoliaceae, Ericaceae, Theaceae	Aquifoliales, Ericales	asterids, euasterids II
<i>Aleurocybotus graminicolus</i>	Poaceae	Poales	commelinids
<i>Aleurocybotus occiduus</i>	Cyperaceae, Poaceae	Poales	commelinids
<i>Aleuroduplidens eucalyptifolia</i>	Myrtaceae	Myrtales	rosids
<i>Aleuroduplidens santali</i>	Santalaceae	Santalales	core eudicots
<i>Aleuroduplidens triangularis</i>	Fabaceae	Fabales	eurosid I
<i>Aleuroduplidens wellsae</i>	Fabaceae	Fabales	eurosid I
<i>Aleuroglandulus magnus</i>	Arecaceae	Arecales	commelinids
<i>Aleuroglandulus malangae</i>	Araceae	Alismatales	monocots
<i>Aleuroglandulus striatus</i>	Polygonaceae	Caryophyllales	core eudicots
<i>Aleuroglandulus subtilis</i>	Arecaceae, Rubiaceae	Arecales, Gentianales	commelinids, euasterids I
<i>Aleuoinanis myrtacei</i>	Myrtaceae	Myrtales	rosids
<i>Aleuolobus diastematus</i>	Combretaceae, Salicaceae	Malpighiales, Myrtales	eurosid I, rosids
<i>Aleuolobus graminicola</i>	Poaceae	Poales	commelinids
<i>Aleuolobus hargreavesi</i>	Poaceae	Poales	commelinids
<i>Aleuolobus marlatti</i>	Araceae, Daphniphyllaceae, Moraceae, Rutaceae, Ulmaceae	Alismatales, Rosales, Sapindales, Saxifragales	core eudicots, eurosid I, eurosid II, monocots
<i>Aleuolobus moundi</i>	Sapotaceae	Ericales	asterids
<i>Aleuolobus onitshae</i>	Euphorbiaceae	Malpighiales	eurosid I
<i>Aleuolobus styraci</i>	Styracaceae	Ericales	asterids
<i>Aleuolobus teucarii</i>	Lamiaceae	Lamiales	euasterids I
<i>Aleuolobus cassiae</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus corbettaiformis</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus dalbergiae</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus kallarensis</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus littoralis</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus marginiquis</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus moundi</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus nenciae</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus nigrus</i>	Fabaceae	Fabales	eurosid I
<i>Aleuomarginatus serdangensis</i>	Host indet.		
<i>Aleuomarginatus tephrosiae</i>	Fabaceae	Fabales	eurosid I
<i>Aleuopapillatus kumariensis</i>	Verbenaceae	Lamiales	euasterids I
<i>Aleuoparadoxus arctostaphyli</i>	Ericaceae	Ericales	asterids

Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Aleuoparadoxus gardemiae</i>	Ebenaceae, Rubiaceae	Ericales, Gentianales	asterids, euasterids I
<i>Aleuoparadoxus ilicicola</i>	Aquifoliaceae	Aquifoliales	euasterids II
<i>Aleuoparadoxus iridescens</i>	Lamiaceae, Rhamnaceae, Rosaceae	Lamiales, Rosales	euasterids I, eurosids I
<i>Aleuoparadoxus punctatus</i>	Anacardiaceae, Euphorbiaceae, Rosaceae, Salicaceae	Malpighiales, Rosales, Sapindales	eurosids I, eurosids II
<i>Aleuoplatus acaciae</i>	Fabaceae	Fabales	eurosids I
<i>Aleuoplatus berbericulus</i>	Aquifoliaceae, Berberidaceae	Aquifoliales, Ranunculales	euasterids II, eudicots
<i>Aleuoplatus bossi</i>	Apocynaceae, Brassicaceae, Canellaceae, Meliaceae, Polygalaceae, Smilacaceae	Brassicales, Canellales, Fabales, Gentianales, Liliales, Sapindales	euasterids I, eurosids I, eurosids II, magnoliids, monocots
<i>Aleuoplatus cadabae</i>	Apocynaceae, Brassicaceae	Brassicales, Gentianales	euasterids I, eurosids II
<i>Aleuoplatus cockerelli</i>	Asteraceae	Asterales	euasterids II
<i>Aleuoplatus cococolus</i>	Arecaceae, Myrtaceae	Arecales, Myrtales	commelinids, rosids
<i>Aleuoplatus dorsipallidus</i>	Conuniaceae, Lauraceae, Moraceae	Laurales, Oxalidales, Rosales	eurosids I, magnoliids
<i>Aleuoplatus epigaeae</i>	Ericaceae, Lauraceae	Ericales, Laurales	asterids, magnoliids
<i>Aleuoplatus evodiae</i>	Rutaceae	Sapindales	eurosids II
<i>Aleuoplatus fici</i>	Moraceae	Rosales	eurosids I
<i>Aleuoplatus ilicis</i>	Aquifoliaceae, Ericaceae, Lauraceae	Aquifoliales, Ericales, Laurales	asterids, euasterids II, magnoliids
<i>Aleuoplatus incisus</i>	Clusiaceae, Euphorbiaceae	Malpighiales	eurosids I
<i>Aleuoplatus pectiniferus</i>	Betulaceae, Euphorbiaceae, Malvaceae, Moraceae, Myrtaceae, Rutaceae, Salicaceae, Theaceae	Ericales, Fagales, Malpighiales, Malvales, Myrtales, Rosales, Sapindales	asterids, eurosids I, eurosids II, rosids
<i>Aleuoplatus periplocae</i>	Apocynaceae	Gentianales	euasterids I
<i>Aleuoplatus perseaphagus</i>	Lauraceae	Laurales	magnoliids
<i>Aleuoplatus quercusaquaticae</i>	Fagaceae	Fagales	eurosids I
<i>Aleuopleurocelus nigrans</i>	Boraginaceae*, Caprifoliaceae, Ericaceae, Fagaceae, Lamiaceae, Lauraceae, Ranunculaceae, Rhamnaceae, Rosaceae	Dipsacales, Ericales, Fagales, Lamiales, Laurales, Ranunculales, Rosales	asterids, euasterids I, euasterids II, eudicots, eurosids I, magnoliids
<i>Aleuopteridis eastopi</i>	Oleandraceae	§Aspleniales	
<i>Aleuopteridis filicicola</i>	Oleandraceae, Pteridaceae, Thelypteridaceae	§Aspleniales, §Parkeriales	
<i>Aleuopteridis hargreavesi</i>	Undetermined fern		
<i>Aleuopteridis jamesi</i>	Pteridaceae	§Parkeriales	
<i>Aleuopterus perseae</i>	Lauraceae, Myrtaceae	Laurales, Myrtales	magnoliids, rosids
<i>Aleurothrix aepim</i>	Asteraceae, Convolvulaceae, Euphorbiaceae, Rubiaceae, Rutaceae	Asterales, Gentianales, Malpighiales, Sapindales, Solanales	euasterids I, euasterids II, eurosids I, eurosids II
<i>Aleurothrix antidesmae</i>	Euphorbiaceae	Malpighiales	eurosids I
<i>Aleurothrix bondari</i>	Malvaceae	Malvales	eurosids II
<i>Aleurothrix floccosus</i>	Anacardiaceae, Annonaceae, Apocynaceae, Asteraceae, Chrysobalanaceae, Ebenaceae, Liliaceae, Lorantheaceae, Malvaceae, Myrtaceae, Nyctaginaceae, Polygonaceae, Rubiaceae, Rutaceae, Sapotaceae, Solanaceae, Zygophyllaceae*	Asterales, Ericales, Caryophyllales, Gentianales, Liliales, Magnoliales, Malpighiales, Malvales, Myrtales, Santalales, Sapindales, Solanales	asterids, core eudicots, euasterids I, euasterids II, eurosids I, eurosids II, magnoliids, monocots, rosids
<i>Aleurothrix interrogationis</i>	Rhamnaceae	Rosales	eurosids I
<i>Aleurothrix lucumai</i>	Sapotaceae	Ericales	asterids
<i>Aleurothrix myrtacei</i>	Myrtaceae	Myrtales	rosids
<i>Aleurothrix porteri</i>	Anacardiaceae, Lauraceae, Myrtaceae, Rutaceae, Sapotaceae, Solanaceae, Verbenaceae	Ericales, Lamiales, Laurales, Myrtales, Sapindales, Solanales	asterids, euasterids I, eurosids II, magnoliids, rosids
<i>Aleurothrix proximans</i>	Lauraceae	Laurales	magnoliids
<i>Aleurothrix smilaceti</i>	Smilacaceae	Liliales	monocots
<i>Aleurotithius timberlakei</i>	Boraginaceae*		euasterids I
<i>Aleurotrachelus alpinus</i>	Rosaceae	Rosales	eurosids I
<i>Aleurotrachelus ampullatus</i>	Fabaceae	Fabales	eurosids I
<i>Aleurotrachelus asparagi</i>	Liliaceae	Liliales	monocots
<i>Aleurotrachelus atratus</i>	Arecaceae	Arecales	commelinids
<i>Aleurotrachelus camelliae</i>	Theaceae	Ericales	asterids
<i>Aleurotrachelus duplicatus</i>	Rubiaceae	Gentianales	euasterids I
<i>Aleurotrachelus fumipennis</i>	Poaceae	Poales	commelinids

Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Aleurotrachelus limbatus</i>	Ericaceae, Fabaceae	Ericales, Fabales	asterids, eurosids I
<i>Aleurotrachelus longispinus</i>	Rubiaceae	Gentianales	euasterids I
<i>Aleurotrachelus minutus</i>	Moraceae	Rosales	eurosids I
<i>Aleurotrachelus multipapillus</i>	Poaceae	Poales	commelinids
<i>Aleurotrachelus obscurus</i>	Fabaceae	Fabales	eurosids I
<i>Aleurotrachelus taremae</i>	Rubiaceae	Gentianales	euasterids I
<i>Aleurotrachelus tracheifer</i>	Loganiaceae, Melastomataceae	Gentianales, Myrtales	euasterids I, rosids
<i>Aleurotrachelus trachoides</i>	Asteraceae, Bignoniaceae, Casuarinaceae, Convolvulaceae, Dioscoreaceae, Melastomataceae, Rubiaceae, Solanaceae, Verbenaceae	Asterales, Dioscoreales, Fagales, Gentianales, Lamiales, Myrtales, Solanales	II, eurosids I, monocots, rosids
<i>Aleoutuba jelinekii</i>	Adoxaceae, Ericaceae, Myrtaceae	Dipsacales, Ericales, Myrtales	asterids, euasterids II, rosids
<i>Aleuotulus anthuricola</i>	Araceae	Alismatales	monocots
<i>Aleuotulus arundinacea</i>	Poaceae	Poales	commelinids
<i>Aleuotulus nephrolepidis</i>	Aspleniaceae, Blechnaceae, Cyatheaceae, Dryopteridaceae (Aspidiaceae), Oleandraceae, Pteridaceae, Schizaeaceae, Thelypteridaceae, Woodsiaceae	§Aspleniales, §Cyatheaales, §Parkeriales, §Schizaeales	
<i>Aleuotulus pteridophytæ</i>	Selaginellaceae, Pteridaceae, Schizaeaceae	§Parkeriales, §Schizaeales, §Selaginellales	
<i>Aleuroviggianus adanaensis</i>	Fagaceae	Fagales	eurosids I
<i>Aleuroviggianus adrianae</i>	Fagaceae	Fagales	eurosids I
<i>Aleuroviggianus graecus</i>	Fagaceae	Fagales	eurosids I
<i>Aleuroviggianus halperini</i>	Fagaceae	Fagales	eurosids I
<i>Aleuroviggianus polymorphus</i>	Fagaceae	Fagales	eurosids I
<i>Aleuroviggianus zonalus</i>	Fagaceae	Fagales	eurosids I
<i>Aleyrodes amnicola</i>	Salicaceae	Malpighiales	eurosids I
<i>Aleyrodes elevatus</i>	Euphorbiaceae, Moraceae	Malpighiales, Rosales	eurosids I
<i>Aleyrodes lomiceræ</i>	Apiaceae, Asteraceae, Balsaminaceae, Brassicaceae, Campanulaceae, Caprifoliaceae, Clusiaceae, Euphorbiaceae, Grossulariaceae, Lamiaceae, Onagraceae, Oxalidaceae, Papaveraceae, Ranunculaceae, Rosaceae, Scrophulariaceae, Urticaceae, Violaceae	Apiales, Asterales, Brassicales, Dipsacales, Ericales, Lamiales, Malpighiales, Myrtales, Oxalidales, Ranunculales, Rosales, Saxifragales	asterids, core eudicots, euasterids I, euasterids II, eudicots, eurosids I, eurosids II, rosids
<i>Aleyrodes prolella</i>	Apiaceae, Asteraceae, Balsaminaceae, Berberidaceae, Brassicaceae, Campanulaceae, Euphorbiaceae, Fabaceae, Fagaceae, Papaveraceae, Ranunculaceae, Scrophulariaceae	Apiales, Asterales, Brassicales, Ericales, Fabales, Fagales, Lamiales, Malpighiales, Ranunculales	asterids, euasterids I, euasterids II, eudicots, eurosids I, eurosids II
<i>Aleyrodes prunosus</i>	Euphorbiaceae, Rosaceae, Rhamnaceae	Malpighiales, Rosales	eurosids I
<i>Aleyrodes singularis</i>	Euphorbiaceae	Malpighiales	eurosids I
<i>Aleyrodiella lamellifera</i>	Ulmaceae	Rosales	eurosids I
<i>Apobemisia kuwanai</i>	Moraceae	Rosales	eurosids I
<i>Arachmaleyrodes insignis</i>	Combretaceae	Myrtales	rosids
<i>Asialeyrodes lushanensis</i>	Myrtaceae	Myrtales	rosids
<i>Asialeyrodes maesae</i>	Myrsinaceae	Ericales	asterids
<i>Asterobemisia carpini</i>	Anacardiaceae, Betulaceae, Cannabaceae, Caprifoliaceae, Ericaceae, Fabaceae, Fagaceae, Grossulariaceae, Malvaceae, Ranunculaceae, Rosaceae, Salicaceae, Sapindaceae, Saxifragaceae, Ulmaceae	Dipsacales, Ericales, Fabales, Fagales, Malpighiales, Malvales, Ranunculales, Rosales, Sapindales, Saxifragales	asterids, core eudicots, euasterids II, eudicots, eurosids I, eurosids II
<i>Asterobemisia dentata</i>	Moraceae	Rosales	eurosids I
<i>Asterobemisia obenbergeri</i>	Lamiaceae	Lamiales	euasterids I
<i>Asterochiton cordiae</i>	Boraginaceae*		euasterids I
<i>Asterochiton pittospori</i>	Pittosporaceae	Apiales	euasterids II
<i>Bellitudo hispaniolae</i>	Polygonaceae	Caryophyllales	core eudicots
<i>Bellitudo jamaicae</i>	Polygonaceae	Caryophyllales	core eudicots

Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Bemisialeyrodes brideliae</i>	Euphorbiaceae	Malpighiales	eurosids I
<i>Bemisialeyrodes pauliani</i>	Moraceae, Ulmaceae	Rosales	eurosids I
<i>Bemisia afer</i>	Lythraceae, Moraceae, Rutaceae	Myrtales, Rosales, Sapindales	eurosids I, eurosids II, rosids
<i>Bemisia antennata</i>	Boraginaceae*, Fabaceae, Moraceae	Fabales, Rosales	euasterids I, eurosids I
<i>Bemisia berbericola</i>	Berberidaceae, Euphorbiaceae, Moraceae, Rhamnaceae, Rosaceae	Malpighiales, Ranunculales, Rosales	eudicots, eurosids I
<i>Bemisia combreticula</i>	Combretaceae	Myrtales	rosids
<i>Bemisia formosana</i>	Poaceae	Poales	commelinids
<i>Bemisia giffardi</i>	Boraginaceae*, Oleaceae, Rutaceae	Lamiales, Sapindales	euasterids I, eurosids II
<i>Bemisia gigantean</i>	Elaeocarpaceae	Oxalidales	eurosids I
<i>Bemisia guieriae</i>	Combretaceae	Myrtales	rosids
<i>Bemisia hirta</i>	Brassicaceae	Brassicales	eurosids II
<i>Bemisia lauracea</i>	Lauraceae	Laurales	magnoliids
<i>Bemisia mesasiatica</i>	Caprifoliaceae, Rosaceae	Dipsacales, Rosales	euasterids II, eurosids I
<i>Bemisia pongamiae</i>	Fabaceae	Fabales	eurosids I
<i>Bemisia silvatica</i>	Anacardiaceae, Betulaceae, Ericaceae, Fabaceae, Fagaceae, Rhamnaceae, Rosaceae	Ericales, Fabales, Fagales, Rosales, Sapindales	asterids, eurosids I, eurosids II
<i>Bemisia subdecipiens</i>	Myrtaceae	Myrtales	rosids
<i>Bemisia tabaci</i>	Acanthaceae, Amaranthaceae, Anacardiaceae, Annonaceae, Apiaceae, Apocynaceae, Araceae, Aristolochiaceae, Asteraceae, Bignoniaceae, Bixaceae, Brassicaceae, Cannabaceae, Caprifoliaceae, Chrysobalanaceae, Cistaceae, Clusiaceae, Commelinaceae, Convolvulaceae, Cucurbitaceae, Ericaceae, Euphorbiaceae, Fabaceae, Fagaceae, Geraniaceae, Grossulariaceae, Lamiaceae, Lauraceae, Linaceae, Loganiaceae, Lythraceae, Malvaceae, Menispermaceae, Moraceae, Moringaceae, Musaceae, Myrtaceae, Nyctaginaceae, Oleaceae, Oxalidaceae, Passifloraceae, Pedaliaceae, Poaceae, Ranunculaceae, Rhamnaceae, Rosaceae, Rubiaceae, Rutaceae, Salicaceae, Sapindaceae, Scrophulariaceae, Solanaceae, Thymelaeaceae, Ulmaceae, Urticaceae, Verbenaceae, Zygophyllaceae*	Alismatales, Apiales, Asterales, Brassicales, Caryophyllales, Commelinales, Cucurbitales, Dipsacales, Ericales, Fabales, Fagales, Gentianales, Geraniales, Lamiales, Laurales, Magnoliales, Malpighiales, Malvales, Myrtales, Oxalidales, Piperales, Poales, Ranunculales, Rosales, Sapindales, Saxifragales, Solanales, Zingiberales	rosids, asterids, commelinids, core eudicots, euasterids I, euasterids II, eudicots, eurosids I, eurosids II, magnoliids, monocots, rosids
<i>Bemisia tuberculata</i>	Euphorbiaceae	Malpighiales	eurosids I
<i>Bemisiella artemisiae</i>	Asteraceae	Asterales	euasterids II
<i>Brazzaleyrodes eriococciformis</i>	Fabaceae	Fabales	eurosids I
<i>Bulgarialeyrodes cotesii</i>	Rosaceae	Rosales	eurosids I
<i>Calluneyrodes callunae</i>	Ericaceae	Ericales	asterids
<i>Chitonaleyrodes canberrensis</i>	Fabaceae	Fabales	eurosids I
<i>Cockerelliella curcuma</i>	Musaceae, Zingiberaceae	Zingiberales	commelinids
<i>Cockerelliella psidii</i>	Dipterocarpaceae, Magnoliaceae, Meliaceae, Moraceae, Myrtaceae, Verbenaceae	Lamiales, Magnoliales, Malvales, Myrtales, Rosales, Sapindales	euasterids I, eurosids I, eurosids II, magnoliids, rosids
<i>Cohicaleyrodes alternans</i>	Clusiaceae, Combretaceae, Connaraceae, Euphorbiaceae, Fabaceae, Loganiaceae, Rubiaceae, Salicaceae, Verbenaceae	Fabales, Gentianales, Lamiales, Malpighiales, Myrtales, Oxalidales	euasterids I, eurosids I, rosids
<i>Cohicaleyrodes crossopterygis</i>	Combretaceae, Rubiaceae	Gentianales, Myrtales	euasterids I, rosids
<i>Cohicaleyrodes obscura</i>	Rubiaceae	Gentianales	euasterids I
<i>Cohicaleyrodes platysepalii</i>	Fabaceae, Zingiberaceae	Fabales, Zingiberales	commelinids, eurosids I

Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Cohicaleyrodes quadrilongispinae</i>	Annonaceae, Bignoniaceae, Combretaceae, Euphorbiaceae, Fabaceae, Loganiaceae, Melastomataceae, Moraceae, Rubiaceae, Sapindaceae, Verbenaceae, Vitaceae*	Fabales, Gentianales, Lamiales, Magnoliales, Malpighiales, Myrtales, Rosales, Sapindales	euasterids I, eucosids I, eucosids II, magnoliids, rosids
<i>Cohicaleyrodes recurvispina</i>	Annonaceae, Dilleniaceae*, Moraceae, Rubiaceae	Gentianales, Magnoliales, Rosales	core eudicots, euasterids I, eucosids I, magnoliids
<i>Combesaleyrodes bouqueti</i>	Rubiaceae	Gentianales	euasterids I
<i>Combesaleyrodes tauffliebi</i>	Dilleniaceae*, Rubiaceae	Gentianales	core eudicots, euasterids I
<i>Corbettia baphiae</i>	Fabaceae	Fabales	eucosids I
<i>Corbettia bauhiniae</i>	Fabaceae	Fabales	eucosids I
<i>Corbettia graminis</i>	Poaceae	Poales	commelinids
<i>Corbettia isoberliniae</i>	Fabaceae	Fabales	eucosids I
<i>Corbettia lonchocarpi</i>	Fabaceae	Fabales	eucosids I
<i>Corbettia millettiaicola</i>	Apocynaceae, Fabaceae	Fabales, Gentianales	euasterids I, eucosids I
<i>Corbettia tamarindi</i>	Fabaceae	Fabales	eucosids I
<i>Crenidorsum aroidephagus</i>	Araceae	Alismatales	monocots
<i>Crenidorsum celebes</i>	Cunoniaceae, Lauraceae	Laurales, Oxalidales	eucosids I, magnoliids
<i>Crenidorsum commune</i>	Polygonaceae	Caryophyllales	core eudicots
<i>Crenidorsum lasangensis</i>	Musaceae	Zingiberales	commelinids
<i>Crenidorsum magnisetae</i>	Polygonaceae	Caryophyllales	core eudicots
<i>Crenidorsum malpighiae</i>	Malpighiaceae	Malpighiales	eucosids I
<i>Crenidorsum micheliae</i>	Magnoliaceae, Myrsinaceae	Ericales, Magnoliales	asterids, magnoliids
<i>Crenidorsum millennium</i>	Myrtaceae	Myrtales	rosids
<i>Crenidorsum morobensis</i>	Myrtaceae	Myrtales	rosids
<i>Crenidorsum stigmaphylli</i>	Malpighiaceae	Malpighiales	eucosids I
<i>Crenidorsum tuberculatum</i>	Polygonaceae	Caryophyllales	core eudicots
<i>Crescentaleyrodes paulianae</i>	Poaceae	Poales	commelinids
<i>Crescentaleyrodes semilunaris</i>	Poaceae	Poales	commelinids
<i>Cryptolingula perplexa</i>	Host indet.		
<i>Dialeurodes agalmae</i>	Araliaceae	Apiales	euasterids II
<i>Dialeurodes ara</i>	Moraceae	Rosales	eucosids I
<i>Dialeurodes armatus</i>	Meliaceae	Sapindales	eucosids II
<i>Dialeurodes bicornicauda</i>	Meliaceae	Sapindales	eucosids II
<i>Dialeurodes cerifera</i>	Celastraceae	Celastrales	eucosids I
<i>Dialeurodes cinnamomi</i>	Lauraceae	Laurales	magnoliids
<i>Dialeurodes citri</i>	Apocynaceae, Araliaceae, Arecaceae, Bignoniaceae, Boraginaceae*, Caprifoliaceae, Ebenaceae, Euphorbiaceae, Fagaceae, Hamamelidaceae, Lauraceae, Lythraceae, Magnoliaceae, Malpighiaceae, Meliaceae, Moraceae, Myrsinaceae, Myrtaceae, Oleaceae, Proteaceae, Rosaceae, Rubiaceae, Rutaceae, Sabiaceae*, Simaroubaceae, Smilacaceae, Staphyleaceae, Theaceae, Vitaceae*	Arecales, Apiales, Crossosomatales, Dipsacales, Ericales, Fagales, Gentianales, Lamiales, Laurales, Liliales, Magnoliales, Malpighiales, Myrtales, Proteales, Rosales, Sapindales, Saxifragales	asterids, commelinids, core eudicots, euasterids I, euasterids II, eudicots, eucosids I, eucosids II, magnoliids, monocots, rosids
<i>Dialeurodes cyathispinifera</i>	Host indet.		
<i>Dialeurodes decaspermii</i>	Myrtaceae	Myrtales	rosids
<i>Dialeurodes dubia</i>	Host indet.		
<i>Dialeurodes emarginata</i>	Loganiaceae, Malvaceae	Gentianales, Malvales	euasterids I, eucosids II
<i>Dialeurodes erythrinae</i>	Fabaceae	Fabales	eucosids I
<i>Dialeurodes ixorae</i>	Rubiaceae, Sapotaceae	Ericales, Gentianales	asterids, euasterids I
<i>Dialeurodes joholensis</i>	Host indet.		
<i>Dialeurodes kirkaldyi</i>	Apocynaceae, Combretaceae, Loganiaceae, Lythraceae, Malpighiaceae, Malvaceae, Oleaceae, Rubiaceae, Rutaceae, Verbenaceae	Gentianales, Lamiales, Malpighiales, Malvales, Myrtales, Sapindales	euasterids I, eucosids I, eucosids II, rosids
<i>Dialeurodes lanceolata</i>	Celastraceae	Celastrales	eucosids I
<i>Dialeurodes minahassai</i>	Myrtaceae	Myrtales	rosids
<i>Dialeurodes mirabilis</i>	Host indet.		

Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Dialeurodes platycus</i>	Myrtaceae	Myrtales	rosids
<i>Dialeurolobus rhamni</i>	Lythraceae, Rhamnaceae	Myrtales, Rosales	eurosids I, rosids
<i>Dialeurolonga africana</i>	Celastraceae	Celastrales	eurosids I
<i>Dialeurolonga communis</i>	Loganiaceae, Rubiaceae, Rutaceae, Sapindaceae	Gentianales, Sapindales	euasterids I, eurosids II
<i>Dialeurolonga elliptica</i>	Apocynaceae	Gentianales	euasterids I
<i>Dialeurolonga elongata</i>	Rubiaceae, Rutaceae, Sapindaceae	Gentianales, Sapindales	euasterids I, eurosids II
<i>Dialeurolonga hoyti</i>	Rubiaceae	Gentianales	euasterids I
<i>Dialeurolonga maculata</i>	Cunoniaceae	Oxalidales	eurosids I
<i>Dialeurolonga nemoralis</i>	Annonaceae	Magnoliales	magnoliids
<i>Dialeurolonga phyllarthromis</i>	Bignoniaceae	Lamiales	euasterids I
<i>Dialeurolonga ravensarae</i>	Lauraceae	Laurales	magnoliids
<i>Dialeurolonga rusostigmoides</i>	Host indet.		
<i>Dialeurolonga swainii</i>	Moraceae	Rosales	eurosids I
<i>Dialeurolonga tenella</i>	Apocynaceae	Gentianales	euasterids I
<i>Dialeurolonga trialeuroides</i>	Host indet.		
<i>Dialeuropora brideliae</i>	Euphorbiaceae, Lardizabalaceae, Lauraceae	Malpighiales, Ranunculales, Laurales	eudicots, eurosids I, magnoliids
<i>Dialeuropora decempuncta</i>	Annonaceae, Araceae, Boraginaceae*, Dipterocarpaceae, Euphorbiaceae, Fabaceae, Lauraceae, Meliaceae, Moraceae, Rosaceae, Salicaceae, Verbenaceae	Alismatales, Fabales, Lamiales, Laurales, Magnoliales, Malpighiales, Malvales, Rosales, Sapindales	euasterids I, eurosids I, eurosids II, magnoliids, monocots
<i>Dialeuropora mangiferae</i>	Anacardiaceae	Sapindales	eurosids II
<i>Dialeuropora papillata</i>	Annonaceae, Arecaceae, Euphorbiaceae, Fabaceae, Linaceae, Malvaceae	Arecales, Fabales, Magnoliales, Malpighiales, Malvales	commelinids, eurosids I, eurosids II, magnoliids
<i>Dialeuropora silvarum</i>	Host indet.		
<i>Dialeuropora viburni</i>	Caprifoliaceae	Dipsacales	euasterids II
<i>Disiphon</i> sp.	Myrtaceae, Piperaceae?	Myrtales, Piperales?	magnoliids?, rosids
<i>Dumbletoniella callistemoni</i>	Myrtaceae	Myrtales	rosids
<i>Dumbletoniella eucalypti</i>	Myrtaceae	Myrtales	rosids
<i>Dumbletoniella pittospori</i>	Pittosporaceae	Apiales	euasterids II
<i>Dumbletoniella rotunda</i>	Myrtaceae	Myrtales	rosids
<i>Dumbletoniella xanthorrhoeae</i>	Liliaceae	Liliales	monocots
<i>Extensaleyrodes akurensis</i>	Host indet.		
<i>Extensaleyrodes falcata</i>	Fabaceae	Fabales	eurosids I
<i>Fascaleyrodes palmae</i>	Arecaceae	Arecales	commelinids
<i>Fascaleyrodes rara</i>	Annonaceae	Magnoliales	magnoliids
<i>Filicaleyrodes bosseri</i>	Undetermined fern		
<i>Filicaleyrodes williamsii</i>	Aspleniaceae, Blechnaceae, Dryopteridaceae (Aspidiaceae), Oleandraceae, Schizaeaceae, Thelypteridaceae, Woodsiaceae	§Aspleniales, §Schizaeales	
<i>Fippataleyrodes indica</i>	Lauraceae, Moraceae, Myrsinaceae, Salicaceae	Ericales, Laurales, Malpighiales, Rosales	asterids, eurosids I, magnoliids
<i>Fippataleyrodes litseae</i>	Lauraceae	Laurales	magnoliids
<i>Gagudjuia allosyncarpiae</i>	Myrtaceae	Myrtales	rosids
<i>Gomenella dryandrae</i>	Proteaceae	Proteales	eudicots
<i>Gomenella multipora</i>	Host indet.		
<i>Harpaleyrodes tuberculata</i>	Combretaceae	Myrtales	rosids
<i>Hempelia chivelensis</i>	Lauraceae	Laurales	magnoliids
<i>Heteraleyrodes bambusicola</i>	Poaceae	Poales	commelinids
<i>Heterobemisia alba</i>	Grossulariaceae	Saxifragales	core eudicots
<i>Indoaleyrodes glochidioni</i>	Euphorbiaceae	Malpighiales	eurosids I
<i>Indoaleyrodes laos</i>	Euphorbiaceae, Rubiaceae	Malpighiales	eurosids I
<i>Indoaleyrodes pseudoculatus</i>	Myrtaceae	Myrtales	rosids
<i>Kanakarajella vulgaris</i>	Asteraceae, Fabaceae, Myrtaceae, Oleaceae, Rubiaceae	Asterales, Fabales, Gentianales, Lamiales, Myrtales	II, eurosids I, rosids
<i>Leucopogonella pallida</i>	Ericaceae	Ericales	asterids
<i>Leucopogonella simila</i>	Ericaceae	Ericales	asterids
<i>Lipaleyrodes atriplex</i>	Amaranthaceae	Caryophyllales	core eudicots
<i>Lipaleyrodes breyniae</i>	Euphorbiaceae	Malpighiales	eurosids I

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Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Lipaleyrodes crossandrae</i>	Acanthaceae, Amaranthaceae	Lamiales, Caryophyllales	core eudicots, euasterids I
<i>Lipaleyrodes euphorbiae</i>	Euphorbiaceae	Malpighiales	eurosid I
<i>Lipaleyrodes phyllanthi</i>	Asteraceae, Euphorbiaceae	Asterales, Malpighiales	euasterids II, eurosid I
<i>Malayaleyrodes ?lumpurensis</i>	Moraceae	Rosales	eurosid I
<i>Marginaleyrodes ixorae</i>	Rubiaceae	Gentianales	euasterids I
<i>Massilieurodes americanus</i>	Aquifoliaceae, Betulaceae, Cornaceae, Ebenaceae, Ericaceae, Hamamelidaceae, Magnoliaceae, Oleaceae, Symplocaceae	Aquifoliales, Cornales, Ericales, Fagales, Lamiales, Magnoliales, Saxifragales	asterids, core eudicots, euasterids I, euasterids II, eurosid I, magnoliids
<i>Massilieurodes chittendeni</i>	Ericaceae	Ericales	asterids
<i>Massilieurodes euryae</i>	Theaceae	Ericales	asterids
<i>Massilieurodes fici</i>	Euphorbiaceae, Moraceae, Myrsinaceae, Rhamnaceae, Rutaceae, Theaceae, Ulmaceae	Ericales, Malpighiales, Rosales, Sapindales	asterids, eurosid I, eurosid II
<i>Massilieurodes setiger</i>	Adoxaceae	Dipsacales	euasterids II
<i>Metabemisia filicis</i>	Davalliaceae, Dryopteridaceae (Aspidiaceae), Oleandraceae, Pteridaceae, Thelypteridaceae	§Aspleniales, §Parkeriales	
<i>Metabemisia palawana</i>	Dryopteridaceae	§Aspleniales	
<i>Minutaleyrodes minuta</i>	Celastraceae, Rubiaceae	Celastrales, Gentianales	euasterids I, eurosid I
<i>Minutaleyrodes suischanus</i>	Lauraceae, Myrsinaceae	Ericales, Laurales	asterids, magnoliids
<i>Mixaleyrodes ?polystichii</i>	Dryopteridaceae (Aspidiaceae), Marattiaceae, Pteridaceae	§Aspleniales, §Marattiales, §Parkeriales	
<i>Neoleurotrachelus bertillonii</i>	Apocynaceae, Bignoniaceae, Dilleniaceae*, Euphorbiaceae, Loganiaceae, Meliaceae, Menispermaceae, Moraceae, Ochnaceae, Rubiaceae	Gentianales, Lamiales, Malpighiales, Ranunculales, Rosales, Sapindales	core eudicots, euasterids I, eudicots, eurosid I, eurosid II
<i>Neoleurotrachelus graberi</i>	Annonaceae, Bignoniaceae, Celastraceae, Combretaceae, Euphorbiaceae, Lythraceae, Meliaceae, Moraceae, Myrtaceae, Rhamnaceae, Rubiaceae	Celastrales, Gentianales, Lamiales, Magnoliales, Malpighiales, Myrtales, Sapindales, Rosales	euasterids I, eurosid I, eurosid II, magnoliids, rosids
<i>Neomaskellia andropogonis</i>	Poaceae	Poales	commelinids
<i>Neomaskellia bergii</i>	Poaceae	Poales	commelinids
<i>Neopealius rubi</i>	Fabaceae, Lamiaceae, Lauraceae, Rosaceae	Fabales, Lamiales, Laurales, Rosales	euasterids I, eurosid I, magnoliids
<i>Nigrasialeyrodes convexus</i>	Myrtaceae	Myrtales	rosids
<i>Orchamoplatus caledonicus</i>	Moraceae, Rutaceae, Ulmaceae	Rosales, Sapindales	eurosid I, eurosid II
<i>Orchamoplatus citri</i>	Rutaceae	Sapindales	eurosid II
<i>Orchamoplatus dentatus</i>	Loranthaceae, Myrtaceae	Myrtales, Santalales	core eudicots, rosids
<i>Orchamoplatus louisierussellae</i>	Loranthaceae, Myrtaceae	Myrtales, Santalales	core eudicots, rosids
<i>Orchamoplatus mammaeferus</i>	Euphorbiaceae, Myrtaceae, Rutaceae	Malpighiales, Myrtales, Sapindales	eurosid I, eurosid II, rosids
<i>Orchamoplatus montanus</i>	Araliaceae, Cunoniaceae, Ericaceae	Apiales, Ericales, Oxalidales	asterids, euasterids II, eurosid I
<i>Orchamoplatus niuginii</i>	Clusiaceae, Linaceae	Malpighiales	eurosid I
<i>Orchamoplatus noumeae</i>	Rutaceae	Sapindales	eurosid II
<i>Orchamoplatus plumensis</i>	Host indet.		
<i>Orchamoplatus porosus</i>	Myrtaceae	Myrtales	rosids
<i>Orientalleyrodes zeylanicus</i>	Brassicaceae	Brassicales	eurosid II
<i>Orstomaleyrodes fimbriata</i>	Bignoniaceae, Combretaceae, Fabaceae, Salicaceae, Sapindaceae, Ulmaceae	Fabales, Lamiales, Malpighiales, Myrtales, Rosales, Sapindales	euasterids I, eurosid I, eurosid II, rosids
<i>Papillipes spinifer</i>	Fabaceae	Fabales	eurosid I
<i>Parabemisia aceris</i>	Sapindaceae	Sapindales	eurosid II
<i>Parabemisia javani</i>	Host indet.		
<i>Parabemisia lushanensis</i>	Host indet.		
<i>Parabemisia myricae</i>	Ebenaceae, Elaeocarpaceae, Ericaceae, Fagaceae, Lauraceae, Moraceae, Myricaceae, Myrsinaceae, Myrtaceae, Rosaceae, Rubiaceae, Rutaceae, Salicaceae, Theaceae	Ericales, Fagales, Gentianales, Laurales, Malpighiales, Myrtales, Oxalidales, Rosales, Sapindales	asterids, euasterids I, eurosid I, eurosid II, magnoliids, rosids

Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Parabemisia myrmecophila</i>	Dipterocarpaceae, Euphorbiaceae, Lauraceae, Rosaceae	Laurales, Malpighiales, Malvaeles, Rosales	eurosid I, eurosid II, magnoliids
<i>Paraleurolobus chamaedoreae</i>	Arecaceae	Arecales	commelinids
<i>Paulianaleyrodes splendens</i>	Dichapetalaceae	Malpighiales	eurosid I
<i>Paulianaleyrodes tetracerae</i>	Annonaceae, Clusiaceae, Cucurbitaceae, Dilleniaceae*	Cucurbitales, Magnoliales	core eudicots, eurosid I, magnoliids
<i>Pealius akebiae</i>	Lardizabalaceae	Ranunculales	eudicots
<i>Pealius azaleae</i>	Ericaceae	Ericales	asterids
<i>Pealius cryptus</i>	Moraceae	Rosales	eurosid I
<i>Pealius euryae</i>	Theaceae	Ericales	asterids
<i>Pealius fici</i>	Moraceae	Rosales	eurosid I
<i>Pealius kelloggi</i>	Fagaceae, Rosaceae	Fagales	eurosid I
<i>Pealius liquidambari</i>	Hamamelidaceae	Saxifragales	core eudicots
<i>Pealius madeirensis</i>	Oleaceae	Lamiales	euasterids I
<i>Pealius mori</i>	Moraceae, Salicaceae	Malpighiales, Rosales	eurosid I
<i>Pealius rhododendri</i>	Ericaceae	Ericales	asterids
<i>Pectinaleyrodes culcasiae</i>	Annonaceae, Araceae, Connaraceae, Malvaceae	Alismatales, Magnoliales, Malvaeles, Oxalidales	eurosid I, eurosid II, magnoliids, monocots
<i>Pectinaleyrodes trichlisiae</i>	Annonaceae, Bignoniaceae, Connaraceae, Dilleniaceae*, Euphorbiaceae, Fabaceae, Meliaceae, Menispermaceae, Rubiaceae, Salicaceae, Verbenaceae	Fabales, Gentianales, Lamiales, Magnoliales, Malpighiales, Oxalidales, Ranunculales, Sapindales	core eudicots, euasterids I, eudicots, eurosid I, eurosid II, magnoliids
<i>Pentaleyrodes cimnamomi</i>	Lauraceae	Laurales	magnoliids
<i>Pentaleyrodes hongkongensis</i>	Lauraceae	Laurales	magnoliids
<i>Pentaleyrodes yasumatsui</i>	Lauraceae	Laurales	magnoliids
<i>Pogonaleyrodes fastuosa</i>	Host indet.		
<i>Pogonaleyrodes zimmermanni</i>	Acanthaceae, Rubiaceae	Gentianales, Lamiales	euasterids I
<i>Pseudaleyroplatus kiensis</i>	Myrtaceae	Myrtales	rosids
<i>Ramsesseus folioli</i>	Fabaceae	Fabales	eurosid I
<i>Rhachisphora alishanensis</i>	Theaceae	Ericales	asterids
<i>Rhachisphora elongatus</i>	Sapotaceae	Ericales	asterids
<i>Rhachisphora franksae</i>	Loganiaceae	Gentianales	euasterids I
<i>Rhachisphora koshunensis</i>	Lauraceae	Laurales	magnoliids
<i>Rhachisphora maesae</i>	Myrsinaceae	Ericales	asterids
<i>Rhachisphora rutherfordi</i>	Loranthaceae	Santalales	core eudicots
<i>Rhachisphora trilobitoides</i>	Boraginaceae*, Melastomataceae, Myrtaceae, Rubiaceae, Sapindaceae, Sapotaceae	Ericales, Gentianales, Myrtales, Sapindales	asterids, euasterids I, eurosid II, rosids
<i>Rositalleyrodes oplismeni</i>	Poaceae	Poales	commelinids
<i>Rugaleyrodes angolensis</i>	Icacinaceae*, Marantaceae, Ochnaceae	Malpighiales, Zingiberales	commelinids, euasterids I, eurosid I
<i>Rugaleyrodes bidentata</i>	Annonaceae	Magnoliales	magnoliids
<i>Rugaleyrodes tetracerae</i>	Dichapetalaceae, Dilleniaceae*, Linaceae, Loganiaceae, Meliaceae, Myrtaceae, Ochnaceae	Gentianales, Malpighiales, Myrtales, Sapindales	core eudicots, euasterids I, eurosid I, eurosid II, rosids
<i>Rusostigma eugeniae</i>	Myrtaceae	Myrtales	rosids
<i>Rusostigma radiirugosa</i>	Anacardiaceae, Myrtaceae	Myrtales, Sapindales	eurosid II, rosids
<i>Rusostigma tokyonis</i>	Aquifoliaceae, Theaceae	Aquifoliales, Ericales	asterids, euasterids II
<i>Russellaleyrodes cumiugum</i>	Host indet.		
<i>Setaleyrodes mirabilis</i>	Menispermaceae, Moraceae	Ranunculales, Rosales	eudicots, eurosid I
<i>Setaleyrodes thretonai</i>	Euphorbiaceae, Moraceae	Malpighiales, Rosales	eurosid I
<i>Setaleyrodes viginiseta</i>	Fabaceae	Fabales	eurosid I
<i>Simplaleyrodes hemisphaerica</i>	Oleaceae	Lamiales	euasterids I
<i>Singhiella bicolor</i>	Myrtaceae	Myrtales	rosids
<i>Singhiella cardamomi</i>	Zingiberaceae	Zingiberales	commelinids
<i>Singhiella chinensis</i>	Lauraceae	Laurales	magnoliids
<i>Singhiella citrifolii</i>	Moraceae, Rubiaceae, Rutaceae	Gentianales, Rosales, Sapindales	euasterids I, eurosid I, eurosid II
<i>Singhiella delamarei</i>	Annonaceae	Magnoliales	magnoliids
<i>Singhiella kuraruensis</i>	Lauraceae	Laurales	magnoliids
<i>Singhiella premmiae</i>	Verbenaceae	Lamiales	euasterids I
<i>Singhiella subrotunda</i>	Lauraceae	Laurales	magnoliids

Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Singhius hibisci</i>	Annonaceae, Asteraceae, Convolvulaceae, Euphorbiaceae, Lauraceae, Malvaceae, Moraceae, Oleaceae, Salicaceae, Solanaceae, Ulmaceae, Vitaceae*	Asterales, Lamiales, Laurales, Magnoliales, Malpighiales, Malvales, Rosales, Solanales	euasterids I, euasterids II, eurosids I, eurosids II, magnoliids, rosids
<i>Singhius russellae</i>	Host indet.		
<i>Siphoninus immaculatus</i>	Araliaceae	Apiales	euasterids II
<i>Siphoninus phillyreae</i>	Fabaceae, Lythraceae, Oleaceae, Rhamnaceae, Rosaceae, Rutaceae	Fabales, Lamiales, Myrtales, Rosales, Sapindales	euasterids I, eurosids I, eurosids II, rosids
<i>Taiwanaleyrodes carpinii</i>	Betulaceae, Lauraceae, Myrsinaceae	Ericales, Fagales, Laurales	asterids, eurosids I, magnoliids
<i>Taiwanaleyrodes meliosmae</i>	Daphniphyllaceae, Lauraceae, Sabiaceae*, Sapindaceae	Laurales, Sapindales, Saxifragales	core eudicots, eudicots, eurosids II, magnoliids
<i>Tetraleurodes acaciae</i>	Anacardiaceae, Bursereaceae, Caprifoliaceae, Euphorbiaceae, Fabaceae, Geraniaceae, Moraceae, Myrtaceae, Nyctaginaceae, Rhamnaceae, Rosaceae, Salicaceae, Solanaceae, Urticaceae	Caryophyllales, Dipsacales, Fabales, Geraniales, Malpighiales, Myrtales, Rosales, Sapindales, Solanales	core eudicots, euasterids I, euasterids II, eurosids I, eurosids II, rosids
<i>Tetraleurodes andropogoni</i>	Annonaceae, Arecaceae, Clusiaceae, Connaraceae, Costaceae, Euphorbiaceae, Fabaceae, Linaceae, Loranthaceae, Poaceae, Rutaceae, Salicaceae	Arecales, Fabales, Magnoliales, Malpighiales, Oxalidales, Poales, Santalales, Sapindales, Zingiberales	commelinids, core eudi- cots, eurosids I, eurosids II, magnoliids
<i>Tetraleurodes banksiae</i>	Proteaceae	Proteales	eudicots
<i>Tetraleurodes bicolor</i>	Myrtaceae	Myrtales	rosids
<i>Tetraleurodes caulicola</i>	Fabaceae	Fabales	eurosids I
<i>Tetraleurodes confusa</i>	Fagaceae, Lauraceae, Magnoliaceae, Theaceae	Ericales, Fagales, Laurales, Magnoliales	asterids, eurosids I, magnoliids
<i>Tetraleurodes granulata</i>	Annonaceae, Combretaceae, Euphorbiaceae, Fabaceae, Loganiaceae, Rubiaceae	Fabales, Gentianales, Magnoliales, Malpighiales, Myrtales	euasterids I, eurosids I, magnoliids, rosids
<i>Tetraleurodes pluto</i>	Fabaceae	Fabales	eurosids I
<i>Tetraleurodes quercicola</i>	Fagaceae	Fagales	eurosids I
<i>Tetraleurodes selachidemata</i>	Anacardiaceae, Annonaceae, Bignoniaceae, Boraginaceae*, Combretaceae, Euphorbiaceae, Fabaceae, Loganiaceae, Rhamnaceae, Rubiaceae, Verbenaceae	Fabales, Gentianales, Lamiales, Magnoliales, Malpighiales, Myrtales, Rosales, Sapindales	euasterids I, eurosids I, eurosids II, magnoliids, rosids
<i>Tetraleurodes stirlingiae</i>	Proteaceae	Proteales	eudicots
<i>Tetraleurodes sulcistriatus</i>	Avicenniaceae, Clusiaceae, Moraceae, Myrtaceae	Lamiales, Malpighiales, Myrtales, Rosales	euasterids I, eurosids I, rosids
<i>Tetralicia ericae</i>	Ericaceae	Ericales	asterids
<i>Tetralicia graminicola</i>	Poaceae	Poales	commelinids
<i>Tetralicia iberica</i>	Ericaceae	Ericales	asterids
<i>Tetralicia tuberculata</i>	Verbenaceae	Lamiales	euasterids I
<i>Trialeurodes abutiloneus</i>	Acanthaceae, Apocynaceae, Asteraceae, Balsaminaceae, Boraginaceae*, Brassicaceae, Convolvulaceae, Cucurbitaceae, Euphorbiaceae, Fabaceae, Geraniaceae, Hamamelidaceae, Lamiaceae, Lythraceae, Malvaceae, Moraceae, Myrtaceae, Onagraceae, Oxalidaceae, Polygonaceae, Portulacaceae, Ranunculaceae, Rhamnaceae, Rosaceae, Rutaceae, Sapindaceae, Scrophulariaceae, Solanaceae, Ulmaceae, Urticaceae, Verbenaceae, Violaceae	Asterales, Brassicales, Caryophyllales, Cucurbitales, Ericales, Fabales, Gentianales, Geraniales, Lamiales, Malpighiales, Malvales, Myrtales, Oxalidales, Ranunculales, Rosales Sapindales, Saxifragales, Solanales	asterids, core eudicots, euasterids I, euasterids II, eudicots, eurosids I, eurosids II, rosids
<i>Trialeurodes bruneiensis</i>	Thelypteridaceae	§Aspleniales	
<i>Trialeurodes darwiniensis</i>	Euphorbiaceae	Malpighiales	eurosids I
<i>Trialeurodes dicksoniae</i>	Dicksoniaceae	§Cyatheales	
<i>Trialeurodes drewsi</i>	Fagaceae	Fagales	eurosids I
<i>Trialeurodes ericae</i>	Ericaceae	Ericales	asterids

Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Trialeurodes floridensis</i>	Anacardiaceae, Annonaceae, Bignoniaceae, Chrysobalanaceae, Convolvulaceae, Dilleniaceae*, Lauraceae, Fabaceae, Lythraceae, Malpighiaceae, Malvaceae, Meliaceae, Myrtaceae, Polygonaceae, Rhamnaceae, Rubiaceae, Rutaceae, Sapotaceae, Verbenaceae, Zygophyllaceae*	Caryophyllales, Ericales, Fabales, Gentianales, Lamiales, Laurales, Magnoliales, Malpighiales, Malvales, Myrtales, Rosales, Sapindales, Solanales	asterids, euasterids I, core eudicots, eurosids I, eurosids II, magnoliids, rosids
<i>Trialeurodes pergandei</i>	Bignoniaceae, Caprifoliaceae, Hydrangeaceae, Oleaceae, Rosaceae, Rutaceae, Smilacaceae	Cornales, Dipsacales, Lamiales, Liliales, Rosales, Sapindales	asterids, euasterids I, euasterids II, eurosids I, eurosids II, monocots
<i>Trialeurodes rex</i>	Undetermined fern		
<i>Trialeurodes vaporariorum</i>	Acanthaceae, Alismataceae, Anacardiaceae, Apocynaceae, Aquifoliaceae, Araceae, Araliaceae, Asteraceae, Balsaminaceae, Begoniaceae, Berberidaceae, Bignoniaceae, Boraginaceae*, Brassicaceae, Campanulaceae, Caprifoliaceae, Caryophyllaceae, Clusiaceae, Convolvulaceae, Cucurbitaceae, Cunoniaceae, Ebenaceae, Ericaceae, Euphorbiaceae, Fabaceae, Fagaceae, Garryaceae, Geraniaceae, Gesneriaceae, Grossulariaceae, Haloragaceae, Hamamelidaceae, Hydrangeaceae, Iridaceae, Juglandaceae, Lamiaceae, Lauraceae, Liliaceae, Loganiaceae, Lythraceae, Magnoliaceae, Malvaceae, Melianthaceae, Myrtaceae, Oleaceae, Onagraceae, Oxalidaceae, Papaveraceae, Passifloraceae, Pedaliaceae, Phytolaccaceae, Pittosporaceae, Platanaceae, Plumbaginaceae, Polemoniaceae, Polygalaceae, Polygonaceae, Primulaceae, Proteaceae, Ranunculaceae, Rhamnaceae, Rosaceae, Rubiaceae, Rutaceae, Salicaceae, Sapindaceae, Sapotaceae, Scrophulariaceae, Smilacaceae, Solanaceae, Thymelaeaceae, Tropaeolaceae, Ulmaceae, Urticaceae, Verbenaceae, Violaceae, Vitaceae*, ‡Zamiaceae	Alismatales, Apiales, Aquifoliales, Asparagales, Asterales, Brassicales, Caryophyllales, Cornales, Cucurbitales, †Cycadales, Dipsacales, Ericales, Fabales, Fagales, Garryales, Gentianales, Geraniales, Lamiales, Laurales, Liliales, Magnoliales, Malpighiales, Malvales, Myrtales, Oxalidales, Proteales, Ranunculales, Rosales, Sapindales, Saxifragales, Solanales	asterids, core eudicots, euasterids I, euasterids II, eudicots, eurosids I, eurosids II, magnoliids, monocots, rosids
<i>Trialeurodes variabilis</i>	Caricaceae, Euphorbiaceae, Polygonaceae, Rubiaceae, Rutaceae, Sapindaceae	Brassicales, Caryophyllales, Gentianales, Malpighiales, Sapindales	core eudicots, euasterids I, eurosids I, eurosids II
<i>Tuberaleyrodes machili</i>	Lauraceae	Laurales	magnoliids
<i>Vasdavidius cobarensis</i>	Poaceae	Poales	commelinids
<i>Vasdavidius concursus</i>	Poaceae	Poales	commelinids
<i>Vasdavidius indicus</i>	Poaceae	Poales	commelinids
<i>Vasdavidius setiferus</i>	Pandanaceae, Poaceae, Rutaceae	Poales	commelinids
<i>Venezaleurodes pisoniae</i>	Nyctaginaceae	Caryophyllales	core eudicots
<i>Viennotaleyrodes bicolorata</i>	Fabaceae	Fabales	eurosids I
<i>Viennotaleyrodes bosciae</i>	Brassicaceae	Brassicales	eurosids II
<i>Viennotaleyrodes curvisetosus</i>	Fabaceae	Fabales	eurosids I
<i>Viennotaleyrodes fallax</i>	Fabaceae	Fabales	eurosids I
<i>Viennotaleyrodes lacunae</i>	Fabaceae	Fabales	eurosids I
<i>Viennotaleyrodes megapapillae</i>	Fabaceae	Fabales	eurosids I
<i>Viennotaleyrodes platysepalii</i>	Fabaceae	Fabales	eurosids I

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Appendix E. (Continued).

Species	Host plant family	Host plant order	Host plant higher group
<i>Xenaleyrodes artocarp</i>	Clusiaceae, Lauraceae, Lecythidaceae, Moraceae, Musaceae, Sapindaceae, Sapotaceae, Verbenaceae	Ericales, Lamiales, Laurales, Malpighiales, Rosales, Sapindales, Zingiberales	asterids, commelinids, euasterids I, eurosids I, eurosids II, magnoliids
<i>Xenaleyrodes broughae</i>	Rutaceae	Sapindales	eurosids II
<i>Xenaleyrodes fauceregus</i>	Fabaceae	Fabales	eurosids I
<i>Yleyrodes isoberliniae</i>	Fabaceae	Fabales	eurosids I
<i>Zaphanera cappar</i>	Brassicaceae	Brassicales	eurosids II
<i>Zaphanera cyanotis</i>	Acanthaceae, Commelinaceae	Commelinales, Lamiales	commelinids, euasterids I
<i>Zaphanera niger</i>	Fabaceae	Fabales	eurosids I
<i>Zaphanera papyrocarpae</i>	Fabaceae	Fabales	eurosids I
<i>Zaphanera rhachisreticulata</i>	Fabaceae	Fabales	eurosids I
New genus 1 sp. 1	Piperaceae	Piperales	magnoliids
New genus 1 sp. 2	Piperaceae	Piperales	magnoliids
New genus 2 sp. 1	Rubiaceae	Gentianales	euasterids I
New genus 2 sp. 2	Host indet.		
<i>Aleurodicus cocois</i>	Anacardiaceae, Annonaceae, Arecaceae, Chrysobalanaceae, Euphorbiaceae, Fabaceae, Lauraceae, Malvaceae, Moraceae, Musaceae, Myrtaceae, Piperaceae, Polygonaceae, Rubiaceae	Arecales, Caryophyllales, Fabales, Gentianales, Laurales, Magnoliales, Malpighiales, Malvales, Myrtales, Piperales, Rosales, Sapindales, Zingiberales	commelinids, core eudicots, euasterids I, eurosids I, eurosids II, magnoliids, rosids
<i>Aleurodicus dispersus</i>	Acanthaceae, Anacardiaceae, Annonaceae, Apocynaceae, Araceae, Araliaceae, Arecaceae, Begoniaceae, Burseraceae, Clusiaceae, Combretaceae, Euphorbiaceae, Fabaceae, Lamiaceae, Lauraceae, Lecythidaceae, Moraceae, Musaceae, Myrtaceae, Orchidaceae, Polygonaceae, Rosaceae, Rubiaceae, Rutaceae, Sapotaceae, Solanaceae	Alismatales, Apiales, Arecales, Asparagales, Caryophyllales, Cucurbitales, Ericales, Fabales, Gentianales, Lamiales, Laurales, Magnoliales, Malpighiales, Myrtales, Rosales, Sapindales, Solanales, Zingiberales	asterids, commelinids, core eudicots, euasterids I, euasterids II, eurosids I, eurosids II, magnoliids, monocots, rosids
<i>Azureurodicus pentarthrus</i>	Rubiaceae	Gentianales	euasterids I
<i>Dialeurodicus cockerellii</i>	Myrtaceae	Myrtales	rosids
<i>Eudialeurodicus bodkimi</i>	Fabaceae	Fabales	eurosids I
<i>Octaleurodicus nitidus</i>	Arecaceae	Arecales	commelinids
<i>Stenaleyrodes vinsoni</i>	Arecaceae	Arecales	commelinids
<i>Synaleurodicus hakeae</i>	Proteaceae	Proteales	eudicots