Anatomical indicators of *Eucalyptus* spp. resistance to *Glycaspis brimblecombei* (Hemiptera: Aphalaridae)

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ABSTRACT

The total area of forest crops in Brazil is 9.55 million hectares, of which 7.5 million hectares are *Eucalyptus*. These crops are the most productive in the world, but may suffer losses due to exotic pests, including *Glycaspis brimblecombei* Moore (Hemiptera: Aphalaridae) found in Brazil since 2003. Interactions between *Eucalyptus* plants and insect pests may led to the selection of resistant genotypes. *Eucalyptus* species are either susceptible or resistant to this pest group, but the damage they suffer needs to be evaluated. The objective was to determine possible leaf anatomy indicators of different *Eucalyptus* species associated with *G. brimblecombei* infestations, focusing on plant resistance to this pest. The study was carried out with *Eucalyptus camaldulensis*, *Eucalyptus grandis*, *Eucalyptus saligna* and *Eucalyptus urophylla* saplings infested or not by *G. brimblecombei* eggs and nymphs. Eighteen anatomical characteristics of the leaves of these plants were analyzed. The number of stomata on the adaxial and abaxial sides and the glandular area in the central leaf vein are associated with greater or lesser infestation by *G. brimblecombei* in the *Eucalyptus* genotypes.

INTRODUCTION

Globally, forest crops cover around 294 million hectares (*Food & Agriculture Organization of the United Nations, 2020*). Brazil accounts for 9.55 million hectares of this area, with 7.5 million being *Eucalyptus*. Forest plantations in Brazil are among the most productive in
the world with 36.8 m³/ha year and with economic, social and environmental importance (Indústria Brasileira de Árvores, 2021). Native and exotic pests can compromise this productivity (Floris et al., 2020; Pereira et al., 2022). Eucalyptus plantations are established in large contiguous areas that provide a significant quantity of food and shelter for insect pests (Wingfield et al., 2008).

Exotic pests, introduced in the last two decades, are causing losses to the Brazilian forestry sector (Paine, Steinbauer & Lawson, 2011; Almeida et al., 2018). In 2003, Glycaspis brimblecombei Moore (Hemiptera: Aphalaridae) was reported in Brazil (Wilcken et al., 2015) and has reduced crop yields (Saliba et al., 2019). This insect feeds only on Eucalyptus species (Wilcken et al., 2015) and leaf rolling and deformation, “witch broom”, dieback and sooty mold are the main features of its infestation (Dittrich-Schroder et al., 2021).

Control methods for G. brimblecombei should focus on breeding and planting resistant eucalypt varieties, especially in areas with large G. brimblecombei populations (Jere et al., 2020). Different environmental conditions influence host plant susceptibility and infestation levels in the field (Ferreira-Filho et al., 2017; Bush, Slippers & Hurley, 2020).

Leaves, allelochemicals (tannins, phenols and waxes), glands that produce essential oils, often rich in terpenoids, hardness (sclorophilia), heterophilia (differentiation between young and mature leaves) and high regrowth of Eucalyptus plants can affect insect damage to this plant, with potential to select for resistant genotypes (Ohmart & Edwards, 1991).

Leaf anatomy is poorly studied and may allow us to understand pest infestations and the development of new tools for their management. Developing integrated psyllid management in Eucalyptus plantations depends on knowledge of plant/insect interactions. The objective of this study was to determine possible indicators based on leaf anatomy of four Eucalyptus species associated with G. brimblecombei infestations. These indicators may be useful in breeding programs for plant resistance to this pest.

MATERIALS AND METHODS

The study was carried out at the Universidade Estadual Paulista (FCA/UNESP) in Botucatu, São Paulo state, Brazil. Eucalyptus camaldulensis, E. grandis, E. saligna and E. urophylla were planted in 1.5 L pots with an autoclaved mixture of soil: sand: manure (2:1:1) and kept in a greenhouse for infestation with G. brimblecombei.

The Eucalyptus species were previously classified according to their response to G. brimblecombei with E. saligna and E. urophylla being resistant, E. grandis tolerant and E. camaldulensis susceptible to damage (Brennan et al., 2001; Pereira et al., 2013; Ribeiro et al., 2015).

Infestation of the Glycaspis brimblecombei on Eucalyptus plants

Glycaspis brimblecombei eggs and nymphs, collected in the field on Eucalyptus leaves, were placed on 25 cm high saplings of this plant in the laboratory. Each of the plants was infested with approximately 40 nymphs and two egg masses (more than 25 eggs each), weekly, for 4 weeks.

Twenty seedlings of each Eucalyptus species were used per treatment, with 10 plants (replications) infested with G. brimblecombei and another 10, as a control, free from the
pest. All the plants in the control were sprayed with systemic insecticide (acephate) and the others only with water, to evaluate the effects of mechanical action of the water.

**Anatomical characterization of *Eucalyptus* leaves**

*Eucalyptus camaldulensis*, *E. grandis*, *E. saligna* and *E. urophylla* leaves, infested or not, were analyzed. The samples were one to two leaves from the middle third of each eucalypt sapling, cut in three parts with the middle third analyzed. These samples were placed in formaldehyde + glacial acetic acid + 50% alcohol fixative solution (FAA-50) for 48 h and stored in 70% ethanol (*Johansen, 1940*). The samples were submersed into glyco-methacrylate resin (*Gerrits, 1991*) and cut, transversely, in a manual microtome, in the internervural region and in the central rib, with 15 to 25 μm thickness. The pieces were cleared, stained with acid fuchsin (*Brennan, Weinbaum & Pinney, 2001*) and toluidine blue pH 4.7 and mounted in synthetic resin (*O’Brien, Feder & Mccully, 1964*).

The thickness and the area with the epidermal, parenchymal and vascular leaf tissues were obtained with the computer program Image Tool 3.0 (UTHSCA) to evaluate the damage by *G. brimblecombei* on infested leaves. The quantitative anatomy was performed for three plants (replications) per species of *Eucalyptus* infested or not by *G. brimblecombei*. Eighteen variables for anatomical characterization of the leaf were evaluated.

**Quantitative variables of leaf anatomical characteristics**

The 18 variables related to leaf anatomy were: percentages of upper (%UE) and lower (%LE) epidermis, collenchyma (%Col), phloem (%Ph), xylem (%Xy), chlorenchyma (%Chl), gland (%Gl), and total cross-sectional area (mm$^2$) (CS) in the region of the central rib, thickness of the upper (TUE) and lower (TLE) epidermis, upper (TUPP) and lower (TLPP) palisade parenchyma, spongy parenchyma (TSP), leaf (TL), mesophyll (TM), the mean area of a gland (MGA), and number of stomata/mm$^2$ of the upper (NUS) and lower (NLS) surfaces in the internervre region (*Sambugaro et al., 2004*).

**Statistical analysis**

The anatomical leaf characterization data were subjected to multivariate statistical tests of Cluster Analysis and Principal Component Analysis (PCA) (*Sneath & Sokal, 1973*) to verify the discriminatory capacity of the quantitative anatomical variables obtained by the measurements of the different leaf tissues, and the means compared by the Tukey test at 5% probability, using R Studio software.

**RESULTS**

**Damage by *Glycaspis brimblecombei***

The infestation of *G. brimblecombei* was constant with low plant mortality. *Eucalyptus camaldulensis* was more infested than *E. urophylla* and *E. grandis* and all *G. brimblecombei* nymphs died in the first instars on *E. saligna* without development on plants of this species. Sooty mold developed on *G. brimblecombei* lerp. The occurrence of leaf spot from
Teratosphaeria epicoccoides was observed on *E. camaldulensis*, *E. grandis* and *E. urophylla* and with greater damage to *E. saligna*.

### Anatomical leaf characterization

The percentage of upper and lower epidermis in the region of the central vein, percentage of collenchyma, thickness of the upper and lower epidermis in the internervure region and thickness of the spongy parenchyma was similar between the *Eucalyptus* species (Table 1). The percentage of chlorenchyma was lowest and that of phloem, xylem and the mean gland area in the central vein region was highest in *E. grandis* leaves than in the other *Eucalyptus* species (Table 1).

The cluster analysis, based on the discriminatory capacity of the quantitative anatomical variables, that is, comparing the elements according to the presence or absence of certain characteristics separated the *Eucalyptus* species into two groups (Fig. 1) based on the low level of 0.32 on the similarity distance scale: group 1—*E. saligna*, *E. urophylla* and *E. grandis*; group 2—*E. camaldulensis*, *E. saligna* and *E. urophylla*.

The graph dispersion of the four *Eucalyptus* species showed *E. saligna*, *E. urophylla* and *E. grandis* forming group 1 and *E. camaldulensis* group 2 for the principal component analysis with contrast between these species (Y1 and Y2) (Fig. 2). The graphic dispersion of the PCA and the dendrogram of the cluster analysis, grouped the four *Eucalyptus* species as follows:

| Variable                              | *E. camaldulensis* | *E. grandis* | *E. urophylla* | *E. saligna* |
|---------------------------------------|--------------------|--------------|----------------|--------------|
| Upper epidermis (%)                  | 2.83 ± 0.67a       | 2.76 ± 1.00a | 3.80 ± 0.57a   | 3.39 ± 0.75a |
| Lower epidermis (%)                  | 2.42 ± 0.36a       | 3.00 ± 0.52a | 4.11 ± 0.69a   | 3.90 ± 0.71a |
| Collenchyma (%)                      | 33.46 ± 10.69a     | 29.44 ± 4.17a| 31.09 ± 4.30a  | 35.70 ± 6.42a|
| Phloem (%)                           | 13.90 ± 3.86a      | 24.74 ± 3.52b| 14.97 ± 6.39a  | 17.41 ± 4.34a|
| Xylem (%)                            | 16.40 ± 0.52a      | 19.88 ± 3.61b| 12.50 ± 3.78a  | 10.22 ± 2.46a|
| Chlorophyll parenchyma (%)           | 30.12 ± 4.41a      | 16.03 ± 3.43b| 31.36 ± 5.22a  | 28.11 ± 3.27a|
| Glands (%)                           | 0.87 ± 0.63a       | 4.15 ± 1.39c | 2.17 ± 2.59b   | 1.26 ± 1.05b |
| Total cross-sectional area (mm²)     | 0.61 ± 0.05a       | 0.57 ± 0.03a | 0.31 ± 0.02a   | 0.46 ± 0.03a |
| Total of the upper epidermis (µm)    | 15.94 ± 3.42a      | 18.44 ± 3.39b| 16.56 ± 2.43b  | 17.19 ± 2.82b|
| Total of the lower epidermis (µm)    | 15.31 ± 3.55a      | 12.19 ± 2.80a| 15.31 ± 2.95a  | 13.75 ± 3.56a|
| Upper palisade parenchyma            | 97.19 ± 12.16a     | 70.94 ± 12.65b| 70.00 ± 11.02b | 58.12 ± 3.37b|
| Lower palisade parenchyma            | 78.44 ± 12.76a     | 0.00 ± 0.00  | 0.00 ± 0.00    | 0.00 ± 0.00  |
| Total of spongy parenchyma (µm)     | 103.75 ± 26.28a    | 121.25 ± 14.39a| 102.81 ± 10.47a| 117.19 ± 14.46a|
| Mesophyll thickness                  | 279.37 ± 77.14a    | 192.19 ± 59.23b| 172.81 ± 32.00b| 175.31 ± 48.85b|
| Leaf thickness (µm)                  | 310.62 ± 52.93a    | 222.81 ± 50.54b| 204.37 ± 23.41b| 206.25 ± 58.97b|
| Mean area of a gland                 | 7.65 ± 2.60a       | 11.68 ± 2.19a| 6.52 ± 0.73a   | 7.39 ± 0.92a |
| Number of stomata of the upper surfaces | 231.73 ± 20.57a  | 1.37 ± 0.06b | 0.00 ± 0.00   | 0.00 ± 0.00 |
| Number of stomata of the lower surfaces     | 256.68 ± 23.89a   | 500.39 ± 35.71b| 527.55 ± 21.01b| 557.06 ± 29.43b|

*Note:* Averages followed by the same lowercase letter per line do not differ by Tukey’s test (*p* ≤ 0.05).

*Teratosphaeria epicoccoides* was observed on *E. camaldulensis*, *E. grandis* and *E. urophylla* and with greater damage to *E. saligna*.

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**Table 1** Values of the 18 quantitative anatomical variables for *Eucalyptus camaldulensis*, *Eucalyptus grandis*, *Eucalyptus urophylla* and *Eucalyptus saligna* leaves infested by *Glycaspis brimblecombei* (Hemiptera: Aphalaridae) in a greenhouse.
species into two main groups, based on the 18 quantitative anatomical characteristics of the Eucalyptus leaves (Fig. 2).

The correlation coefficients among the 18 quantitative anatomical variables of the Eucalyptus leaves and the first two principal components (Y1 and Y2) were found to be thickness variables of the lower palisade parenchyma, mesophyll, leaf, upper palisade parenchyma, upper epidermis, as well as the number of stomata of the upper and lower surfaces. These were the main variables that served to discriminate the four Eucalyptus species, based on the first principal component (Y1) (Table 2). The discriminatory power of the absolute value of Y1 for these variables, was high. The information retained for the second principal component (Y2) was low (26.43%), which meant that analysis of this component was unsatisfactory. The combined analysis of the first principal component (Table 2) and the graphic dispersion (Fig. 2) showed that the number of stomata on the lower side, percentage of lower epidermis, thickness of the upper epidermis, and
percentage of gland in the central vein of the group 2 species (E. camaldulensis) were lower than those of the group 1 species (E. saligna, E. grandis and E. urophylla) (Table 1).

The values of the thickness characteristics of the upper and lower palisade parenchyma, mesophyll and leaf and the number of stomata on the upper surface of E. camaldulensis were higher than those for other species. The E. camaldulensis leaf profile was classified (Fig. 3). Signs of stylet insertion by G. brimblecombei nymphs were found in E. camaldulensis leaf sections, passing through the collenchyma, near the central leaf vein and the palisade parenchyma (Fig. 4).

**DISCUSSION**

*Glycaspis brimblecombei* damages young plants, from 6 months to mature ones, up to cutting age, causing serious damage throughout its cycle (Saliba *et al.*, 2019). The damage in younger plantations, between 6 months up to 2 years, results in greater losses when compared to more mature plantations (5 years or more) (Wardlaw *et al.*, 2018). *Glycaspis brimblecombei* is a sucking insect and its nymphs produce a large amount of honeydew, causing the development of sooty mold (Reguia & Peris-Felipo, 2013). *Teratosphaeria epicoccoides* on *Eucalyptus* leaves, with greater damage to *E. saligna*, is generally associated with stressed plants (Andjic *et al.*, 2019).

The more intense *G. brimblecombei* infestation on *E. camaldulensis* than on other species tested here is related to its susceptibility to this insect (Firmino-Wincker *et al.*, 2009; Ribeiro *et al.*, 2015). The lack of development of *G. brimblecombei* nymphs on *E. saligna* plants is due to the resistance related to epicuticular wax on the leaves, reducing the presence of eggs and nymphs and the severity of *G. brimblecombei* infestation (Brennan *et al.*, 2001).

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**Table 2** Correlations between the 18 quantitative anatomical variables retained and accumulated in $Y_1$ and $Y_2$ for the leaf of *Eucalyptus camaldulensis*, *Eucalyptus grandis*, *Eucalyptus urophylla* and *Eucalyptus saligna* and the first two main components ($Y_1$ and $Y_2$).

| Original variables | $Y_1$ | $Y_2$ | Original variables | $Y_1$ | $Y_2$ |
|-------------------|-------|-------|-------------------|-------|-------|
| TLPP              | 0.9987| 0.0492| %Ph               | −0.5640| 0.8062|
| NUS               | 0.9984| 0.0548| CS                | 0.5582 | 0.7198|
| TL                | 0.9772| 0.2124| TSP               | −0.5534| 0.5648|
| NLS               | −0.9762| −0.1881| TLE               | 0.5527 | −0.7287|
| TM                | 0.9758| 0.2186| %UE               | −0.4584| −0.8365|
| TUPP              | 0.9204| 0.2303| %Chl              | 0.3943 | −0.9055|
| %LE               | −0.7627| −0.6427| %Col              | 0.2953 | −0.6390|
| TUE               | −0.7122| 0.6655| MGA               | −0.2374| 0.9642|
| %Gl               | −0.6077| 0.7445| %Xy               | 0.2077 | 0.9431|
| %Retained         | 70.17 | 26.43 | %Accumulated      | 70.17 | 96.6  |

**Note:**  
TLPP, lower palisade parenchyma thickness; NUS, number of stomata/mm$^2$ of upper face; TL, leaf thickness (μm); NLS, number of stomata/mm$^2$ of the lower face in the internervural region; TM, mesophyll thickness; TUPP, upper palisade parenchyma thickness; %LE, Percentage of lower epidermis; TUE, thickness of the upper epidermis; %Gl, gland; %Ph, phloem; CS, total cross-sectional area (mm$^2$) in the central rib region; TSP, spongy parenchyma thickness (μm); TLE, lower palisade parenchyma thickness; %UE, percentage of upper epidermis; %Chl, chlorenchyma; %Col, collenchyma; MGA, mean gland area; %Xy, xylem.
The similar percentage of epidermis in the central vein region, collenchyma and epidermis thickness in the internervural region, and thickness of spongy parenchyma for the resistant and susceptible *Eucalyptus* species (*Brennan et al., 2001; Pereira et al., 2013; Ribeiro et al., 2015*), indicates that these anatomical variables are not associated with the plant resistance or susceptibility to *G. brimblecombei*. The percentage of chlorenchyma, responsible for photosynthesis, is lower in *E. grandis* leaves than in the other *Eucalyptus* species. This is related to a reduction of leaf area, similar to that caused by *Costalimaia ferruginea* (Coleoptera: Chrysomelidae) on shoots and apical parts of *Eucalyptus*, which may reduce chlorenchyma, impairing plant development (*Santos, Gonçalves & Silva, 2016*). The higher percentage of glands on *E. grandis* leaves in the central vein region, and phloem and xylem in the central vein than in other species may be
related to the presence and production of phenolic compounds in the epidermis (Santos et al., 2008), as a result of plant defense to insect pests, including G. brimblecombei.

Differences in the number of stomata on the upper surface, and thickness of the upper and lower palisade parenchyma on *E. camaldulensis* due to stomata on the adaxial surface and a double layer of palisade parenchyma on both sides of its leaves. The single layer of palisade parenchyma was found only on the adaxial surface of the other species (James & Bell, 1995).

The palisade parenchyma probably does not confer resistance on *Eucalyptus* spp. to *G. brimblecombei*, because this structure is duplicated on the adaxial and abaxial surfaces of *E. camaldulensis* leaves and single in the adaxial surface of *E. grandis*, *E. saligna* and *E. urophylla*, as well as thicker, on both sides, in *E. camaldulensis* than in the other species. The signs of stylet insertion by *G. brimblecombei* nymphs through the *E. camaldulensis* leaf sections indicates that they passed through the parenchyma cells rather than between them. Cell-degrading proteins such as amylase, cellulase, pectinase and pectinesterase enable stylet entry into the plant tissue (Wu et al., 2021). Stomata are absent or in low numbers in the adaxial surface of *E. grandis*, *E. saligna* and *E. urophylla*, whereas they are
present on *E. camaldulensis* leaf side surfaces. The total number of stomata is similar between these species, but this may explain the similar infestation on the abaxial and adaxial surfaces of *E. camaldulensis* compared to *E. urophylla*, with greater infestation on the abaxial surfaces. Styles of *G. brimblecombei* nymphs penetrated the mesophyll, crossing between the guard cells of the stomata, similar to that observed for this insect in *E. globulus* ([Brennan & Weinbaum, 2001a, 2001b]) and, for this reason, stomata on both sides of *E. camaldulensis* may confer greater susceptibility to *G. brimblecombei*.

Defense strategies of *Eucalyptus* trees for insects include physical barriers and constitutive and inducible chemical defenses ([Patton et al., 2017](#)). The concentration and variability of terpenes, the presence of specific compounds ([Silveira et al., 2021](#)), amounts of epicuticular wax in the leaves, and the occurrence of antibiosis, related to longer insect development stages or life cycles, and/or antixenosis resistance, related to extended developmental stages due to lower food intake of insects, are characteristics normally associated with *Eucalyptus* resistance to *G. brimblecombei* ([Pereira et al., 2020](#)).

The proportional area and number of stomata occupying the epidermis may also be important for *G. brimblecombei* nymph infestation and to explain *E. camaldulensis* susceptibility to this pest. The thinner epidermis of the adaxial surface and lower percentage of epidermal tissue on the abaxial surface of *E. camaldulensis* leaves are possibly related to the higher susceptibility to *G. brimblecombei*. This is a pioneering study evaluating anatomical foliar indicators in relation to *Eucalyptus* pests, and allows us to better understand pest infestation patterns, and concomitantly, the morphological characteristics that normally confer resistance, such as waxy coating, trichoids, and stomata in these plants.

**CONCLUSIONS**

The number of stomata in the adaxial and abaxial leaf surfaces and percentage of gland area in the central vein of the leaves are related to the resistance or susceptibility of *Eucalyptus* plants to *G. brimblecombei*.

*Eucalyptus grandis*, *E. urophylla* and *E. saligna*, with higher values of the leaf characteristics evaluated, may be considered resistant or moderately resistant to *G. brimblecombei*.

**ACKNOWLEDGEMENTS**

Dr. Phillip John Villani (University of Melbourne, Australia) revised and corrected the English language used in this manuscript.

**ADDITIONAL INFORMATION AND DECLARATIONS**

**Funding**

The study was financially supported by the following Brazilian agencies “Coordenação de Aperfeiçoamento de Pessoal de Nível Superior-Brasil (CAPES-Finance Code 001)”, “Fundação de Amparo à Pesquisa do Estado de Minas Gerais (FAPEMIG)” and “Programa Cooperativo sobre Proteção Florestal/PROTEF do Instituto de Pesquisas e Estudos...”
The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Grant Disclosures
The following grant information was disclosed by the authors:
Coordenação de Aperfeiçoamento de Pessoal de Nível Superior-Brasil: 001.
Fundação de Amparo à Pesquisa do Estado de Minas Gerais.
Programa Cooperativo sobre Proteção Florestal/PROTEF do Instituto de Pesquisas e Estudos Florestais/IPEF.

Competing Interests
The authors declare that they have no competing interests.

Author Contributions
- Fernando Henrique Moreno de Oliveira Del Piero conceived and designed the experiments, performed the experiments, analyzed the data, prepared figures and/or tables, and approved the final draft.
- Carlos Frederico Wilcken conceived and designed the experiments, performed the experiments, analyzed the data, prepared figures and/or tables, and approved the final draft.
- Maurício Magalhães Domingues analyzed the data, prepared figures and/or tables, authored or reviewed drafts of the paper, and approved the final draft.
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- Roberto Antonio Rodella conceived and designed the experiments, performed the experiments, analyzed the data, prepared figures and/or tables, and approved the final draft.
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- José Eduardo Serrão analyzed the data, prepared figures and/or tables, authored or reviewed drafts of the paper, and approved the final draft.
- José Cola Zanuncio analyzed the data, prepared figures and/or tables, authored or reviewed drafts of the paper, and approved the final draft.

Data Availability
The following information was supplied regarding data availability:
- The raw measurements are available in the Supplemental Files.

Supplemental Information
Supplemental information for this article can be found online at http://dx.doi.org/10.7717/peerj.13346#supplemental-information.
REFERENCES

Almeida KEC, Silva JGS, Silva IMA, Costa AL, Laia ML. 2018. Ecophysiological analysis of Eucalyptus camaldulensis (Dehnh) submitted to attack from Thaumastocoris peregrinus (Carpintero & Dellape). Revista Árvore 42(1):1 DOI 10.1590/1806-90882018000100020.

Andjic V, Carnegie AJ, Pegg GS, Hardy GSJ, Maxwell A, Crous PW, Pérez C, Wingfield MJ, Burgess TJ. 2019. 23 years of research on Teratosphaeria leaf blight of Eucalyptus. Forest Ecology and Management 443(5):1–27 DOI 10.1016/j.foreco.2019.04.013.

Brennan EB, Levison W Jr, Hrusa GF, Weinbaum SA. 2001. Resistance of Eucalyptus species to red gum lerp psyllid (Glycaspis brimblecombei) (Homoptera: Psyllidae) in San Francisco Bay area. The Pan-Pacific Entomologist 77:249–253.

Brennan EB, Weinbaum SA. 2001a. Psyllid responses to colored sticky traps and the colors of juvenile and adult leaves of the heteroblastic host plant Eucalyptus globulus. Environmental Entomology 30(2):365–370 DOI 10.1603/0046-225X-30.2.365.

Brennan EB, Weinbaum SA. 2001b. Stylet penetration and survival of three psyllid species on adult leaves and 'waxy' and 'de-waxed' juvenile leaves of Eucalyptus globulus. Enomologia Experimentalis et Applicata 100(3):355–363 DOI 10.1046/j.1570-7458.2001.00883.x.

Brennan EB, Weinbaum SA, Pinney K. 2001. A new technique for studying the stylet the tracks of homopteran insects in hand-sectioned plant tissue under light or epifluorescence microscopy. Biotechnic & Histochemistry 76(2):59–66 DOI 10.1080/bih.76.2.59.66.

Bush SJ, Slippers B, Hurley BP. 2020. Eucalypt susceptibility towards the invasive Glycaspis brimblecombei Moore (Hemiptera: Aphabetaria) in South Africa. Southern Forests: A Journal of Forest Science 82(3):243–252 DOI 10.2989/20702620.2020.1824556.

Dittrich-Schroder G, Garnas J, Arriagada-Cares D, Ahumada R, Hurley BP, Lawson SA, Slippers B. 2021. Global diversity and introduction history of Glycaspis brimblecombei reflects a history of bridgeheads and distinct invasions. Frontiers in Forests and Global Change 4:2357 DOI 10.3389/ffgc.2021.783603.

Ferreira-Filho PJ, Wilcken CF, Masson MV, Tavares WS, Guerreiro JC, Carmo JB, Prado EP, Zanuncio JC. 2017. Influence of temperature and rainfall on the population dynamics of Glycaspis brimblecombei and Psyllaephagus bletteus in Eucalyptus camaldulensis plantations. Revista Colombiana de Entomologia 43:1–6 DOI 10.25100/socolen.v43l1.6638.

Firmino-Wincker DC, Wilcken CF, Matos CAO, Oliveira NC. 2009. Biologia do psilídeo-de-concha Glycaspis brimblecombei Moore (Hemiptera: Psyllidae) em Eucalyptus spp. Revista Brasileira de Entomologia 53(1):144–146 DOI 10.1590/S0085-56262009000100030.

Floris I, Pusceddu M, Mannu R, Buffa F, Quaranta M, Satta A. 2020. Impact of sap-sucking insect pests (Blastosylla occidentalis Taylor and Glycaspis brimblecombei Moore, Hemiptera: Psyllidae) on unifloral eucalyptus honey. Annals of Silvicultural Research 44:66–70 DOI 10.12899/asr-1848.

Food and Agriculture Organization of the United Nations. 2020. Faostat. Available at http://www.fao.org/faostat/en/#data/RL (accessed 26 March 2020).

Gerrits PO. 1991. The application of glycol metacrylate in histotechnology: some fundamental principles. Germany: LeicaGmbH, 80.

Indústria Brasileira de Árvores. 2021. IBÁ annual report. Available at https://iba.org/datafiles/publicacoes/relatorios/relatorioiba2021-compactado.pdf?utm_source=akna&utm_medium=email&utm_campaign=Iba-lanca-Relatorio-Anual-2021 (accessed 1 February 2022).

James SA, Bell DT. 1995. Morphology and anatomy of leaves of Eucalyptus camaldulensis clones: variation between geographically separated locations. Australian Journal of Botany 43(4):415–433 DOI 10.1071/BT9950415.
Jere V, Mhango J, Njera D, Jenya H. 2020. Infestation of *Glycaspis brimblecombei* (Hemiptera: Psyllidae) on three *Eucalyptus* species in selected ecological zones in Malawi. *African Journal of Ecology* 58:251–259 DOI 10.1111/aje.12686.

Johansen DA. 1940. *Plant microtechnique*. New York and London: McGraw-Hill Book Company, Inc, 523.

O’Brien TP, Feder N, McCully ME. 1964. Polychromatic staining of plant cell walls by Toluidine Blue O. *Protoplasma* 59(2):367–373 DOI 10.1007/BF01248568.

Ohmart CP, Edwards PB. 1991. Insect herbivory on *Eucalyptus*. *Annual Review of Entomology* 36(1):637–657 DOI 10.1146/annurev.en.36.010191.003225.

Paine TD, Steinbauer MJ, Lawson SA. 2011. Native and exotic pests of *Eucalyptus*: a worldwide perspective. *Annual Review of Entomology* 56(1):181–201 DOI 10.1146/annurev-ento-120709-144817.

Patton MF, Arena GD, Salminen JP, Steinbauer MJ, Casteel CL. 2017. Transcriptome and defence response in *Eucalyptus camaldulensis* leaves to feeding by *Glycaspis brimblecombei* Moore (Hemiptera: Aphalaridae): a stealthy psyllid does not go unnoticed. *Austral Entomology* 57(2):247–254 DOI 10.1111/aen.12319.

Pereira JM, Baldin ELL, Soliman EP, Wilcken CF. 2013. Attractiveness and oviposition preference of *Glycaspis brimblecombei* Moore in *Eucalyptus* spp. *Phytoparasitica* 41(2):117–124 DOI 10.1007/s12600-012-0268-7.

Pereira JM, Baldin ELL, Soliman EP, Wilcken CF. 2020. Development of the red gum lerp psyllid *Glycaspis brimblecombei* (Hemiptera: Aphalaridae) in *Eucalyptus* spp. *Scientia Forestalis* 48(127):e3283 DOI 10.18671/scifor.v48i127.18.

Pereira JM, Santos TTM, Soliman EP, Dias TKR, Baldin ELL, Wilcken CF. 2022. Survival and performance of *Sarsina violascens* (Lepidoptera: Lymantriidae) larvae on *Eucalyptus* species and hybrids. *Phytoparasitica* 50(1):13–20 DOI 10.1007/s12600-021-00933-9.

Reguia K, Peris-Felipo FJ. 2013. *Glycaspis brimblecombei* Moore, 1964 (Hemiptera: Psyllidae) invasion and new records in the Mediterranean area. *Biodiversity Journal* 4:501–506.

Ribeiro ZA, Souza BHS, Costa EN, Mendes JEP, Mafia RG, Júnior ALB. 2015. *Glycaspis brimblecombei* Moore, 1964 (Hemiptera: Psyllidae) on *Eucalyptus*: oviposition non-preference and antibiosis. *Euphytica* 202(2):285–295 DOI 10.1007/s10681-014-1298-7.

Saliba IL, Lunz AM, Batista TF, Schwartz G, Queiroz DL. 2019. First record of *Glycaspis brimblecombei* (Moore, 1964) and *Blastopsylla occidentalis* (Taylor, 1985) (Hemiptera, Aphalaridae) in *Eucalyptus* plantations in State of Pará, Brazil. Embrapa Florestas-Nota Técnica/Nota Científica (ALICE). Available at https://www.alice.cnptia.embrapa.br/bitstream/doc/1116971/1/2019DalvaECFirstrecord.pdf (accessed 18 April 2020).

Sambubaro R, Furtado EL, Rodella RA, Mattos CRR. 2004. Anatomia foliar de seringueira (*Hevea* spp.) e desenvolvimento da infecção por *Microcyclus ulei*. *Summa Phytopathologica* 30:51–56.

Santos RS, Gonçalves R, Silva NDA. 2016. Primeiro registro do besouro-amarelo-do-eucalipto em plantio de eucalipto no Estado do Acre. *Revista Ceres* 63(4):584–587 DOI 10.1590/0034-737X201663040020.

Santos LDT, Thadeo M, Iarema L, Meira RMSA, Ferreira FA. 2008. Foliar anatomy and histochemistry in seven species of *Eucalyptus*. *Revista Árvore* 32(4):769–779 DOI 10.1590/S0100-67622008000400019.

Silveira AC, Siqueira GLA, Mayer FM, Lazzarotto SRS, Miguel OG, Zini CA, Queiroz DL, Lazzarotto M. 2021. Thermal tool to evaluate essential oil composition of different *Eucalyptus*
genotypes in relation to *Glycaspis brimblecombei* susceptibility (Hemiptera: Aplalaridae). *Journal of Thermal Analysis and Calorimetry* 30:1–9 DOI 10.1007/s10973-021-11027-3.

Sneath PHA, Sokal RR. 1973. *Numerical taxonomy*. San Francisco: W.H. Freeman, 573.

Wardlaw T, Cameron N, Carnegie A, Lawson S, Venn T. 2018. Costs and benefits of a leaf beetle Integrated Pest Management (IPM) program. I. Modelling changes in wood volume yields from pest management. *Australian Forestry* 81(1):46–52 DOI 10.1080/00049158.2018.1425969.

Wilcken CF, Firmino-Winckler DC, Dal Pogetto MHFA, Dias TKR, Lima ACV, Sá LD, Ferreira Filho PJ. 2015. Psilídeo de concha do eucalipto, *Glycaspis brimblecombei* Moore. In: Vilela Filho E, Zucchi RA, eds. *Pragas Introduzidas no Brasil: Insetos e Ácaros*. Piracicaba: FEALQ, 883–897.

Wingfield MJ, Slippers B, Hurley BP, Coutinho TA, Wingfield BD, Roux J. 2008. Eucalypt pests and diseases: growing threats to plantation productivity. *Southern Forests: A Journal of Forest Science* 70(2):139–144 DOI 10.2989/SOUTH.FOR.2008.70.2.9.537.

Wu ZZ, Qu MQ, Chen MS, Lin JT. 2021. Proteomic and transcriptomic analyses of saliva and salivary glands from the Asian citrus psyllid, *Diaphorina citri*. *Journal of Proteomics* 238(54):104136 DOI 10.1016/j.jprot.2021.104136.