Relative host plant species use by the lantana biological control agent *Aconophora compressa* (Membracidae) across its native and introduced ranges

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**Abstract**

*Aconophora compressa* Walker (Hemiptera: Membracidae) was released in 1995 against the weed *Lantana camara* in Australia, and is now found on multiple host plant species. The intensity and regularity at which *A. compressa* uses different host species was quantified in its introduced Australian range and also its native Mexican range. In Australia, host plants fell into three statistically defined categories, as indicated by the relative rates and intensities at which they were used in the field. Fiddlewood (*Citharexylum spinosum* L.; Verbenaceae) was used much more regularly and at higher densities than any other host sampled, and alone made up the first group. The second group, lantana (*Lantana camara* L.; Verbenaceae; pink variety) and geisha girl (*Duranta erecta* L.; Verbenaceae), were used less regularly and at much lower densities than fiddlewood. The third group, Sheena's gold (another variety of *D. erecta*), jacaranda (*Jacaranda mimosifolia* D. Don: Bignoniaceae) and myoporum (*Myoporum acuminatum* R. Br.: Myoporaceae), were used infrequently and at even lower densities. In Mexico, the insect was found at relatively low densities on all hosts relative to those in Australia. Densities were highest on *L. urticifolia*, *D. erecta* and *Tecoma stans* (L.) Juss. ex Kunth (Bignoniaceae), which were used at similar rates to one another. It was found also on a few other verbenaceous and non-verbenaceous host species but at even lower densities. The relative rate at which *Citharexylum* spp. and *L. urticifolia* were used could not be assessed in Mexico because *A. compressa* was found on only one plant of each species in areas where these host species co-occurred. The low rate at which *A. compressa* occurred on fiddlewood in Mexico is likely to be an artefact of the short-term nature of the surveys or differences in the suites of *Citharexylum* and *Lantana* species available there. These results provide further incentive to insist on structured and quantified surveys of non-target host use in the native range of potential biological control agents prior to host testing studies in quarantine.

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1. Introduction

Field surveys of herbivorous species on their host plants are essential for identifying candidates for weed biological control agents (Goolsby et al., 2006; Sheppard et al., 2006a). When host-testing results suggest the biocontrol agent is host specific, its release, once approved, is usually made on the assumption it will attack only the target weed. Unanticipated non-target effects have the potential to be devastating to the environment, damage the reputation of biological control (Simberloff and Stiling, 1996; Louda et al., 2003) and jeopardise permission for future releases of other agents (Sheppard et al., 2006b). Past mistakes of this nature do, however, provide a basis for understanding how and why such errors were made and should provide insights to improve the process by which biocontrol agents are selected and sanctioned for release (Briese, 2005).

Host specificity testing of candidate biocontrol agents relies mostly on data collected from glasshouse and laboratory studies in which a range of potential hosts are exposed to the herbivore species of interest. Plant species that are economically and ecologically important, and which are deemed to be at potential risk, must be included. So, too, must plant species, or close relatives, with which that agent has been associated in its native range. All other available information on the herbivore species, in relation to its association with the target weed (and other host species), is normally gathered prior to host testing, and this includes data from insect collections, published literature and surveys and personal knowledge of field collectors. This approach to host testing has several recognised shortcomings, including the inadvertent
Aconophora compressa was introduced into Australia from Mexico in 1995 for the biological control of lantana (Lantana camara L.: Verbenaceae) (Palmer et al., 1996) but in Australia it is mostly associated with fiddlewood (Citharexylum spinosum L.: Verbenaceae) (Dhileepan et al., 2006; Manners and Walter, 2009; Manners et al., 2010), a tree native to the Caribbean. The available data for A. compressa from field and laboratory studies indicate that fiddlewood is a primary host, whereas lantana and the varieties of D. erecta, (i.e. geisha girl and Sheena’s gold) are secondary hosts (Dhileepan et al., 2006; Manners and Walter, 2009; Manners et al., 2010).

A. compressa is ideal for investigating aspects of the herbivore-host interaction that are relevant to the pre-release evaluation of biocontrol agents because it uses multiple hosts, much of its basic biology on a number of hosts is known (Manners and Walter, 2009; Manners et al., 2010), and it has already been introduced into a new area where it interacts with hosts differently from the way originally anticipated. Furthermore, a number of plant species exist in both Mexico and Australia for which the insect plant interaction has been studied in Australia, including D. erecta (variety not known in Mexico), jacaranda (Jacaranda mimosifolia), lantana (L. camara in Australia is thought to be most closely related to L. urticifolia from Mexico (Scott 2002), although recent evidence suggests that it may be more closely related to lantana from Venezuela and the Dominican Republic (R. Watts, CSIRO, Plant Industries, Canberra, personal communication). In addition, many species of Citharexylum exist in Mexico, although C. spinosum has not been recorded there (Moldenke, 1942). Meaningful comparisons of the host plant relationships of A. compressa can thus be made across the native and introduced ranges.

In particular, the following questions were addressed through quantified field sampling in Australia and Mexico. Does field host use in the native range correlate to that in the introduced range of A. compressa? Could the quantified host use of A. compressa in Mexico across all host plants have thus been used to predict the observed behaviour of this species after its release in Australia and, if not, what are the possible explanations? Answers to these questions should improve our understanding of insect–host relationships in general and lead to recommendations for host testing methods that should reduce the risk associated with releasing an organism into a new environment.
2. Methods

2.1. Surveys in Australia

2.1.1. General methods

A survey was conducted to quantify the regularity and intensity at which *A. compressa* used different host species in South East Queensland (SEQ), Australia. Six plant taxa were monitored at each of five sites, Bracken Ridge (27°19’33”S 153°01’28”E), St Lucia (27°29’54”S 153°00’04”E), Sherwood (27°31’53”S 152°58’53”E), Durack (27°35’19”S 153°00’04”E) and Mt Tamborine (27°55’20”S 153°12’21”E) (Fig. 1). Four taxa were verbenaceous (fiddlewood, lantana – pink-edged red variety which is common in SEQ, and two varieties of *D. erecta*, namely geisha girl and Sheena’s gold) and two were non-verbenaceous (jacaranda and myoporum, *Myoporum acuminatum*). Each month at each site, the numbers of adults, females attending egg batches, early instar nymphs (first to third) and late instar nymphs (fourth and fifth) were counted on 15 randomly selected branches on each of 15 trees of each taxon that had been marked with aluminium tags. Specific branches were not marked or re-sampled each month, although re-sampling may

![Fig. 2. Mean number of each stage of *Aconophora compressa* per branch (±SE) (labels across top of figure) on six host taxa (labelled in each row on the extreme right) sampled monthly in South East Queensland, Australia (*n* = 15 branches on each of 15 trees per plant species per month at each of five sites). Different scales are used for different host plants and stages of insect. Sample number (*x*-axis) represents successive months between August 2006 and November 2007, except October 2007.](image)

<table>
<thead>
<tr>
<th>Treatment</th>
<th>t-Value</th>
<th>df</th>
<th>P-value</th>
<th>Total proportion of branches</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intercept (fiddlewood)</td>
<td>15.82</td>
<td>226</td>
<td>&lt;0.0001a</td>
<td>0.236</td>
</tr>
<tr>
<td>Lantana</td>
<td>-8.83</td>
<td>226</td>
<td>&lt;0.0001b</td>
<td>0.050</td>
</tr>
<tr>
<td>Geisha girl</td>
<td>-7.46</td>
<td>226</td>
<td>&lt;0.0001b</td>
<td>0.040</td>
</tr>
<tr>
<td>Sheena’s gold</td>
<td>-10.89</td>
<td>226</td>
<td>&lt;0.0001c</td>
<td>0.009</td>
</tr>
<tr>
<td>Myoporum</td>
<td>-10.11</td>
<td>226</td>
<td>&lt;0.0001c</td>
<td>0.002</td>
</tr>
<tr>
<td>Jacaranda</td>
<td>-12.26</td>
<td>226</td>
<td>&lt;0.0001c</td>
<td>0.002</td>
</tr>
<tr>
<td>Sine (time)</td>
<td>3.46</td>
<td>226</td>
<td>0.0006</td>
<td>NA</td>
</tr>
</tbody>
</table>

Table 1

Linear mixed effects ANOVA, with site modelled as a random factor, on the proportion of branches with *Aconophora compressa* in any developmental stage on different host plants. Different letters next to *p*-values indicate significant differences across host taxa.
have occurred incidentally. All branches from all plant taxa sampled were less than 2.5 m from the ground.

Plants were sampled monthly in a repeated measures design between August 2006 and November 2007 (except Bracken Ridge, which was not sampled in August 2006, and no sites were sampled in October 2007). All plants at each site were within a 2 km radius of the site centre (Fig. 1) and were most often within a 1 km radius of co-ordinates stated above. All sites, except Mt Tamborine, were chosen because sufficient plants of each species were present. Mt Tamborine was chosen because of its relatively high altitude and cooler climate and these conditions may contribute to relatively high survival by A. compressa during high summer temperatures (Dhileepan et al., 2005). Only three plants of myoporum were found at Mt Tamborine, and then only in February 2007.

Most plants were in suburban gardens and subject to periodic pruning. Plants that were heavily pruned during the study were omitted until sufficient regrowth was available. A few plants that were removed by gardeners and, for sampling purposes, were re-placed with another nearby plant that was sampled monthly thereafter. Where individual plants could not be distinguished, e.g. in thickets of lantana and myoporum, a 1 m section of the entire growth was marked and monitored consistently. The leaf rachis of jacaranda was treated as a “branch” for the purposes of the survey and analysis because A. compressa uses these parts almost exclusively when on jacaranda.

All analyses were conducted in R version 2.6.2. Survey methods were designed with the intent of conducting repeated measures statistical analysis with random effects for site and individual plant. More than 90% of the data were, however, represented by zeroes and consequently broke the assumptions of all such analyses attempted (i.e. a variety of models using lme, glmmPQL, gamlss and gam in R), so the data were analysed as described in the following subsections.

Climatic data were collected for each site from the Australian Bureau of Meteorology (http://www.longpaddock.qld.gov.au/silo/datadrill/index.frames.html). Temperatures above 30 °C cause sig-
significant levels of mortality for *A. compressa* (Dhileepan et al., 2005). Therefore, for each month and site, the maximum temperature, the total number of days with 30 °C or above, the longest period with consecutive days with 30 °C or above and the mean rainfall were retrieved. Means and standard errors were calculated for each variable across sites.

2.1.2. Intensity

To examine differences in the intensity of host use across the plant taxa, the proportion of branches with *A. compressa* (any number and any developmental stage) was calculated for each month, site and taxon and arc-sine square root transformed. A linear mixed effects ANOVA was used to determine differences in host use across the plant taxa. The transformed proportion of branches with *A. compressa* was modelled by host taxon and the sine of time (to account for oscillating abundance of the insects over time) and site was modelled as a random factor. A power variance structure, representing a variance covariate given by the fitted values (the default setting of the varPower function), was also modelled. A generalised linear hypothesis test based on a Tukey test was used to determine significant differences across host taxa.

The mean number of each stage of *A. compressa* (unsexed adults, females on egg batches, early nymphs and late nymphs) per branch was plotted for each host plant per month to illustrate relative field host use across the plant taxa sampled. The pattern of infestation that was typical for individual plants was obscured when data were averaged across those individuals. Therefore the mean number of each life-stage of *A. compressa* was also plotted for each individual plant, of a selected subset of plants over time for fiddlewood, lantana and geisha girl to highlight typical temporal patterns of host use.

The proportion of plants with *A. compressa* was plotted for each plant taxon each month. In addition, the proportion of adult and nymphal *A. compressa* on each host plant taxon was plotted relative to fiddlewood for each month sampled.

2.1.3. Regularity

The presence of each stage of *A. compressa* on individual plants of each taxon was plotted to measure the regularity of host use over time and to calculate various summary statistics. Individual plants sampled for at least eight months (about half the entire sampling period) were included in the following analysis. The percentages of plants upon which *A. compressa* were present and absent in every month sampled were calculated separately. The percentage of months in which *A. compressa* was present on each plant and the longest continuous sequence of months through which *A. compressa* was present and absent were also calculated separately. These statistics were presented as a percentage of the total number of months sampled for each individual plant, to account for differences in the number of months sampled across individuals. Individual plants were scored positively every month that at least one *A. compressa* individual was present.

2.2. Survey in Mexico

A 3 week field survey was conducted in Mexico (within the native range of *A. compressa*) between January 23 and February 14, 2007. Plants were sampled as per the Australian surveys, but the number of plants sampled in any given area was not limited to 15 plants (as in Australian surveys) because repeated measures were not possible. The field survey was concentrated in the state of Morelos (Fig. 1) as this is considered an area of relatively high *A. compressa* abundance (Palmer et al., 1996). Plants were also searched in the following states: Mexico, Puebla, Tlaxcala and Guerrero (Fig. 1). Surveys were conducted along major and minor roadways and were more concentrated in areas with relatively high soil moisture, as indicated by lush, green foliage. *Lantana urticifolia, Citharexylum* spp. (probably two species that were 2–3 m shrubs, unlike *C. spinosum* in Australia which is a tall tree), *jacaranda, T. stans* and other verbenaceous plants were actively sought, although plants in moderate abundance in any given area were searched for *A. compressa*. The locations of plants of various *Citharexylum* spp. were sought with reference to herbarium specimens from the National Herbarium of Mexico, Mexico City. Insect specimens from all host species were identified by Chris Dietrich of the Illinois Natural History Survey. The data were split into two groups based on geographic location: Morelos and Puebla. Since there were no repeated measures and distinct sites were not identified (and which acted as replicates in the Australian analysis), formal statistical analysis was not conducted. Non-overlapping 95% confidence intervals were therefore used to indicate differences in abundance across host taxa.

3. Results

3.1. Surveys in Australia

3.1.1. Intensity and regularity

Three distinct and significantly different groups of plant species could be circumscribed by the numbers of *A. compressa* they hosted in the field. Fiddlewood hosted significantly more insects per branch than any other host (Fig. 2), had consistently more branches with *A. compressa* (Table 1) and alone made up the first group. The second group consisted of lantana and geisha girl and the third Sheena’s gold, myoporum and jacaranda (Fig. 2; Table 1). The proportion of individual plants of each taxon with *A. compressa* (Fig. 3), and the proportion of all *A. compressa* adults and nymphs collected

![Graph](image-url)
at each sampling time on each host species, relative to the proportion on fiddlewood plants (Fig. 4), followed this same trend.

On average, *A. compressa* was present continuously on fiddlewood, lantana and geisha girl (for all but 2 months on the last plant species) (Figs 2 and 4). When individual plants were inspected for continuity of *A. compressa* presence, however, it was only fiddlewood that hosted them persistently (Table 2), as indicated by the data from selected individual plants in Fig. 5. On lantana and geisha girl, by contrast, individual plants were never used continuously (Fig. 5; Table 2). The longest continuous stretch during which *A. compressa* was recorded on lantana and geisha girl was, on average, about 2–3 months (Table 2), much shorter than that observed on fiddlewood, about 7 months (Table 2). Fiddlewood was the only plant taxon for which *A. compressa* was found on every plant over the sample period (Table 2) and, on average, individual trees harboured *A. compressa* about 10 of the 15 months sampled, i.e. 66.6% (Table 2). *A. compressa* was not found on any plant, besides fiddlewood, for the entire sample period and was never found on about 13% of lantana plants sampled (Table 2). In summary, the data indicate that fiddlewood was used much more regularly (Table 2) and at much higher intensities (Figs 2–5) than any

<table>
<thead>
<tr>
<th>Treatment (n)</th>
<th>Plants that had <em>A. compressa</em> for the entire sampling period (%)</th>
<th>Plants that had no <em>A. compressa</em> for entire sampling period (%)</th>
<th>Mean longest continuous time period without <em>A. compressa</em></th>
<th>Mean number of months that plants had <em>A. compressa</em></th>
<th>Mean longest continuous time period with <em>A. compressa</em></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Plants</td>
<td>Mean length</td>
<td>Plants</td>
<td>Mean length</td>
<td>Plants</td>
</tr>
<tr>
<td></td>
<td>overall</td>
<td>continuous</td>
<td>overall</td>
<td>continuous</td>
<td>overall</td>
</tr>
<tr>
<td>Fiddlewood (74)</td>
<td>8.1</td>
<td>28.9 ± 2.2</td>
<td>0.0</td>
<td>4.3 ± 0.3</td>
<td>66.6 ± 2.3</td>
</tr>
<tr>
<td>Lantana (85)</td>
<td>0.0</td>
<td>62.2 ± 2.6</td>
<td>12.9</td>
<td>9.3 ± 0.4</td>
<td>26.0 ± 1.9</td>
</tr>
<tr>
<td>Geisha girl (78)</td>
<td>0.0</td>
<td>72.5 ± 2.5</td>
<td>18.0</td>
<td>10.9 ± 0.4</td>
<td>20.9 ± 1.9</td>
</tr>
<tr>
<td>Sheena’s gold (72)</td>
<td>0.0</td>
<td>90.7 ± 1.5</td>
<td>54.2</td>
<td>13.6 ± 0.2</td>
<td>7.2 ± 1.2</td>
</tr>
<tr>
<td>Myoporum (65)</td>
<td>0.0</td>
<td>96.4 ± 1.3</td>
<td>80.0</td>
<td>14.5 ± 0.2</td>
<td>2.4 ± 0.7</td>
</tr>
<tr>
<td>Jacaranda (73)</td>
<td>0.0</td>
<td>97.3 ± 1.0</td>
<td>82.2</td>
<td>14.6 ± 0.2</td>
<td>1.8 ± 0.5</td>
</tr>
</tbody>
</table>

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**Fig. 5.** Bar plots illustrating that *Aconophora compressa* was present continuously (August 2006–November 2007) on some fiddlewood individuals, and that on lantana and geisha girl it was not. Segments of the x-axis without bars above them indicate that no *A. compressa* individuals were on that individual plant in that month. The plant represented by Fiddlewood 2 (top right hand plot) was not sampled in August or September 2006.
other host species. Regularity and intensity of host use were lower on lantana, followed by geisha girl, Sheena’s gold, jacaranda and myoporum.

Protracted periods of high summer temperatures (≥ 30 °C) were recorded between November 2006 and March 2007 (Table 3). Numbers of A. compressa dropped considerably during this time period across all host plants (Fig. 2). The numbers of A. compressa

Table 3
Monthly climate data over the survey period. Means (±S.E.) were calculated across sites for each month. No survey was conducted in October 2007.

<table>
<thead>
<tr>
<th>Month (survey month)</th>
<th>Maximum temperature (°C)</th>
<th>Total number of days &gt;30 °C</th>
<th>Longest number of consecutive days &gt;30 °C</th>
<th>Rainfall (mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>August 2006 (1)</td>
<td>28.3 ± 0.85</td>
<td>0.0 ± 0.0</td>
<td>0.0 ± 0.0</td>
<td>45.6 ± 3.52</td>
</tr>
<tr>
<td>September 2006 (2)</td>
<td>28.3 ± 0.75</td>
<td>0.0 ± 0.0</td>
<td>0.0 ± 0.0</td>
<td>63.2 ± 4.97</td>
</tr>
<tr>
<td>October 2006 (3)</td>
<td>32.9 ± 1.30</td>
<td>1.2 ± 0.37</td>
<td>0.8 ± 0.2</td>
<td>204.2 ± 2.25</td>
</tr>
<tr>
<td>November 2006 (4)</td>
<td>34.0 ± 0.98</td>
<td>6.4 ± 1.33</td>
<td>3.4 ± 0.6</td>
<td>63.3 ± 2.33</td>
</tr>
<tr>
<td>December 2006 (5)</td>
<td>31.4 ± 0.75</td>
<td>3.4 ± 0.93</td>
<td>1.4 ± 0.40</td>
<td>75.8 ± 6.48</td>
</tr>
<tr>
<td>January 2007 (6)</td>
<td>34.0 ± 0.76</td>
<td>16.8 ± 2.67</td>
<td>10.0 ± 1.76</td>
<td>70.4 ± 2.71</td>
</tr>
<tr>
<td>February 2007 (7)</td>
<td>30.8 ± 0.75</td>
<td>10.0 ± 3.10</td>
<td>5.2 ± 2.60</td>
<td>74.7 ± 10.54</td>
</tr>
<tr>
<td>March 2007 (8)</td>
<td>35.6 ± 1.04</td>
<td>15.8 ± 3.87</td>
<td>6.4 ± 1.44</td>
<td>42.1 ± 13.07</td>
</tr>
<tr>
<td>April 2007 (9)</td>
<td>28.7 ± 0.60</td>
<td>0.2 ± 0.20</td>
<td>0.2 ± 0.20</td>
<td>8.7 ± 2.43</td>
</tr>
<tr>
<td>May 2007 (10)</td>
<td>29.8 ± 0.75</td>
<td>1.2 ± 0.49</td>
<td>1.2 ± 0.49</td>
<td>36.3 ± 4.10</td>
</tr>
<tr>
<td>June 2007 (11)</td>
<td>25.3 ± 0.85</td>
<td>0.0 ± 0.0</td>
<td>0.0 ± 0.0</td>
<td>109.9 ± 2.38</td>
</tr>
<tr>
<td>July 2007 (12)</td>
<td>25.6 ± 0.78</td>
<td>0.0 ± 0.0</td>
<td>0.0 ± 0.0</td>
<td>1.5 ± 0.50</td>
</tr>
<tr>
<td>August 2007 (13)</td>
<td>27.4 ± 0.86</td>
<td>0.0 ± 0.0</td>
<td>0.0 ± 0.0</td>
<td>103.4 ± 5.35</td>
</tr>
<tr>
<td>September 2007 (14)</td>
<td>30.8 ± 0.97</td>
<td>4.2 ± 1.11</td>
<td>2.2 ± 0.58</td>
<td>36.2 ± 4.24</td>
</tr>
<tr>
<td>October 2007</td>
<td>32.9 ± 0.93</td>
<td>7.2 ± 2.27</td>
<td>2.2 ± 0.8</td>
<td>90.9 ± 9.00</td>
</tr>
<tr>
<td>November 2007 (15)</td>
<td>31.1 ± 0.80</td>
<td>1.8 ± 0.58</td>
<td>1.8 ± 0.58</td>
<td>80.1 ± 3.12</td>
</tr>
</tbody>
</table>

3.2. Survey in Mexico

Overall, about 450 plants were sampled from about 15 host taxa (Table 4). The abundance of A. compressa in Mexico was relatively low, on all host plants, compared to its abundance in Australia. A. compressa was more abundant in Morelos than any other area surveyed (Fig. 6; Table 4). In Morelos it was found at similar relatively high rates on D. erecta, L. urticifolia and T. stans (Fig. 6a). Abundance was very low on lemon verbena (Aloysia citriodora (Cav.) Ort.: Verbenaceae) and jacaranda (Fig. 6a). Citharexylum spp. were not found in Morelos despite extensive search.

In Puebla, A. compressa was found on three plants, one each of C. cinereum L., L. urticifolia and Senecio spp. (colloquial name ‘jarilla’, which possibly represents a number of plant species across the different regions surveyed) (Fig. 6b). A single male A. compressa was found on Wigandia spp. (Hydrophyllaceae) in the state of Mexico but was not found on this plant anywhere else despite sampling this species in areas in which A. compressa was in relatively high abundance (Table 4). Otherwise, A. compressa was not found outside Morelos and Puebla (Table 4).

4. Discussion

Early determination of multiple host use by herbivore species can be made and hosts can be categorised in an ecologically meaningful way from quantitative field surveys of multiple host plant species over time. The categorisations derived here correlate well with those derived from nymphal rearing experiments across the host plants sampled in Australia (Manners and Walter, 2009) and also from adult performance tests on these plants (Manners et al., 2010). The results presented here also demonstrate that the host plant associations of A. compressa are consistent across geographic regions sampled, in terms of which plant species are used as primary, secondary or incidental hosts (Figs. 2–5, Tables 1 and 2). Further sampling is needed in the native range, but the results have clear implications for biocontrol practice, as explained below.

4.1. A. compressa in Australia

The abundance of A. compressa on fiddlewood was always higher than on any other host sampled, in terms of the mean number of A. compressa per branch (Fig. 2) and proportion of trees with A. compressa increased on fiddlewood, lantana and geisha girl from April 2007: numbers on Sheena’s gold, jacaranda and myoporum remained very low (Fig. 2).
compressa (Fig. 3). The relative proportion of A. compressa adults and nymphs, compared to fiddlewood, on each host species was never more than 15% and 30%, respectively, of that on fiddlewood in any given month (Fig. 4), but was usually much lower. Results indicate that lantana, geisha girl, Sheena’s gold, myoporum and jacaranda were used mostly as a “spill-over” from fiddlewood (Figs 2–4). Only fiddlewood trees harboured A. compressa continuously (Table 2, Fig. 5) and recolonisation of secondary host plants occurred repeatedly over the course of the surveys (Figs. 2 and 5). Furthermore, the use of secondary and incidental host plants appears ephemeral, perhaps as a result of adults that developed from eggs laid on the secondary and incidental hosts emigrating soon after eclosion. However, this possibility would require a separate test.

In general, the results presented here provide further support that fiddlewood is the primary host plant of A. compressa in Australia and that all other hosts, including lantana, are secondary or incidental hosts (Manners and Walter, 2009; Manners et al., 2010). A. compressa has been recorded on particular lantana plants for periods of about 26 weeks (double that recorded in this study – Table 2) and at moderately high levels (about three to five times that recorded in this study) in 2009 and 2010, at the Alan Fletcher Research Station, Sherwood, Brisbane, Australia. Such populations on lantana have been recorded only as relatively isolated patches of these insects in New South Wales and Queensland (W.A. Palmer, unpublished data). The relative abundance of A. compressa on fiddlewood and other host species near these relatively large populations of A. compressa on lantana is not known.

The statistical analysis grouped Sheena’s gold with jacaranda and myoporum, rather than with lantana and geisha girl, even though Sheena’s gold is used more regularly and at somewhat higher abundance than jacaranda and myoporum (Figs. 2–4; Tables 2). Sheena’s gold may thus be a relatively low ranking secondary host.

4.2. A. compressa in Mexico

The overall abundances of A. compressa on host plants in Mexico were much lower than those recorded in Australia (Fig. 6), but patterns of host use were similar across the two regions (Figs. 2–4). In the state of Morelos, A. compressa was found on a number of plant species, mainly from the family Verbenaceae, but also some from relatively distantly related families (Table 4). A. compressa was found on D. erecta and T. stans at levels similar to lantana (Table 4; Fig. 6). Counts of A. compressa on Lantana spp. in Mexico between 1989 and 1993 indicate that numbers are sometimes much higher (e.g. 40–100 adults per plant) than results presented here (W.A. Palmer, unpublished data). The results support the notion that host plant surveys in the native range should not focus exclusively on the target plant species (Walter, 2003; Sheppard et al., 2006a). The extent to which non-target plant species are surveyed remains problematic. It may be possible to formalise a native range ‘search list’ similar to those plants included for host testing. In any case, effort should be made to survey all possible plant species from the same family and of similar growth habit to the suspected primary host species in regions where the herbivore is relatively abundant.

Host plant use by A. compressa in Morelos also provides evidence that a number of the plant species used in Australia would have been anticipated if non-target plants had been surveyed prior to its release. These include D. erecta, jacaranda, A. citriodora and T. stans (quantitative surveys have not been conducted on the latter two species in Australia, although Palmer et al. (2004) documented the presence of A. compressa on these hosts). The original host testing indicated that D. erecta was suitable for development by A.
compressa (Palmer et al. 1996) and had been found on D. erecta in Mexico prior to its release in Australia (Palmer and Pullen, 1995; Palmer et al., 1996).

This field survey of A. compressa in Morelos indicates that L. urticifolia, D. erecta and T. stans are in the same category of hosts (i.e. either primary or secondary) and jacaranda and A. citriodora are in a lesser category (i.e. either secondary or incidental hosts). More information is required to determine whether L. urticifolia, D. erecta and T. stans are primary or secondary hosts in Mexico. Essentially, data over multiple seasons on each host plant are required to understand this aspect of A. compressa ecology in Mexico.

The relationship between A. compressa and Citharexylum spp. in Mexico remains somewhat ambiguous, perhaps influenced by a number of factors. Many Citharexylum spp. are present in Mexico, but C. spinosum has not been recorded (Moldenke, 1942). C. spinosum is native to the Caribbean and Florida but A. compressa is not known from these regions (Dietrich and Deitz, 1991). Fiddlewood plants in Mexico were, however, rare or had low appearance during the survey. Since Citharexylum spp. are deciduous and January to February is dry in Mexico, these plants possibly harbour A. compressa at other times. Long term field surveys, perhaps including adjacent geographic regions in the range of A. compressa (e.g. Guatemala), would help clarify the status of the relationship between A. compressa and fiddlewood in its native range. This would also aid in the determination of whether lantana and fiddlewood are primary or secondary host plants in Mexico.

Lantana in Australia was thought to be most closely related to lantana from Mexico (Scott 2002), although recent evidence suggests that it may be more closely related to lantana from Venezuela and the Dominican Republic (R. Watts, CSIRO, Plant Industries, Canberra, personal communication). In general, the points made above indicate just how crucial are full field surveys across all seasons in the native range of a weed to make an informed release decision. That is, long-term field surveys are more likely to provide an accurate indication of the host relationships of the herbivore with its host plants than will sole reliance on laboratory or greenhouse based tests in quarantine. The latter, however, are important to interpret the field results and will provide the sole information on plants not present in the native range.

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