

A meta-analytic investigation of the potential for plant volatiles and sex pheromones to enhance detection and management of Lepidopteran pests

Article

Accepted Version

Staton, T. ORCID: <https://orcid.org/0000-0003-0597-0121> and Williams, D. T. (2023) A meta-analytic investigation of the potential for plant volatiles and sex pheromones to enhance detection and management of Lepidopteran pests. *Bulletin of Entomological Research*. ISSN 0007-4853 doi: <https://doi.org/10.1017/S0007485323000457> Available at <https://centaur.reading.ac.uk/113733/>

It is advisable to refer to the publisher's version if you intend to cite from the work. See [Guidance on citing](#).

To link to this article DOI: <http://dx.doi.org/10.1017/S0007485323000457>

Publisher: Cambridge University Press

All outputs in CentAUR are protected by Intellectual Property Rights law, including copyright law. Copyright and IPR is retained by the creators or other copyright holders. Terms and conditions for use of this material are defined in the [End User Agreement](#).

www.reading.ac.uk/centaur

CentAUR

Central Archive at the University of Reading

Reading's research outputs online

1 **Title Page**

2 **Title:**

3 A meta-analytic investigation of the potential for plant volatiles and sex pheromones to
4 enhance detection and management of Lepidopteran pests

5

6 **Authors:** Tom Staton^{1*}, David T. Williams¹

7 ¹ Forest Research, Alice Holt Lodge, Farnham, GU10 4LH, UK

8 * Correspondence author: tom.staton@reading.ac.uk; current address: University of
9 Reading, School of Agriculture Policy & Development, Earley Gate, Reading, RG6 7BE, UK

10

11 Revision 1 (submitted 05 July 2023)

12 **Abstract**

13 Effective early detection, monitoring and management methods are critical for reducing the
14 impacts of insect pests in agriculture and forestry. Combining host plant volatiles with sex
15 pheromones could enhance trapping methodologies, whilst the use of non-host volatiles could
16 improve the effectiveness of pest management through repellency effects. In this
17 meta-analysis approach, we analysed 51 studies that used electroantennograms (EAG), wind
18 tunnels, and/or field traps, to evaluate the antennal and behavioural responses of Lepidoptera
19 to sex pheromones combined with attractant or repellent plant volatiles. Proposed attractant
20 plant volatiles had a positive association with female Lepidoptera responses to sex
21 pheromone, but effects on males were highly variable, with unexpected repellency reported in
22 some studies. Repellent plant volatiles were significantly or near-significantly negatively
23 associated with male attraction to sex pheromones but were scarcely studied. Sub-group
24 analysis identified that male responses to sex pheromone were reduced when the dose of
25 attractant plant volatile relative to sex pheromone was increased. Green-leaf volatiles were
26 associated with the strongest positive effects for males in field traps. Multiple-compound
27 attractant plant volatile blends were less effective than single compounds in field studies. Our
28 analysis demonstrates, i) the potential value of combining host plant volatiles with sex
29 pheromones to capture females rather than only males, ii) the importance of identifying
30 appropriate host plant volatiles and optimal relative doses, and iii) the potential for non-host
31 plant volatile use in pest management strategies.

32

33 Running head: Plant volatiles and Lepidopteran pests

34 Keywords: electroantennogram, insects, kairomone, semiochemical, trapping, wind tunnel

35

36 Introduction

37 Insect pests have major economic and environmental impacts within both forestry and
38 agricultural systems worldwide (Culliney, 2014; van Lierop et al., 2015), which is predicted to
39 be exacerbated because of projected climate change scenarios and increased risk of the
40 establishment of non-native invasive species through global trade (Deutsch et al., 2018;
41 Lehmann et al., 2020). The identification of pheromones and plant volatile chemicals which
42 may attract or repel insect pests is a critical research area that could facilitate the development
43 of more effective early detection and monitoring tools and improved or alternative
44 management approaches (Larson et al., 2020; Mafra-Neto et al., 2022). In the Coleoptera for
45 example, successful applications of plant volatiles in insect pest management include the
46 combination of a host plant volatile with a sex pheromone to attract emerald ash borer *Agrilus*
47 *planipennis* Fairmaire to traps (Ryall et al., 2012; Wittman et al., 2021), and anti-aggregation
48 pheromones combined with non-host volatiles to repel mountain pine beetle *Dendroctonus*
49 *ponderosae* Hopkins from forestry crops (Fettig & Munson, 2020).

50 Lepidoptera (butterflies and moths) include many of the most serious pests of plants and have
51 been reported to make up eight of the top 20 most studied arthropod pests, which includes
52 diamondback moth *Plutella xylostella* L., cotton bollworm *Helicoverpa armigera* Hübner, and
53 codling moth *Cydia pomonella* L. (Willis, 2017). Plant volatile lures, typically derived from the
54 insect's favoured host plant species, have previously been shown to be effective in attracting
55 adult Lepidoptera, especially females (Szendrei & Rodriguez-Saona, 2010). Furthermore, the
56 identification of sex pheromones for many Lepidopteran pest species in recent years has led
57 to the development of effective species-specific traps, although these typically only catch
58 males (Witzgall et al., 2010; Rizvi et al., 2021). The combination of host plant volatile lures
59 with sex pheromones therefore has potential to improve capture rates of males whilst also
60 attracting females, on the premise that females are attracted to volatiles associated with
61 favoured host plants, and males are attracted to females located on optimal hosts (Reddy &
62 Guerrero, 2004; Szendrei & Rodriguez-Saona, 2010; Bruce & Pickett, 2011; Xu & Turlings,

63 2018). In addition, the identification of non-host plant volatiles which repel pests could lead to
64 more effective management options, such as 'push-pull' strategies which aim to 'push' pests
65 away from high priority areas using repellent plant volatiles or appropriate non-host plants,
66 whilst simultaneously using attractant volatiles to 'pull' the target pest into traps or to lower-
67 risk areas (Cook et al., 2007; Eigenbrode et al., 2016).

68 Early studies in the 1980s and 1990s found that combining host plant volatiles with sex
69 pheromone lures did improve capture rates of male moths compared with pheromone-only
70 lures (reviewed in Landolt & Phillips, 1997). A number of subsequent studies, however, have
71 found that proposed attractant plant volatiles unexpectedly reduce male capture rates (e.g.
72 Meagher, 2001; Hu et al., 2013; Barros-Parada et al., 2018) or fail to catch females (Tang et
73 al., 2012). Several possible explanations have been suggested to explain the variable
74 effectiveness of candidate attractant plant volatiles. The choice of host plant volatile used
75 within combined lures is an important factor which depends on target species (e.g. Tang et
76 al., 2012; Hu et al., 2013; Mujica et al., 2018), and in some cases effects have been classified
77 according to the type of compound such as green leaf volatiles, floral aromatics or terpenes
78 (Fang et al., 2018). In addition, the dosage of host plant volatiles and their ratio to the sex
79 pheromone has been shown to be an important factor in some cases, with several studies
80 demonstrating that that the dosage of plant volatile can determine whether there is an
81 attractant or repellency effect of a certain plant volatile on male response of a target species
82 to its sex pheromone (e.g. Hu et al., 2013; Yu et al., 2015; Xiang et al., 2019). In the field, trap
83 design can also influence the effectiveness of combined lures (Gregg et al., 2018), because
84 trapping area (i.e. the trap's behaviourally effective plume reach) could depend on trap type,
85 and a plant volatile which is attractive at distance could act as a repellent at close proximity to
86 the insect (Kvedaras et al., 2007). Other factors which might conceivably influence the effects
87 of plant volatiles combined with sex pheromones include; (i) host specialism of the target
88 species, since host-specialists might be more attracted to or repelled by particular volatiles;
89 (ii) blend complexity, given that blends of host volatiles are often more attractive than a single

90 component (Gregg et al., 2018); and (iii) habitat, given that background odour influences the
91 interaction between plant volatiles and pheromones (Cai et al., 2017). However, these factors
92 have yet to be systematically investigated in the context of combining plant volatiles with sex
93 pheromone lures.

94 The current lack of systematic evidence to explain why host plant volatiles can unexpectedly
95 reduce the attraction of male Lepidoptera to sex pheromones and/or fail to attract females,
96 and whether non-host volatiles can repel male Lepidoptera from sex pheromones, is a
97 constraint to ongoing research of pest management strategies for some Lepidoptera species.
98 As such, the aim of this study was to evaluate the effect of proposed attractant or repellent
99 plant volatiles on the antennal and behavioural responses of adult Lepidoptera to sex
100 pheromone in laboratory and field studies. In addition, we considered which factors might be
101 most important in designing effective combined pheromone-plant volatile lures, by evaluating
102 the influence of plant volatile category, plant volatile blend complexity, ratio of the plant volatile
103 to sex pheromone, host specialism of the target insect, habitat type, and trap type, on the
104 attraction of male Lepidoptera to plant volatiles. We also investigated spatial patterns of
105 attraction of males and females to combined pheromone-plant volatile lures in wind tunnels.
106 We then discuss how our findings relate to previous studies which have investigated the
107 effects of plant volatiles alone (e.g., Szendrei & Rodriguez-Saona, 2010).

108

109 **Materials and Methods**

110 *Literature search*

111 A search of the literature was undertaken using Scopus in early 2022. The following search
112 term was used within article title, abstract and keywords: pheromone AND (volatile OR
113 kairomone OR allomone OR semiochemical) AND (“wind tunnel” OR olfactometer OR EAG
114 OR electroantennogram OR trap*). Studies were initially screened by reviewing their titles,
115 abstracts, and main text as appropriate, and included in the meta-analysis if they met all of

116 the following criteria: (i) results presented for a species of Lepidoptera; (ii) the study compared
117 the effects of plant volatile(s) combined with a conspecific sex pheromone simultaneously at
118 the same source, versus a sex pheromone-only control, on the response of the target species.
119 The two treatments otherwise comprised exactly the same specifications (e.g. trap type) and
120 the same sex pheromone constituents; (iii) for field trapping, sites which were not under mating
121 disruption or sterile moth release treatments; (iv) sufficient text in English to decipher the
122 results, and (v) adequate and clearly presented statistical information (e.g. means, standard
123 errors, and sample sizes) for inclusion in the meta-analysis models. Studies which failed the
124 final criterion were included in the Discussion. A non-systematic supplemental search of
125 Google Scholar was also undertaken to identify any additional references including grey
126 literature. Upon completion of the systematic search, references cited in the most recent
127 publications were screened to obtain any additional relevant studies.

128 Volatiles which were tested as both attractants (e.g. host volatiles) and repellents (e.g. non-
129 host volatiles) were included but were evaluated separately. Both male and female adult
130 Lepidoptera were included. The literature search was global with no geographic restrictions.
131 We aimed to include behavioural laboratory studies such as electroantennogram (EAG)
132 responses, choice experiments such as olfactometers, and wind tunnels, in addition to field
133 trapping studies. However, only three choice experiment publications were identified (Xiao et
134 al., 2002; Ma et al., 2016; McCormick et al., 2017), therefore, this method was not included in
135 the meta-analysis.

136 A total of 1699 publications were returned by the search term. The majority of these did not
137 report data for Lepidoptera. Therefore, after screening according to the above criteria, 51
138 studies were selected for inclusion in the meta-analysis (listed in Supplementary Material 1),
139 comprising 9 EAG studies, 23 wind tunnel studies, and 35 field-trapping studies (note that
140 some of the 51 studies included more than one method).

141 *Calculation of effect sizes*

142 Data was extracted from each publication and compiled into separate databases for EAG,
143 wind tunnel, and field-trapping studies. Where necessary, data was extracted from figures
144 using WebPlotDigitizer Version 4.5 (Rohatgi, 2021). The response of Lepidoptera to combined
145 plant volatile-sex pheromone lures versus pheromone-only lures was measured as follows
146 according to each experimental method: for EAG studies, response was measured as
147 electrical antennal signal (which could equally represent an attractant or repellent response);
148 for wind tunnel studies, response was measured as the number of individuals achieving a
149 certain stage in the tunnel (e.g. take-off, half-way, approach to lure; recorded in separate
150 rows); and for field trapping studies, the response was defined as the number of individuals
151 captured in traps. Sex of the target insect was recorded, where stated. Where no female
152 captures were reported, the entire paper was reviewed for any reference to female captures;
153 if the paper stated that no females were captured, this was added into the effect size database
154 for each experiment.

155 All analysis was undertaken using the 'metafor' package (Viechtbauer, 2010) in R version
156 4.0.2 (R Core Team, 2020). Effect sizes were calculated using standardized mean difference
157 (Hedges, 1981) for EAG and field studies, or log odds ratio for wind tunnel studies because of
158 the proportion data. The 'treatment' was insect response to the combined plant volatile and
159 pheromone lure, and the 'control' was response to the pheromone-only lure.

160 *Meta-analysis models*

161 Meta-analysis models were built using the 'rma.mv' function in the 'metafor' package. To
162 account for the non-independence of experiments and studies, a hierarchical random effects
163 structure was specified comprising each effect size nested within experiment number (where
164 specified), nested within study ID. In addition, for wind tunnel data, stage of the wind tunnel
165 (e.g. take-off, halfway, approach to lure) was nested within experiment number. Confidence

166 intervals and significance tests were calculated using the t -distribution, which is more
167 conservative than the Z-distribution with less risk of Type I error.

168 Initially, overall meta-analysis models were built to test the effect of attractant or repellent plant
169 volatiles on adult Lepidoptera responses to sex pheromone according to each of the three
170 methods (EAG, wind tunnel, and field-trapping). Subgroup analysis was then used to
171 investigate causes of inconsistent effects among studies (i.e. heterogeneity) for attractant
172 volatiles.

173 *Subgroup analysis*

174 Subgroup analysis was performed to identify any differences in responses to combined versus
175 pheromone-only lures according to sex, host specialism, trap type, habitat, complexity of the
176 volatile blend, ratio of plant volatile to pheromone, and type of volatile chemical. These
177 subgroups were selected based on their expected influence on effect sizes, either
178 hypothetically or from previous evidence (as explained in the Introduction). A separate model
179 was built for each subgroup and method (EAG, wind tunnel, field-trapping), because different
180 data subsets were needed for each subgroup (Table 1). In each model, the subgroup was
181 specified as a moderator in the model.

182 In each subgroup model, the intercept term was removed to provide effect sizes for each level.
183 Data was filtered to only include males (apart from the sex subgroup), because males and
184 females responded differently, and males were much better represented in the data than
185 females. The significance of levels within each subgroup was tested using the omnibus
186 moderator test in the metafor package for models with the intercept term included, to test the
187 null hypothesis that the subgroup has no influence on effect sizes.

188 In addition, sex-specific differences in responses to combined attractant lures versus
189 pheromone-only lures at varying stages in wind tunnels was analysed by including the
190 interaction between sex and wind tunnel stage as a moderator, with the intercept removed.

191 Subgroup data was extracted from each publication where provided. Specialism was classified
192 according to the main host range of the study species according to CABI's Invasive Species
193 Compendium (CABI, 2022), with other sources used where necessary (Supplementary
194 Material 2). Specialists were defined as having main host plants within one taxonomic family,
195 while generalists had main hosts within more than one taxonomic family. Plant volatile types
196 were classified into seven categories (Table 1) based on their chemical composition. Values
197 for ratio of plant volatile to sex pheromone were quantified by standardising the amount of
198 plant volatile per one equivalent unit of sex pheromone. This quantity varied enormously
199 among studies, therefore, the amount of plant volatile relative to sex pheromone was further
200 standardised within each study using the 'scale' function in R.

201 *Sex ratios*

202 Ratios of female to male Lepidoptera were calculated for each field trapping study which
203 reported data for both sexes. The effect of combined attractant lures versus pheromone-only
204 lures on sex ratio was tested using a mixed model in the 'lme4' package (Bates et al., 2015),
205 where the sex ratio was $\log(x+1)$ transformed to account for positive skew. Random effects
206 comprised experiment number nested within study ID. Marginal means were calculated using
207 the 'emmeans' package (Lenth, 2021) and back-transformed.

208 *Publication bias*

209 Publication bias was assessed by visual inspection of funnel plots, and via a multilevel
210 meta-regression model with the square-root of the inverse effective sample size as a
211 moderator. This model tests whether larger effect sizes are associated with larger standard
212 errors, which would indicate publication bias (Nakagawa et al., 2022). Evidence of publication
213 bias was found for the EAG models according to both funnel plots and the standard error
214 moderator models, although given that only nine studies were available for this method, the
215 apparent publication bias could be a chance effect. Therefore, the findings for the EAG models
216 are presented as preliminary results, which require further research.

217 *Influential cases*

218 The influence of individual effect sizes on the model outputs was tested by calculating
219 DFBETAS values in the 'metafor' package. In accordance with the guidance for this function,
220 DFBETAS values greater than 1 were considered to be influential effect sizes. This was
221 detected in one model, the sex subgroup model for EAG studies. Omitting the five influential
222 effect sizes from this model did not substantially alter the results; the moderator and each sex
223 was still significant and the contrast between the sexes was greater. Therefore, the full model
224 is presented.

225

226 **Results**

227 A total of 1561 effect sizes were extracted from 51 studies, comprising 126 from 9
228 electroantennogram (EAG) studies, 993 from 23 wind-tunnel studies, and 442 from 35 field-
229 trapping studies (note that some papers fell into more than one category). The majority of
230 these studies reported effects for plant volatiles tested as attractants, while only 3 wind tunnel
231 studies and 4 field trapping studies tested proposed repellents. The number of publications
232 rose sharply after 2010 (Figure 1). The publications reported data from a total of 27
233 Lepidopteran species, comprising 6 in EAG studies, 15 in wind tunnel studies, and 21 in field-
234 trapping studies. The most frequently studied species were oriental fruit moth *Grapholita*
235 *molesta* Busck (included in 11 of 57 studies) and codling moth *Cydia pomonella* L. (10 of 57
236 studies). The field-trapping data originated from 11 countries across Asia (14 studies), North
237 America (14 studies), Europe (4 studies), Australia/New Zealand (2 studies), and South
238 America (1 study).

239 Attractant plant volatiles had a significant positive association with Lepidoptera responses
240 (both sexes) to sex pheromones in field-trapping studies (0.210 ± 0.069 SE, $t = 3.028$, p -value
241 = 0.003) and in the more limited number of EAG studies (effect size 1.347 ± 0.395 SE, $t =$
242 3.413 , p -value < 0.001), but the effect was not significant in wind tunnel studies (0.241 ± 0.149

243 SE, $t = 1.619$, $p\text{-value} = 0.106$). Repellents had a significant negative association with male
244 Lepidoptera responses to sex pheromone lures in wind tunnels (effect size -1.120 ± 0.177 SE,
245 $t = -6.338$, $p\text{-value} < 0.001$), and a near-significant negative association with male capture
246 rates in pheromone traps in the field (-1.243 ± 0.646 SE, $t = -1.924$, $p\text{-value} = 0.062$). No data
247 on combining repellents with sex pheromones was available for EAG studies or for females.

248 There was significant heterogeneity in the effect of combined attractant lures (i.e. plant volatile
249 and pheromone) versus pheromone-only lures on adult Lepidoptera responses, for all three
250 methods (EAG: $Q = 609$ ($df = 125$), $p\text{-value} < 0.001$; wind tunnel: $Q = 3702$ ($df = 935$), $p\text{-value}$
251 < 0.001 ; field-trapping: $Q = 669$ ($df = 352$), $p\text{-value} < 0.001$), indicating that variation in effect
252 sizes was greater than expected based on sampling error alone. Subgroup analysis explained
253 some of this variability, demonstrated by the significant moderator tests for five of the seven
254 subgroups, according to at least one method (Figure 2). The five significant effects comprised;
255 (1) females responded more strongly than males to the combined lures compared with sex
256 pheromone lures alone, which was consistent across all three methods (EAG, wind tunnel,
257 and field trapping) but only significant for field trapping; (2) significant differences among plant
258 volatile categories on male responses to sex pheromone in wind tunnels, where fruit volatiles
259 were associated with the strongest attraction effects. In field traps, effect sizes were
260 significantly higher for green leaf volatiles, although plant volatile category was not significant
261 overall; (3) significantly stronger effects of male responses to combined host plant volatiles
262 with pheromone lures versus pheromone-only lures in agricultural rather than orchard
263 habitats; (4) significant effects of blend complexity on male responses to combined lures in
264 the field, with stronger attractant effects associated with single-component plant volatile
265 blends; and (5) significantly stronger associations of male responses to sex pheromone when
266 combined with lower doses of plant volatiles across all three methods (EAG, wind tunnel, and
267 field trapping). Trap type and host specialism were not significant sub-groups for any method,
268 although consistently stronger associations of male responses to combined lures were
269 observed in specialist rather than generalist species across methods.

270 Captures in field traps were dominated by males in both the pheromone-only and combined
271 lures, although the latter captured significantly higher proportions of females ($t = 3.767$ ($df =$
272 205.7), p -value < 0.001). Estimated marginal mean sex ratios were 0.051 females per male in
273 pheromone-only traps, increasing to 0.100 females per male in the combined lure traps.

274 A comparison of sex-specific responses within wind tunnels revealed a strong differentiation
275 in female responses to combined plant volatile and sex pheromone versus pheromone-only
276 sources at the activation and take-off stages (Figure 3). By contrast, significant differences in
277 male responses to combined versus pheromone-only lures were only seen at the final
278 approach stage (Figure 3).

279

280 **Discussion**

281 The indications from our meta-analysis study highlighted that proposed attractant plant
282 volatiles had a significant positive association with adult Lepidoptera responses to sex
283 pheromones in EAG and field-trapping studies. Similarly, proposed repellents, such as non-
284 host volatiles, had a significant negative association with male responses to sex pheromones
285 in wind-tunnel trials, and a near-significant negative association in field trapping studies,
286 although only 3 and 4 studies were available, respectively.

287 A number of sub-group factors significantly influenced the effect of combined plant volatile-
288 sex pheromone versus pheromone-only lures across EAG, wind tunnel and field-trapping
289 studies (Figure 2): (i) responses of females to the combined versus pheromone-only lures
290 were consistently stronger than males (but only significantly so in field-trapping studies), (ii)
291 lower ratios of plant volatile to sex pheromone were associated with a stronger response to
292 combined lures in males, and (iii) there was an indication that host-specialist male Lepidoptera
293 responded more strongly to the combined lures than generalists. The properties of the plant
294 volatile, in terms of blend complexity and chemical category, were significant factors for certain
295 methods. Attractant volatiles had a stronger effect on male responses to sex pheromones in

296 agricultural than in orchard habitats. Analysis of wind tunnel studies revealed that the
297 combination of plant volatiles with sex pheromones increased female attraction at an early
298 stage, but aided male attraction at the final approach stage (Figure 3).

299 *Interactions between attractant plant volatiles and sex pheromones*

300 A previous meta-analysis reported that attractant plant volatiles alone have a significant effect
301 on insect herbivore captures, particularly in Lepidoptera, with stronger effects on females than
302 males (Szendrei & Rodriguez-Saona, 2010). This is supported by a more recent study
303 included in our meta-analysis (Judd et al., 2017a), albeit other studies reported similar
304 responses of both sexes to attractant plant volatiles alone (Li et al., 2018; Kong et al., 2020).
305 In contrast, sex pheromone lures are designed to be highly effective for males, but are typically
306 ineffective for females. Our meta-analysis indicates that combined sex pheromone-plant
307 volatile lures are more effective than pheromone-only lures at attracting both sexes, although
308 captures continue to be dominated by males due to the potency of sex pheromones. However,
309 our findings support those of individual studies which reported highly variable success in
310 combining proposed attractant plant volatiles with sex pheromones to attract males (e.g.
311 Meagher, 2001; Hu et al., 2013; Barros-Parada et al., 2018).

312 *Attractant vs repellent plant volatiles*

313 Repellent plant volatiles were far less studied than attractants, with only 4 field-trapping
314 studies included in our meta-analysis, compared with 27 studies on proposed attractants.
315 Despite the limited research in this area, plant volatiles tested as repellents relatively
316 consistently deterred males from sex pheromones, particularly in wind tunnel studies, in
317 contrast to proposed attractants which had highly variable outcomes. A similar pattern has
318 been reported for studies of the effects of plant volatiles alone on herbivorous insects
319 (Szendrei & Rodriguez-Saona, 2010). Non-host plant volatiles can interfere with male
320 attraction to females, and suppress female egg-laying (McNair et al., 2000; Jactel et al., 2011),
321 demonstrating their potential application in push-pull strategies (McNair et al., 2000; Cook et

322 al., 2007) or mating disruption (Wang et al., 2016). However, more field studies are needed,
323 given the lack of overall significant effect (p -value = 0.062) in our meta-analysis.

324 *Sex-specific differences*

325 Females consistently responded more strongly than males to combined lures compared with
326 sex pheromone-only lures, especially in field studies. This is perhaps unsurprising given the
327 low effectiveness of sex pheromone lures on female attraction; the addition of plant volatiles
328 to sex pheromone traps increased the proportion of females from ca. 5% to 10% relative to
329 male captures. As such, a key advantage of combined plant volatile and sex pheromone lures
330 is the potential to attract both female and male Lepidoptera, albeit males typically remain
331 dominant. This could have applications in mass trapping or monitoring programmes where
332 detecting females is advantageous. However, the number of field studies reporting data for
333 female captures was limited (16, compared to 27 for males), and only 4 of these 16 specified
334 that no females were captured. As such, the significant positive effect of combined attractant
335 lures on female capture rates could be affected by reporting bias. Where combined lures do
336 not capture females, this could be explained by the limited mobility of females in some species
337 (Li et al., 2012a; Miluch et al., 2014) and the repellent effect of conspecific sex pheromone
338 (Barnes et al., 1992; Weissling & Knight, 1996; Judd et al., 2017b), although this appears to
339 vary among species (Jósvai et al., 2016; Judd et al., 2017a).

340 *Relative dosage: less is more?*

341 Our findings strongly corroborate previous reports from individual studies that higher
342 concentrations of host plant volatile relative to sex pheromone can inhibit male Lepidopteran
343 responses to sex pheromones (e.g. Hu et al., 2013; Yu et al., 2015; Xiang et al., 2019). This
344 effect was significant even for the limited number of EAG studies, suggesting that the addition
345 of host plant volatiles at high relative doses do not repel males, given that EAG responses do
346 not differentiate between attractant and repellent effects, but instead interfere with the
347 attraction effect of sex pheromones (Deisig et al., 2014). This response could potentially help

348 males avoid heavily defoliated host plants. If the relative concentration of plant volatile is too
349 low however, it might not have any observable effect (Varela et al., 2011; Barros-Parada et
350 al., 2018). As such, there is likely to be an optimal ratio of plant volatile to sex pheromone,
351 which will depend on the component chemicals and target species. Although ratio of repellent
352 plant volatile to sex pheromone was not included in this meta-analysis due to limited previous
353 research, some evidence suggests that the effect could be simpler, with stronger repellent
354 effects at higher relative doses of plant volatile (Jactel et al., 2011; Wang et al., 2016).
355 Similarly, the effect of relative plant volatile dose on female attraction could be simpler, for
356 example, higher doses of pear ester can increase female capture rates (Knight et al., 2005;
357 Mitchell et al., 2008).

358 *Blend complexity: keep it simple?*

359 Unexpectedly, the use of single plant volatiles with sex pheromones outperformed more
360 complex plant volatile blends in terms of male capture rates in the field. Although a previous
361 meta-analysis of plant volatile effects without sex pheromones found that blend complexity
362 increased effectiveness (Szendrei & Rodriguez-Saona, 2010), our findings suggest that using
363 complex blends with sex pheromones could increase the risk of plant volatiles interfering with
364 male responses to sex pheromones, as discussed above. For instance, in the case of
365 *Grapholita molesta*, the lowest doses of (Z)-3-hexenyl acetate and 1-undecanol increased
366 male capture rates by approximately 4.8 and 3.1 times, respectively, compared with
367 pheromone-only traps (Yu et al., 2015). However, when these two plant volatiles were
368 combined, capture rates were only 2.4 times higher than the pheromone-only controls. The
369 authors hypothesise that this decrease in effectiveness was due to interference among the
370 plant volatile compounds.

371 *Plant volatile categories*

372 There were no consistent differences in the effect of plant volatile compound type on male
373 responses to pheromones across laboratory and field studies. Our findings indicated that

374 green leaf volatiles were associated with the strongest responses of males to sex pheromones
375 in the field, and were the most commonly studied chemical category. In contrast, fruit volatiles
376 (limited to pear ester) were associated with the strongest attractant effects in wind tunnel
377 studies.

378 A wide variety of plant volatiles were reported in the literature. For example, 37 individual
379 compounds were included in our meta-analysis of field trapping studies, in addition to blends
380 of multiple compounds. These were typically selected based on their identified presence in
381 host plants or previous success in attracting the same or similar species. (*Z*)-3-hexenyl acetate
382 was the most frequently studied plant volatile, appearing in 9 of the 35 field trapping studies,
383 while pear ester and phenylacetaldehyde were tested in 7 and 6 field studies, respectively.
384 Other alcohols and aldehydes were also frequently studied, such as (*Z*)-3-hexen-1-ol and (*E*)-
385 2-hexenal which were each tested as unblended volatiles in three field studies. The
386 relationship between plant volatile and growth stage or condition of the host plant (defoliated
387 or undefoliated) could also be relevant in determining their interactive effects with sex
388 pheromones on Lepidoptera (Tang et al., 2012).

389 The variable effects of plant volatile categories are unsurprising given the diversity of
390 compounds within each category, while their effects also depend on target species and
391 background odour. For example, linalool enhanced the attraction of codling moth *Cydia*
392 *pomonella*, but inhibited the attraction of tobacco cutworm *Spodoptera litura* Fabricius, to their
393 respective sex pheromones (Yang et al., 2004; Fang et al., 2018). In addition, the effects of
394 herbivore-induced plant volatiles on phytophagous insect behaviour may depend on their
395 survival strategy, e.g. gregarious versus solitary species (Guo & Wang, 2019). Therefore,
396 understanding the host plant volatile profiles of the target insect species, and their attraction
397 to damaged versus healthy plants, may be important to assemble informed hypotheses for
398 potential future lure development.

399 *Influence of habitat and host specialism*

400 The positive effects (either synergistic or additive) of adding plant volatiles to pheromone traps
401 on male capture rates were higher in herbaceous-crop agriculture than in orchard habitats,
402 despite a similar number of studies in the meta-analysis. This could be because in herbaceous
403 crops, combined traps are often used to increase male captures (e.g. Li et al., 2012b; Miluch
404 et al., 2014; Fang et al., 2018), while in orchard pest management, the aim is often to attract
405 both sexes (e.g. Light, 2016; Knight et al., 2017; Mujica et al., 2018). Only 4 studies were
406 available in forestry contexts, and whilst results were generally positive overall, but non-
407 significant, only one of the studies reported female capture rates (Jósvai et al., 2016).
408 Pheromone traps positioned within host plant tree species have been shown to capture
409 significantly more males of a specialist Lepidoptera than those in suboptimal congeneric hosts
410 (Williams & Jonusas, 2019), while single-species forest stands tend to be more susceptible to
411 specialist pests than generalists (Jactel et al., 2021), indicating potential applications for
412 attractant and repellent plant volatiles for host-specialist Lepidopteran tree pests.

413 Our hypothesis that males of host-specialist Lepidoptera species would exhibit stronger
414 responses to plant volatiles than generalist species was tentatively supported across all three
415 methods but was not statistically significant. As such, the use of plant volatiles to increase
416 male capture rates in pheromone traps might be easier to achieve for host specialist species.

417 *Other potential causes of heterogeneity*

418 Although our analysis provides some insights into the lack of effect or unexpected repellent
419 effects of proposed attractant plant volatiles on male Lepidoptera responses to sex
420 pheromones, other potentially important variables could not be investigated due to insufficient
421 data. Mating experience is one such variable that can influence Lepidopteran responses to
422 sex pheromones and plant volatiles, because newly-mated males can stop responding to sex
423 pheromone and may also change their response to plant volatiles (Deisig et al., 2014). For
424 example, green leaf volatiles of host plants increased the response of unmated but not mated

425 diamondback moth *Plutella xylostella* males to sex pheromone in a wind tunnel experiment
426 (Reddy & Guerrero, 2000). This factor could potentially confound field studies where mating
427 experience is uncontrolled.

428 Weather conditions such as temperature, humidity, and wind speed are also likely to affect
429 success of combining plant volatiles with sex pheromones, and could explain some of the
430 discrepancy between wind tunnel and field trapping results. For example, flight activity of
431 moths is dependent on suitable temperatures and wind speeds, while wind speed also affects
432 trapping area (Elkinton & Cardé, 1988; Schouest Jnr & Miller, 1994; Reardon et al., 2006).

433 The choice and dosage of sex pheromone can also be relevant, although studies have found
434 conflicting evidence of how these factors interact with plant volatiles. Stronger attractant
435 effects have been demonstrated for plant volatiles combined with less effective pheromones
436 (Knight et al., 2014; Miluch et al., 2014; Sans et al., 2016; Borrero-Echeverry et al., 2018), and
437 with pheromones at underdosed or overdosed concentrations (Schmidt-Büsser et al., 2009).
438 However, other studies have found the reverse pattern, where host plant volatiles reduce male
439 attraction to incomplete synthetic pheromones, but increase attraction to optimal pheromones
440 (Sans et al., 2016; Borrero-Echeverry et al., 2018). It is likely that this interaction depends on
441 the combination of pheromone and plant volatile and whether the plant volatile stimulates
442 receptors for missing pheromone components or interferes with pheromone detection (Deisig
443 et al., 2014; Miluch et al., 2014). Clearly initial dosage and release rates of both sex
444 pheromones and plant volatiles are likely to be important factors influencing trap efficacy.

445 Finally, the effect of plant volatiles on Lepidopteran responses to sex pheromone appears to
446 depend on background odour, which can interfere with plant volatile lures when their
447 components overlap (Cai et al., 2017). This perhaps might explain the contrasting results from
448 field studies in different crop types, while findings from laboratory studies in controlled
449 environments are often not corroborated by field studies (Deng et al., 2004; Li et al., 2012b;
450 Tang et al., 2012; Miluch et al., 2014).

451 *Conclusion*

452 This meta-analytic review provides evidence that the addition of attractant plant volatiles to
453 sex pheromone traps leads to higher captures rates of adult Lepidoptera. A key advantage of
454 combined plant volatile-sex pheromone lures is the potential to develop trapping approaches
455 that attract both sexes, which could improve early detection, monitoring, and mass trapping
456 programmes. Although research on proposed repellent plant volatiles is limited, we found
457 preliminary evidence of repellent effects on males towards sex pheromones, which presents
458 opportunities to develop natural pest management strategies such as push-pull and mating
459 disruption approaches.

460 However, effects of attractant plant volatiles on male responses to sex pheromones were
461 highly variable, and in some cases resulted in unexpected repellent effects. We found
462 evidence that this effect depends on factors such as relative concentrations of plant volatile to
463 sex pheromone, category of plant volatile tested, and blend complexity. Hence, our findings
464 demonstrate the potential applications of both attractant and repellent plant volatiles in
465 Lepidoptera pest management, but that careful consideration of attractant lures is critical to
466 minimise interference of plant volatiles on male attraction to sex pheromones. In addition,
467 further research, particularly field trials, is urgently needed to investigate repellent volatiles.

468

469 **Acknowledgements**

470 The funding for the Forest Research work reported in this paper came from a core programme
471 responding to the Science and Innovation Strategy for Great Britain. We are grateful for
472 statistical advice provided by Dr Toni Clarke (Forest Research), and for the comments
473 received from anonymous reviewers that helped to improve the manuscript.

474

475 **Competing Interests**

476 Competing interests: The authors declare none.

477

478 **Supplementary Materials**

479 Supplementary Material 1: List of studies included in the meta-analysis, according to the three
480 methodological categories: electroantennograms (EAG), wind tunnels, and field trapping.

481 Supplementary Material 2: List of Lepidoptera species included in the meta-analysis and their
482 host specialism.

483

484 **References**

485 **Barnes, M. M., Millar, J. G., Kirsch, P. A. & Hawks, D. C.** (1992) Codling Moth
486 (Lepidoptera: Tortricidae) Control by Dissemination of Synthetic Female Sex
487 Pheromone *Journal of Economic Entomology*, **85**(4), 1274–1277.

488 **Barros-Parada, W., Ammagarahalli, B., Basoalto, E., Fuentes-Contreras, E. & Gemeno,**
489 **C.** (2018) Captures of oriental fruit moth, *Grapholita molesta* (Lepidoptera: Tortricidae),
490 in traps baited with host-plant volatiles in Chile *Applied Entomology and Zoology*, **53**(2),
491 193–204.

492 **Bates, D., Mächler, M., Bolker, B. & Walker, S.** (2015) Fitting Linear Mixed-Effects Models
493 Using lme4 *Journal of Statistical Software*, **67**(1), 1–48.

494 **Borrero-Echeverry, F., Bengtsson, M., Nakamuta, K. & Witzgall, P.** (2018) Plant odor and
495 sex pheromone are integral elements of specific mate recognition in an insect herbivore
496 *Evolution*, **72**(10), 2225–2233.

497 **Bruce, T. J. A. & Pickett, J. A.** (2011) Perception of plant volatile blends by herbivorous

- 498 insects – Finding the right mix *Phytochemistry*, **72**(13), 1605–1611.
- 499 **CABI** (2022) *Invasive Species Compendium*, <https://www.cabi.org/isc/>. Retrieved from
500 <https://www.cabi.org/isc/>
- 501 **Cai, X., Bian, L., Xu, X., Luo, Z., Li, Z. & Chen, Z.** (2017) Field background odour should be
502 taken into account when formulating a pest attractant based on plant volatiles *Scientific*
503 *Reports*, **7**(1), 1–10.
- 504 **Cook, S. M., Khan, Z. R. & Pickett, J. A.** (2007) The use of push-pull strategies in
505 integrated pest management *Annual Review of Entomology*, **52**, 375–400.
- 506 **Culliney, T. W.** (2014) Crop losses to arthropods In D. Pimentel & P. Rajinder, eds.,
507 *Integrated Pest Management: Pesticide Problems*, Vol.3, Dordrecht: Springer, , pp.
508 201–225.
- 509 **Deisig, N., Dupuy, F., Anton, S. & Renou, M.** (2014) Responses to pheromones in a
510 complex odor world: sensory processing and behavior *Insects*, **5**(2), 399–422.
- 511 **Deng, J. Y., Wei, H. Y., Huang, Y. P. & Du, J. W.** (2004) Enhancement of attraction to sex
512 pheromones of *Spodoptera exigua* by volatile compounds produced by host plants
513 *Journal of Chemical Ecology*, **30**(10), 2037–2045.
- 514 **Deutsch, C. A., Tewksbury, J. J., Tigchelaar, M., Battisti, D. S., Merrill, S. C., Huey, R.**
515 **B. & Naylor, R. L.** (2018) Increase in crop losses to insect pests in a warming climate
516 *Science*, **361**(6405), 916–919.
- 517 **Eigenbrode, S. D., Birch, A. N. E., Lindzey, S., Meadow, R. & Snyder, W. E.** (2016) A
518 mechanistic framework to improve understanding and applications of push-pull systems
519 in pest management *Journal of Applied Ecology*, **53**(1), 202–212.
- 520 **Elkinton, J. S. & Cardé, R. T.** (1988) Effects of Intertrap Distance and Wind Direction on the
521 Interaction of Gypsy Moth (Lepidoptera: Lymantriidae) Pheromone- Baited Traps
522 *Environmental Entomology*, **17**(5), 764–769.

- 523 **Fang, Y., Zeng, R., Lu, S., Dai, L. & Wan, X.** (2018) The synergistic attractiveness effect of
524 plant volatiles to sex pheromones in a moth *Journal of Asia-Pacific Entomology*, **21**(1),
525 380–387.
- 526 **Fettig, C. J. & Munson, A. S.** (2020) Efficacy of verbenone and a blend of verbenone and
527 nonhost volatiles for protecting lodgepole pine from mountain pine beetle (Coleoptera:
528 Curculionidae) *Agricultural and Forest Entomology*, **22**(4), 373–378.
- 529 **Gregg, P. C., Socorro, A. P. Del & Landolt, P. J.** (2018) Advances in attract-and-kill for
530 agricultural pests: beyond pheromones *Annual Review of Entomology*, **63**, 453–470.
- 531 **Guo, H. & Wang, C. Z.** (2019) The ethological significance and olfactory detection of
532 herbivore-induced plant volatiles in interactions of plants, herbivorous insects, and
533 parasitoids *Arthropod-Plant Interactions* 2019 13:2, **13**(2), 161–179.
- 534 **Hedges, L. V.** (1981) Distribution theory for Glass's estimator of effect size and related
535 estimators *Journal of Educational and Behavioral Statistics*, **6**(2), 107–128.
- 536 **Hu, D., Feng, J., Wang, Z., Wu, H. & Zhang, X.** (2013) Effect of nine plant volatiles in the
537 field on the sex pheromones of *Leguminivora glycinivorella* *Natural Product*
538 *Communications*, **8**(3), 393–396.
- 539 **Jactel, H., Birgersson, G., Andersson, S. & Schlyter, F.** (2011) Non-host volatiles mediate
540 associational resistance to the pine processionary moth *Oecologia*, **166**, 703–711.
- 541 **Jactel, H., Moreira, X. & Castagneyrol, B.** (2021) Tree diversity and forest resistance to
542 insect pests: patterns, mechanisms, and prospects *Annual Review of Entomology*, **66**,
543 277–296.
- 544 **Jósvai, J. K., Koczor, S. & Tóth, M.** (2016) Traps baited with pear ester and acetic acid
545 attract both sexes of *Hedya nubiferana* (Lepidoptera: Tortricidae) *Journal of Applied*
546 *Entomology*, **140**(1–2), 81–90.
- 547 **Judd, G. J. R., Knight, A. L. & El-Sayed, A. M.** (2017a) Development of kairomone-based

- 548 lures and traps targeting *Spilonota ocellana* (Lepidoptera: Tortricidae) in apple orchards
549 treated with sex pheromones *The Canadian Entomologist*, **149**(5), 662–676.
- 550 **Judd, G. J. R., Knight, A. L. & El-Sayed, A. M.** (2017b) Trapping *Pandemis limitata*
551 (Lepidoptera: Tortricidae) moths with mixtures of acetic acid, caterpillar-induced apple-
552 leaf volatiles, and sex pheromone *The Canadian Entomologist*, **149**(6), 813–822.
- 553 **Knight, A., Cichon, L., Lago, J., Fuentes-Contreras, E., Barros-Parada, W., Hull, L.,**
554 **Krawczyk, G., Zoller, B., Hansen, R., Hilton, R. & Basoalto, E.** (2014) Monitoring
555 oriental fruit moth and codling moth (Lepidoptera: Tortricidae) with combinations of
556 pheromones and kairomones *Journal of Applied Entomology*, **138**(10), 783–794.
- 557 **Knight, A. L., El-Sayed, A. M., Judd, G. J. R. & Basoalto, E.** (2017) Development of 2-
558 phenylethanol plus acetic acid lures to monitor obliquebanded leafroller (Lepidoptera:
559 Tortricidae) under mating disruption *Journal of Applied Entomology*, **141**(9), 729–739.
- 560 **Knight, A. L., Hilton, R. & Light, D. M.** (2005) Monitoring Codling Moth (Lepidoptera:
561 Tortricidae) in apple with blends of ethyl (E, Z)-2,4-decadienoate and codlemone
562 *Environmental Entomology*, **34**(3), 598–603.
- 563 **Kong, W. N., Wang, Y., Guo, Y. F., Chai, X. H., Li, J. & Ma, R. Y.** (2020) Behavioral effects
564 of different attractants on adult male and female oriental fruit moths, *Grapholita molesta*
565 *Pest Management Science*, **76**(9), 3225–3235.
- 566 **Kvedaras, O. L., Del Socorro, A. P. & Gregg, P. C.** (2007) Effects of phenylacetaldehyde
567 and (Z)-3-hexenyl acetate on male response to synthetic sex pheromone in *Helicoverpa*
568 *armigera* (Hübner) (Lepidoptera: Noctuidae) *Australian Journal of Entomology*, **46**(3),
569 224–230.
- 570 **Landolt, P. J. & Phillips, T. W.** (1997) Host plant influences on sex pheromone behavior of
571 phytophagous insects *Annual Review of Entomology*, **42**, 371–391.
- 572 **Larson, E. R., Graham, B. M., Achury, R., Coon, J. J., Daniels, M. K., Gambrell, D. K.,**

- 573 **Jonasen, K. L., King, G. D., LaRacuenta, N., Perrin-Stowe, T. I. N., Reed, E. M.,**
574 **Rice, C. J., Ruzi, S. A., Thairu, M. W., Wilson, J. C. & Suarez, A. V.** (2020) From
575 eDNA to citizen science: emerging tools for the early detection of invasive species
576 *Frontiers in Ecology and the Environment*, **18**(4), 194–202.
- 577 **Lehmann, P., Ammunét, T., Barton, M., Battisti, A., Eigenbrode, S. D., Jepsen, J. U.,**
578 **Kalinkat, G., Neuvonen, S., Niemelä, P., Terblanche, J. S., Økland, B. & Björkman,**
579 **C.** (2020) Complex responses of global insect pests to climate warming *Frontiers in*
580 *Ecology and the Environment*, **18**(3), 141–150.
- 581 **Lenth, R. V.** (2021) emmeans: Estimated Marginal Means, aka Least-Squares Means. R
582 package version 1.5.5-1., <https://CRAN.R-project.org/package=emmeans>.
- 583 **Li, J., Valimaki, S., Shi, J., Zong, S., Luo, Y. & Heliovaara, K.** (2012a) Attraction of the
584 gypsy moth to volatile organic compounds (VOCs) of damaged Dahurian larch
585 *Zeitschrift Fur Naturforschung - Section C Journal of Biosciences*, **67 C**(7–8), 437–444.
- 586 **Li, P., Zhu, J. & Qin, Y.** (2012b) Enhanced attraction of *Plutella xylostella* (Lepidoptera:
587 Plutellidae) to pheromone-baited traps with the addition of green leaf volatiles *Journal*
588 *of Economic Entomology*, **105**(4), 1149–1156.
- 589 **Li, X., Liu, Y., Duan, M., Yu, Z. & Axmacher, J. C.** (2018) Different response patterns of
590 epigaeic spiders and carabid beetles to varying environmental conditions in fields and
591 semi-natural habitats of an intensively cultivated agricultural landscape *Agriculture,*
592 *Ecosystems and Environment*, **264**, 54–62.
- 593 **Light, D. M.** (2016) Control and monitoring of codling moth (Lepidoptera: Tortricidae) in
594 walnut orchards treated with novel high-load, low-density “meso” dispensers of sex
595 pheromone and pear ester *Environmental Entomology*, **45**(3), 700–707.
- 596 **Ma, M., Chang, M. M., Lei, C. L. & Yang, F. L.** (2016) A garlic substance disrupts odorant-
597 binding protein recognition of insect pheromones released from adults of the

598 angoumois grain moth, *Sitotroga cerealella* (Lepidoptera: Gelechiidae) Insect Molecular
599 Biology, **25**(5), 530–540.

600 **Mafra-Neto, A., Wright, M., Fettig, C., Progar, R., Munson, S., Blackford, D., Moan, J.,**
601 **Graham, E., Foote, G., Borges, R., Silva, R., Lake, R., Bernardi, C., Saroli, J.,**
602 **Clarke, S., Meeker, J., Nowak, J., Agnello, A., Martini, X., Rivera, M. J. & Stelinski,**
603 **L. L. (2022)** Repellent semiochemical solutions to mitigate the impacts of global climate
604 change on arthropod pests In J. Coats, C. Corona, & M. Debboun, eds., *Advances in*
605 *Arthropod Repellents*, London: Academic Press, , pp. 279–322.

606 **McCormick, A. C., Heyer, J., Sims, J. W., Mescher, M. C. & De Moraes, C. M. (2017)**
607 Exploring the effects of plant odors, from tree species of differing host quality, on the
608 response of *Lymantria dispar* males to female sex pheromones *Journal of Chemical*
609 *Ecology*, **43**, 243–253.

610 **McNair, C., Gries, G. & Gries, R. (2000)** Cherry bark tortrix, *Enarmonia formosana*:
611 olfactory recognition of and behavioral difference by nonhost angio- and gymnosperm
612 volatiles *Journal of Chemical Ecology*, **26**(4), 809–821.

613 **Meagher, J. (2001)** Trapping fall armyworm (Lepidoptera: Noctuidae) adults in traps baited
614 with pheromone and a synthetic floral volatile compound *Florida Entomologist*, **84**(2),
615 288–292.

616 **Miluch, C. E., Dossall, L. M. & Evenden, M. L. (2014)** Factors influencing male *Plutella*
617 *xylostella* (Lepidoptera: Plutellidae) capture rates in sex pheromone-baited traps on
618 canola in western Canada *Journal of Economic Entomology*, **107**(6), 2067–2076.

619 **Mitchell, V. J., Manning, L.-A., Cole, L., Suckling, D. M. & El-Sayed, A. M. (2008)** Efficacy
620 of the pear ester as a monitoring tool for codling moth *Cydia pomonella* (Lepidoptera:
621 Tortricidae) in New Zealand apple orchards *Pest Management Science*, **64**, 209–214.

622 **Mujica, V., Preti, M., Basoalto, E., Cichon, L., Fuentes-Contreras, E., Barros-Parada,**

- 623 **W., Krawczyk, G., Nunes, M. Z., Walgenbach, J. F., Hansen, R. & Knight, A. L.**
624 (2018) Improved monitoring of oriental fruit moth (Lepidoptera: Tortricidae) with terpinyl
625 acetate plus acetic acid membrane lures Journal of Applied Entomology, **142**(8), 731–
626 744.
- 627 **Nakagawa, S., Lagisz, M., Jennions, M. D., Koricheva, J., Noble, D. W. A., Parker, T. H.,**
628 **Sánchez-Tójar, A., Yang, Y. & O’Dea, R. E.** (2022) Methods for testing publication
629 bias in ecological and evolutionary meta-analyses Methods in Ecology and Evolution,
630 **13**(1), 4–21.
- 631 **R Core Team** (2020) R: A language and environment for statistical computing., Vienna: R
632 Foundation for Statistical Computing. Retrieved from <https://www.r-project.org/>
- 633 **Reardon, B. J., Sumerford, D. V. & Sappington, T. W.** (2006) Impact of Trap Design,
634 Windbreaks, and Weather on Captures of European Corn Borer (Lepidoptera:
635 Crambidae) in Pheromone-Baited Traps Journal of Economic Entomology, **99**(6),
636 2002–2009.
- 637 **Reddy, G. V. P. & Guerrero, A.** (2000) Behavioral responses of the diamondback moth,
638 *Plutella xylostella*, to green leaf volatiles of *Brassica oleracea* subsp. *capitata* Journal of
639 Agricultural and Food Chemistry, **48**(12), 6025–6029.
- 640 **Reddy, G. V. P. & Guerrero, A.** (2004) Interactions of insect pheromones and plant
641 semiochemicals Trends in Plant Science, **9**(5), 253–261.
- 642 **Rizvi, S. A. H., George, J., Reddy, G. V. P., Zeng, X. & Guerrero, A.** (2021) Latest
643 developments in insect sex pheromone research and its application in agricultural pest
644 management Insects, **12**(484), 1–26.
- 645 **Rohatgi, A.** (2021) WebPlotDigitizer, Pacifica, California, USA:
646 <https://automeris.io/WebPlotDigitizer>.
- 647 **Ryall, K. L., Silk, P. J., Mayo, P., Crook, D., Khrimian, A., Coss, A. A., Sweeney, J. &**

- 648 **Scarr, T.** (2012) Attraction of *Agrilus planipennis* (Coleoptera: Buprestidae) to a volatile
649 pheromone: effects of release rate, host volatile, and trap placement *Environmental*
650 *Entomology*, **41**(3), 648–656.
- 651 **Sans, A., Morán, M., Riba, M., Guerrero, Á., Roig, J. & Gemeno, C.** (2016) Plant volatiles
652 challenge inhibition by structural analogs of the sex pheromone in *Lobesia botrana*
653 (Lepidoptera: Tortricidae) *European Journal of Entomology*, **113**, 579–586.
- 654 **Schmidt-Büsser, D., Von Arx, M. & Guerin, P. M.** (2009) Host plant volatiles serve to
655 increase the response of male European grape berry moths, *Eupoecilia ambiguella*, to
656 their sex pheromone *J Comp Physiol A*, **195**, 853–864.
- 657 **Schouest Jnr, L. P. & Miller, T. A.** (1994) Automated Pheromone Traps Show Male Pink
658 Bollworm (Lepidoptera: Gelechiidae) Mating Response Is Dependent on Weather
659 Conditions *Journal of Economic Entomology*, **87**(4), 965–974.
- 660 **Szendrei, Z. & Rodriguez-Saona, C.** (2010) A meta-analysis of insect pest behavioral
661 manipulation with plant volatiles *Entomologia Experimentalis et Applicata*, **134**(3), 201–
662 210.
- 663 **Tang, R., Zhang, J. P. & Zhang, Z. N.** (2012) Electrophysiological and behavioral
664 responses of male fall webworm moths (*Hyphantria cunea*) to herbivory-induced
665 mulberry (*Morus alba*) leaf volatiles *PLoS ONE*, **7**(11), 1–7.
- 666 **van Lierop, P., Lindquist, E., Sathyapala, S. & Franceschini, G.** (2015) Global forest area
667 disturbance from fire, insect pests, diseases and severe weather events *Forest Ecology*
668 *and Management*, **352**, 78–88.
- 669 **Varela, N., Avilla, J., Anton, S. & Gemeno, C.** (2011) Synergism of pheromone and host-
670 plant volatile blends in the attraction of *Grapholita molesta* males *Entomologia*
671 *Experimentalis et Applicata*, **141**(2), 114–122.
- 672 **Viechtbauer, W.** (2010) Conducting meta-analyses in R with the metafor package *Journal of*

- 673 Statistical Software, **36**(3), 1–48.
- 674 **Wang, F., Deng, J., Schal, C., Lou, Y., Zhou, G., Ye, B., Yin, X., Xu, Z. & Shen, L.** (2016)
675 Non-host plant volatiles disrupt sex pheromone communication in a specialist herbivore
676 Scientific Reports, **6**(1), 1–8.
- 677 **Weissling, T. J. & Knight, A. L.** (1996) Oviposition and Calling Behavior of Codling Moth
678 (Lepidoptera: Tortricidae) in the Presence of Codlemone Annals of the Entomological
679 Society of America, **89**(1), 142–147.
- 680 **Williams, D. T. & Jonusas, G.** (2019) The influence of tree species and edge effects on
681 pheromone trap catches of oak processionary moth *Thaumetopoea processionea* (L.)
682 in the U.K. Agricultural and Forest Entomology, **21**(1), 28–37.
- 683 **Willis, K. J.** (Ed.) (2017) State of the World's Plants 2017, Royal Botanic Gardens, Kew.
- 684 **Wittman, J. T., Silk, P., Parker, K. & Aukema, B. H.** (2021) Optimizing early detection
685 strategies: defining the effective attraction radius of attractants for emerald ash borer
686 *Agrilus planipennis* Fairmaire Agricultural and Forest Entomology, **23**(4), 527–535.
- 687 **Witzgall, P., Kirsch, P. & Cork, A.** (2010) Sex pheromones and their impact on pest
688 management Journal of Chemical Ecology, **36**(1), 80–100.
- 689 **Xiang, H. M., Chen, Z., Li, X. W., Guo, Y. Q., Li, X. C. & Ma, R. Y.** (2019) Two terpenoids
690 activates close mating behavior and enhances trap efficiency of sex pheromone of
691 *Grapholita molesta* Journal of Asia-Pacific Entomology, **22**(4), 1109–1114.
- 692 **Xiao, C., Gregg, P. C., Hu, W., Yang, Z. & Zhang, Z.** (2002) Attraction of the cotton
693 bollworm, *Helicoverpa armigera* (Hübner) (Lepidoptera: Noctuidae), to volatiles from
694 wilted leaves of a non-host plant, *Pterocarya stenoptera* Applied Entomology and
695 Zoology, **37**(1), 1–6.
- 696 **Xu, H. & Turlings, T. C. J.** (2018) Plant volatiles as mate-finding cues for insects Trends in
697 Plant Science, **23**(2), 100–111.

698 **Yang, Z., Bengtsson, M. & Witzgall, P.** (2004) Host plant volatiles synergize response to
699 sex pheromone in codling moth, *Cydia pomonella* *Journal of Chemical Ecology*, **30**(3),
700 619–629.

701 **Yu, H., Feng, J., Zhang, Q. & Xu, H.** (2015) (Z)-3-hexenyl acetate and 1-undecanol
702 increase male attraction to sex pheromone trap in *Grapholita molesta* (Busck)
703 (Lepidoptera: Tortricidae) *International Journal of Pest Management*, **61**(1), 30–35.

704

705

706 **Tables**

707 **Table 1** Subgroup analysis for attractant plant volatiles, showing the subsets of data applied to each
 708 subgroup and level of replication.

Subgroup	Levels	Data subset	Number of data points (and studies)		
			EAG	Wind tunnel	Field trapping
Sex	Male	Excludes unsexed data	87 (9)	807 (21)	213 (27)
	Female		39 (2)	129 (3)	134 (20)
Plant volatile blend	Single	Males	78 (8)	572 (17)	143 (23)
	Multiple		9 (4)	235 (13)	70 (15)
Habitat	Agriculture (herbaceous crops)		N/A	N/A	108 (12)
	Forest				19 (4)
	Orchard				86 (13)
Specialism	Generalist		30 (4)	352 (10)	86 (14)
	Specialist		57 (5)	455 (11)	127 (13)
Trap type	Delta	N/A	N/A	73 (14)	
	Inverted cone			10 (2)	
	Panel			2 (1)	
	Unitrap			22 (3)	
	Water trap			100 (6)	
Plant volatile category	Fruit volatile	Males, excludes blends of >1 type	0	13 (2)	18 (5)
	GLV		34 (5)	256 (14)	83 (12)
	Homoterpene		0	14 (3)	1 (1)
	Monoterpene/oid		16 (5)	100 (11)	18 (6)
	Organosulfur		0	0	3 (1)
	Phenylpropanoid		22 (4)	125 (9)	28 (7)
Sesquiterpene	7 (2)	68 (7)	7 (2)		
Ratio (standardised amount of plant volatile (PV) per unit of sex pheromone (SP))	PV:SP > mean*	Males, experiments where >1 ratio tested	18 (2)	123 (10)	50 (13)
	PV:SP < mean*		56 (2)	370 (10)	73 (13)

709 * mean ratio calculated separately for each study

710

711

712 **Figure Captions**

713 **Figure 1.** Cumulative number of studies included in the meta-analysis by publication date, for the three
714 different methods (electroantennogram, wind tunnel, and field trapping). Each study reported the effects
715 of combined sex pheromone and plant volatile lures versus pheromone-only lures on antennal or
716 behavioural responses of Lepidopteran pests.

717 **Fig. 2** Effect (with 95% confidence intervals) of combined plant volatile and sex pheromone lures,
718 versus pheromone only, on the responses of adult Lepidoptera according to six subgroups (rows) and
719 three methods (columns). Positive effects represent higher responses to the combined lures compared
720 with pheromone-only lures. One subgroup, trap type in field-trapping studies, is not shown because
721 none of the levels were significant and the results were not informative. Data was filtered according to
722 each subgroup, as shown in Table 1. Asterisks denote where there was a significant difference between
723 levels of the subgroup (*F* test, with intercepts), where *** = $p < 0.001$, ** = $p < 0.01$, and * = $p < 0.05$.
724 Abbreviations: PV = plant volatile, SP = sex pheromone.

725 **Fig. 3** Effect (with 95% confidence intervals) of combined attractant plant volatile and sex pheromone
726 lures, versus pheromone-only lures, on male and female Lepidoptera according to wind tunnel stage.
727 Positive values are associated with stronger effects of the combined lure versus the pheromone.

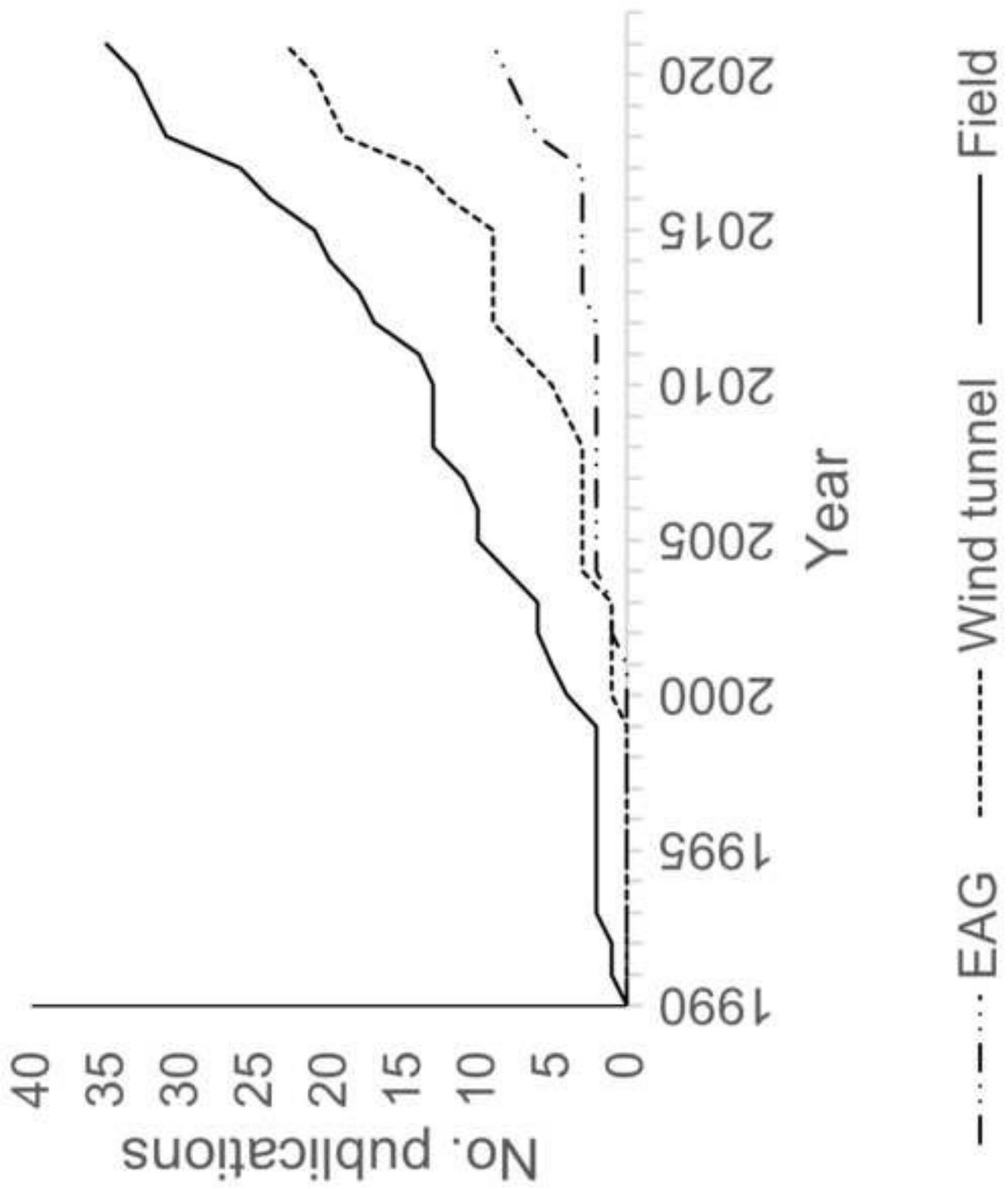
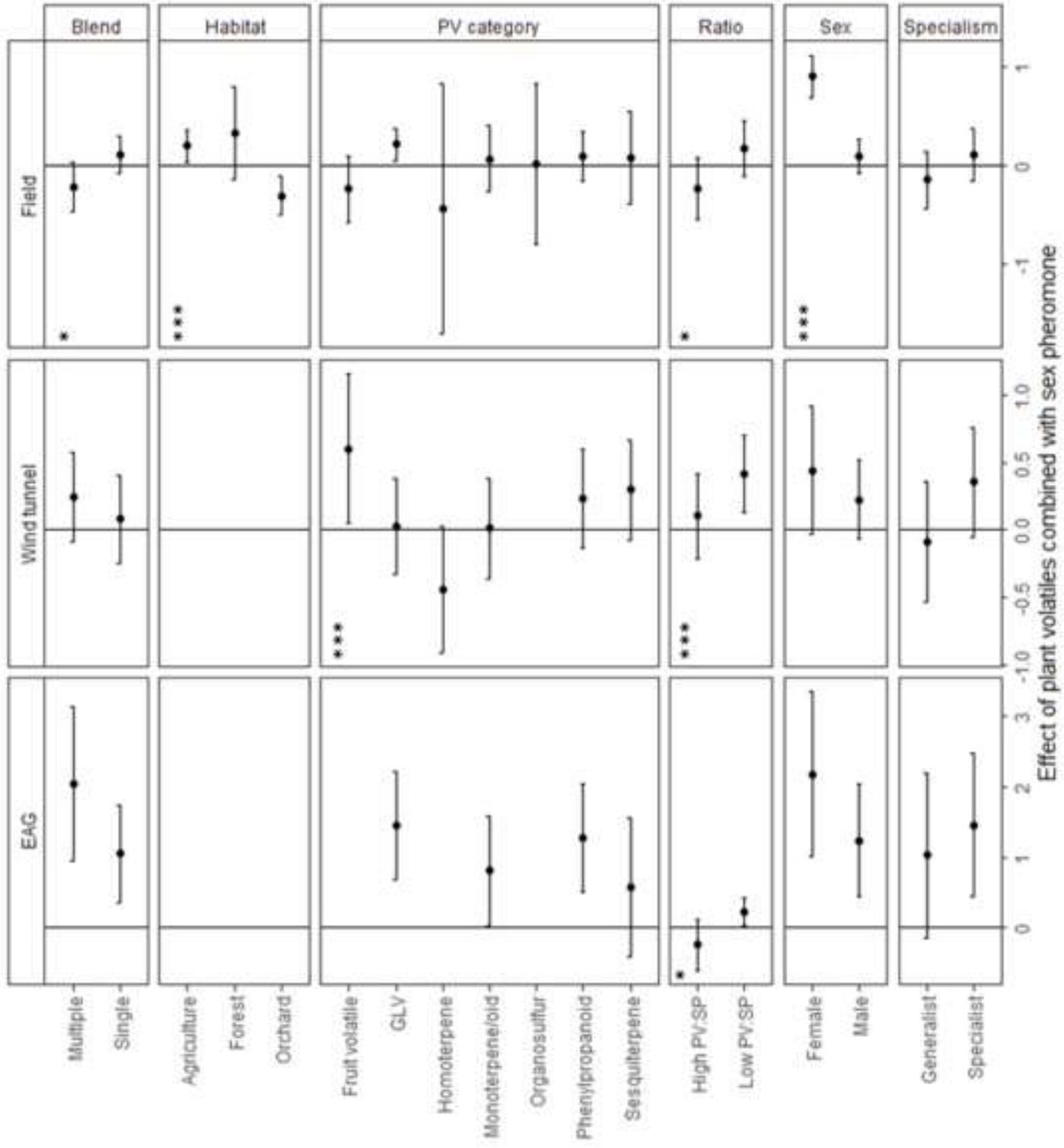
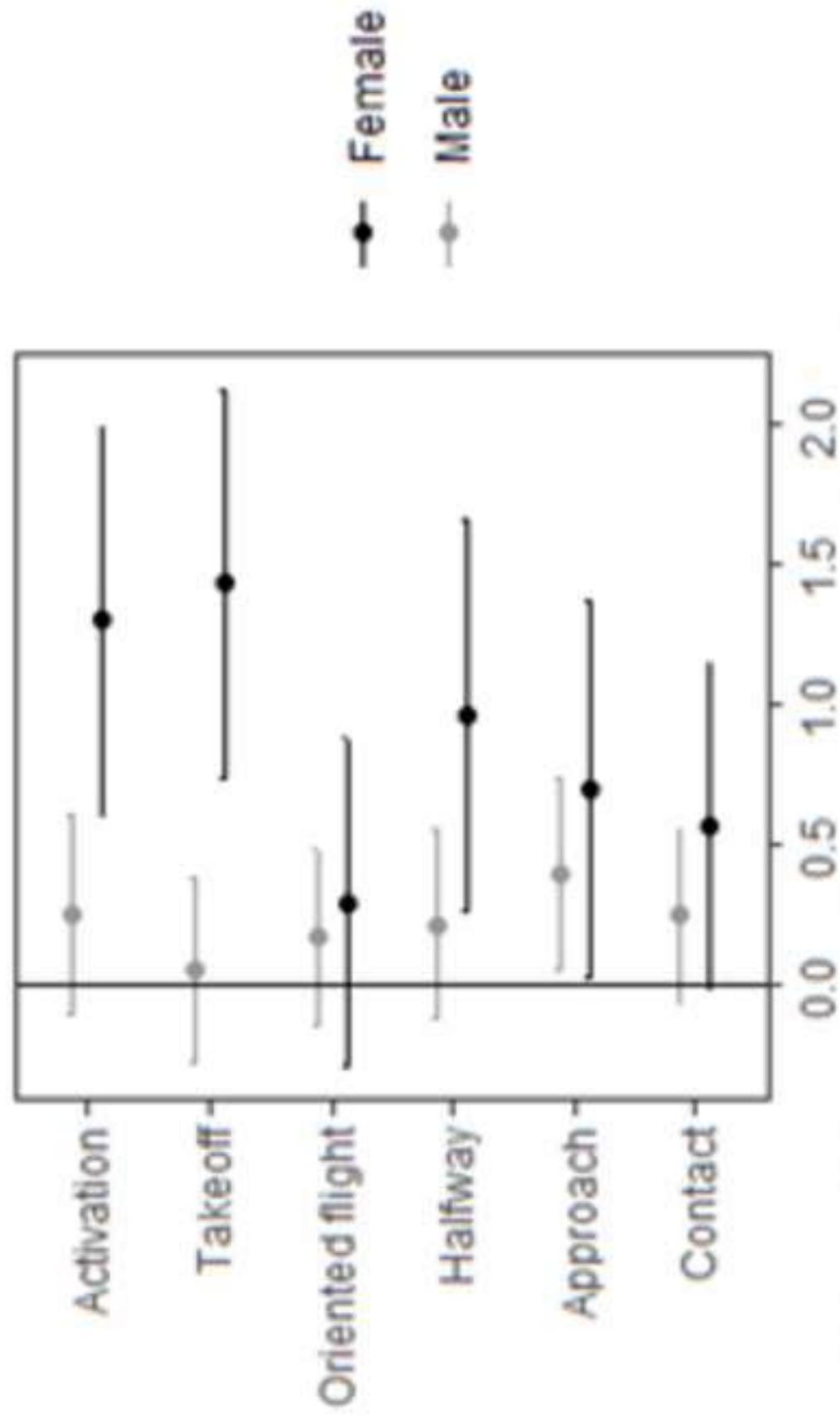


Figure 1

Figure 2





Effect of plant volatiles combined with sex pheromone