

1 Juvenile hormone regulates reproductive physiology and the production of fertility cues in the
2 swarm-founding wasp *Polybia occidentalis*

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16 **Abstract**

17 Juvenile hormone (JH) has important functions that regulate insect life. In adult individuals, it
18 induces gonadotropic and behavioral changes. Manipulating JH levels helps to understand
19 how it influences insect physiology. The effects of JH on Epiponini swarm-founding wasps
20 have shown contrasting results, affecting reproduction, chemical compound expression,
21 behavior, and age polyethism. In this study, we investigated whether JH affects reproductive
22 physiology and production of fertility cues in a swarm-founding wasp species *Polybia*
23 *occidentalis* in an age-controlled experimental setup. We treated newly emerged females with
24 methoprene (JH analogue) and precocene-I (JH inhibitor) to determine their effects on ovary
25 activation and cuticular hydrocarbon (CHC) expression. Furthermore, we compared the
26 chemical profiles of treated workers with those of queens. Our results show that methoprene
27 and precocene-I affected the CHC production in *P. occidentalis*. Additionally, females treated
28 with methoprene were chemically more similar to queens than precocene- and acetone-treated
29 females. Methoprene affected ovarian status (increasing ovary activation). These results
30 suggest that different levels of JH reflect changes in chemical and reproductive traits in *P.*
31 *occidentalis* females. Furthermore, we point out the existence of fertility cues in this
32 Epiponini wasp.

33 **Keywords:** Epiponini wasp, hormonal treatment, methoprene, precocene.

34 **1- Introduction**

35 Juvenile hormone (JH) is a lipid hormone from the sesquiterpenoid class of terpenes,
36 found in diverse natural products. It is synthesized by the *corpora allata*, a pair of glands
37 localized in the insect brains (Nijhout 1994; Hartfelder. 2000; Huang 2020). JH has important
38 functions in the regulation of adult life in social Hymenoptera (Hartfelder, 2000; Huang
39 2020). In social wasps, there is evidence that JH regulates age polyethism and acts as a
40 gonadotropic agent, increasing ovarian activation and controlling cuticular hydrocarbon
41 (CHCs) expression, which in turn acts as a fertility cue (O'Donnell and Jeanne, 1993; Oi et al.
42 2015a; Oliveira et al. 2017; Walton et al., 2020).

43 Manipulating JH levels (using JH analogue methoprene and anti-JH precocene) helps
44 to understand how JH influences insect physiology, especially given the diversification of
45 associated functions across different insect lineages (Slamá, 1971; Robinson, 1992;
46 Ramaseshadri et al., 2012; Kestrup et al., 2014a; Kestrup et al., 2014b). Methoprene is a JH
47 analogue that triggers similar effects as pure JH in the physiology and behavior of insects. On
48 the other hand, precocene is a compound that acts as an anti-JH, destroying the *corpora allata*
49 and consequently inhibiting JH production (Bowers, 1976; O'Donnell and Jeanne, 1993;
50 Robinson and Vargo, 1997; Giray et al., 2005; Burns et al., 2007; Gotoh et al., 2008). Social
51 insect workers treated with methoprene are known to upregulate the production of queen-like
52 chemical profiles, and in contrast, precocene-treated workers express worker-like chemical
53 profiles (Oliveira et al., 2017; Oi et al., 2021a,b). Some specific CHs act as queen signals in
54 different social insect species and the comparison of their chemical profiles reveal that
55 specific CHCs are conserved across different social insect lineages (Van Oystaeyen et al.,
56 2014; Kather and Martin, 2015; Holman, 2018). Indeed, the JH has the function of controlling
57 some features of solitary ancestral species, such as reproduction and the production of fertility
58 cues (Oi et al. 2015b). In social wasps, JH furthermore displays a gonadotropic role in
59 females by increasing ovary activation and oviposition (Tibbetts and Sheehan, 2012 – *Polistes*
60 *metricus*; Kelstrup et al., 2014b – *Synoeca surinama*; Oi et al., 2021a – *Vespula vulgaris*;
61 Prato et al., 2021 – *Polybia occidentalis*; Ferreira et al., 2022 - *Mischocyttarus cerberus* and
62 *Myschocyttarus cassununga*), and is associated with colony ontogeny and pheromone
63 regulation (Oliveira et al., 2017 – *V. vulgaris*; Walton et al., 2020 – *Polistes fuscatus*; Oi et al.
64 2021b – *Polistes dominula* and *Polistes satan*; reviewed in Tibbetts et al., 2020; Ferreira et
65 al., 2022 – *M. cerberus* and *M. cassununga*). Additionally, JH is a pleiotropic hormone,
66 meaning that the endocrine system regulates two or more features in the same individual
67 (Dantzer and Swanson, 2017; Oliveira et al. 2017; Oi et al., 2020; Oi et al., 2021a, b).

68 JH may not have a conserved role in Epiponini swarm-forming wasps. Previous
69 studies demonstrated that JH regulates reproduction in *S. surinama* but not in *Polybia micans*
70 (Kelstrup et al. 2014a, b). In *P. occidentalis*, it was reported that JH influences ovarian status,
71 behavior, and chemical profile (O'Donnell and Jeanne 1993; Prato et al. 2021), and we
72 confirmed that JH influenced age polyethism (Prato et al. 2021). Additionally, the
73 morphological caste differentiation evolved several times in the Epiponini to different degrees
74 (Noll and Wenzel, 2008; Noll et al., 2020) and it is likely to be under JH influence. It is thus
75 not clear how JH acts on Epiponini wasps. This poorly studied yet species-rich tribe presents
76 a wide phylogenetic range of biological characteristics. For instance, morphological castes
77 and ovarian activation in workers are present in some species (*Protopolybia* and
78 *Chartergellus*, respectively) but not in others (*Synoeca*) (Noll et al., 2020). The nests of
79 swarm-forming wasps are polygynic and, in some species, queens and workers express
80 morphological and physiological differences (Noll et al., 2020). For instance, in *P.*
81 *occidentalis*, workers organize themselves in an age-polyethism system, which means that
82 newly emerged females perform tasks within the nest, while older workers perform high-cost
83 tasks, such as foraging (Jeanne, 1991; Jeanne et al., 1992; O'Donnell and Jeanne, 1992).
84 Newly emerged females exhibit ovary activation which decreases as they age (O'Donnell,
85 2001). Another group of females, namely the intermediaries, show ovarian activation, but are
86 not inseminated (Noll and Zucchi, 2000). The number of queens varies depending on the size
87 and stage of the nest, and workers often eliminate queens during nest development (Forsyth,
88 1978). In summary, these biological characteristics provide an interesting opportunity for
89 intra-genus comparative studies and to understand how the endocrine system can affect the
90 morphological development type (post-imaginal and pre-imaginal) (Kelstrup et al., 2014a).

91 Here, we focus on the effects that JH triggers in Epiponini wasps: induction of ovarian
92 activation, acceleration of age polyethism, and cuticular chemical production. We investigate
93 whether JH is responsible for regulating ovary activation and the production of CHCs in *P.*
94 *occidentalis* females kept under controlled conditions. This contrasts from our previous study
95 (Prato et al. 2021), which was conducted in the field and focused on the influence of JH on
96 the division of tasks. Studies have indeed demonstrated that experiments conducted under
97 field and laboratory conditions affect differently the traits under investigation such as
98 differences in physiology, behavior, gene expression, and development time (Calisi and
99 Bentley, 2009; Campbell et al., 2009; Jandt et al., 2015). We thus performed our experiments
100 in the laboratory to control for the influence of age, interaction among workers, and food
101 intake. Furthermore, we investigated whether JH is responsible for upregulating the

102 production of CHCs that may be linked with ovary activation, and thus might represent key
103 compounds to inform fertility status in the species. To do this, we compared the chemical
104 profile of treated workers with those of fertile queens. We hypothesized that reproductive and
105 chemical traits of *P. occidentalis* are directly affected by their circulating levels of JH. We
106 predicted that females treated with methoprene would express higher levels of ovary
107 activation and their cuticular surface would become more queen-like. Conversely, we
108 predicted the opposite effects for females treated with precocene that is low levels of ovarian
109 activation and chemical profile contrasting from queen-like.

110

111 **2- Material and Methods**

112 2.1 – Study site and Experiments

113 The experiments were conducted in the Laboratório de Comportamento e Ecologia de
114 Insetos Sociais at the Universidade de São Paulo (USP), Ribeirão Preto Campus
115 (21°09'50.7"S, 47°51'32.1"W) between August and October 2016. Two nests of *P.*
116 *occidentalis* were collected in the field and kept under laboratory conditions to sample newly
117 emerged females. The newly emerged females were treated topically on the abdomen with
118 either a single dose of 1 µL of a solution at 5 µg/µl of methoprene (MT) Pestanal®
119 (SUPELCO, analytical standard) (25 individuals) or a single dose of 1 µL of a solution at 5
120 µg/µl of precocene-I (PC) (SIGMA-ALDRICH) (25 individuals) diluted in acetone. The
121 solvent group (25 individuals) was treated with acetone (AC) (2 µl of acetone). The choice of
122 solution concentration was based on previous toxicity tests (Prato et al., 2021). The females
123 from each group were paint-marked with non-toxic ink (Magic®) and kept in a plastic box
124 with sucrose solution, mealworm larvae, and a piece of the nest for 10 days. We chose 10
125 days based on a previously published work, which reported that at a later age, almost all the
126 females of *P. occidentalis* have filamentous, not activated ovaries (O'Donnell, 2001). After
127 10 days, the treated females were euthanized in the temperature freezer for posterior
128 extraction of CHs, and dissection to check their ovarian activation. For comparison, six
129 queens (Q) were collected from a third nest belonging to the same region to analyze the
130 CHCs, ovarian status, and insemination.

131 2.2 - CHCs analyses

132 Cuticular hydrocarbons were extracted in hexane solvent (Macron Fine Chemicals,
133 95% n-Hexane) by immersing the body for two minutes. The solvent from the extract was left
134 to evaporate in a flow chamber for 24 hours. The vial with the extract was resuspended in 50

135 μL of hexane and 2 μL of this solution was injected (Splitless mode) in a Gas
136 Chromatography - Mass Spectrometer GC-MS (Shimadzu, model QP2010), using a DB-5MS
137 column (length 30 m), with the injector temperature at 280 °C and the helium carrier gas flow
138 set at 1 mL/min. The oven temperature started at 150 °C, rising by 7 °C/min until 260 °C, and
139 was kept at this temperature for 5 min. After, the temperature was further increased to 310 °C
140 at a 5 °C/min rate and held for 10 min. An external alkane standard solution C21-C40 (Fluka
141 analytical) was used to identify the linear hydrocarbons. The methylated compounds were
142 identified based on the mass spectrometric fragmentation patterns (ion and molecular mass)
143 (Carlson et al., 1998).

144 2.3 – Reproductive trait categorization

145 The ovaries of treated females were dissected in saline solution and categorized into
146 two types, activated or non-activated (Noll and Zucchi, 2000). The ovary of the queens were
147 categorized as activated (mature oocytes ready for laying), shown in Fig. 1a. Treated females
148 were categorized in the following way: activated ovary of workers (small cells and oocytes
149 and few mature oocytes) (Fig. 1b), and non-activated ovary of workers (filamentous
150 ovarioles) (Fig. 1c).

151 2.4 – Statistical analyses

153 All statistical analyses were performed using R software (version 4.0.2, R core team
154 2018). The area of the peaks in the chromatograms were transformed in relative amounts. The
155 CHs from the different treatments were compared using the Permanova test (Bray-Curtis
156 distance). For that, we used the *adonis* function from the *vegan* package (Oksanen et al.,
157 2013). To visually represent our multivariate chemical dataset and check for cluster formation
158 based on their CHs according to different treatments, a principal component analysis (PCA)
159 was performed with the *prcomp* function of the *stats* package (Team RC et al. 2013). To
160 identify the compounds that most contributed to the separation of the groups, a discriminant
161 analysis (SIMPER) was performed using 999 permutations. We used the *simper* function from
162 the *vegan* package (Oksanen et al. 2013). Furthermore, the main compounds that were
163 important in the group separations (results from SIMPER analysis) were divided into three
164 different chemical classes (linear alkanes, methyl alkanes, and alkenes). The classes of
165 compounds and the individual compounds per group were analyzed using a Mann-Whitney
166 test.

167 The Generalized Linear Model was done to analyze the ovarian status of the
168 workers between the treatments. The ovary was the dependent variable, and the treatment and
169 origin nest as the explanatory variables. A post hoc Tukey test was used to compare the
170 treatments. Those statistical analyses used the *lme4*, *emmeans*, *effects*, and *HH* packages (R
171 Development Core Team 2008; Bates et al. 2020).

172

173 3- Results

174

175 3.1- CH differences

176 The cuticle of *P. occidentalis* expressed a chemical profile that included 65
177 compounds, which ranged from 22 to 40 carbons chain length (Table 1 supplementary
178 material). The differences between treated workers and queens were both qualitative and
179 quantitative. The CHs profiles from workers treated with methoprene and precocene were
180 significantly different when compared to the solvent group (Permanova: $F_{93,3}$: 38.66; $p < 0.01$)
181 (Fig 2; Table 2 supplementary material). The most important compounds that contributed to
182 the separation of the treatments were methyl alkanes (66.7%), linear alkanes (21%), alkenes
183 (8.8%), and unknown compounds (3.5%). The main compounds responsible for groups'
184 separation varied mainly between the chain length from n -C₂₅ to n -C₃₉.

185 The relative abundances of each chemical compound class varied among the treated
186 workers. The proportion of linear alkanes differed among all the treatments ($p < 0.01$), except
187 acetone and precocene (Mann-Whitney - $W = 356$; $p = 0.40$) (Fig. 3a; Table 1). In this case,
188 workers from the methoprene treated group showed the highest proportion of linear alkanes
189 (Table 3 supplementary material). The linear alkanes n -C₂₇, n -C₂₈, n -C₂₉, and n -C₃₁ were
190 higher and statistically different in the methoprene treated group when compared with the
191 precocene and acetone treatment groups (Fig. 3b; Table 2). The proportion of methylated
192 alkanes varied among all the treatments (Fig. 3a; Table 1). Considering methylated
193 compounds only, there were also specific methylated compounds that were higher and
194 statistically different in workers from the methoprene treated group when compared with the
195 two others (e.g. 13-;11-;9MeC₂₇; 3MeC₂₇; 3.13-;3.11diMeC₂₇ and 3.15-;3.13-
196 ;3.11diMeC₂₉) (Fig. 4; Table 3). Workers from the precocene-treated group presented a
197 higher proportion of methylated compounds than the other treatments (Table 3 supplementary
198 material). Workers from the methoprene treated group expressed the highest proportion of
199 alkenes and this class of compound varied among all the treatments, but it was not different
200 between workers from the acetone and precocene treated groups ($W = 223.5$; $p = 0.08$) (Fig

201 3a; Table 1 and Table 3 supplementary material). Interestingly, the alkene *z*-C₂₉-1 occurred in
202 a higher proportion in workers from the methoprene treated group when compared with
203 precocene and acetone groups. The alkene *z*-C₂₉-2 was exclusively present in workers from
204 the methoprene group. Finally, some compounds remained unidentified due to their low-
205 quality mass spectra. These compounds were likely methylated hydrocarbons and were
206 present in workers from the methoprene and acetone groups ($W = 515.5$; $p < 0.01$) and
207 appeared in a higher proportion in the acetone group (Fig. 3a; Table 1 and Table 3
208 supplementary material).

209 The chemical profile of queens was statistically different to the chemical profile of
210 workers (Permanova: $F_{98,4} = 34.75$; $p < 0.01$) (Fig 2; Table 4 supplementary material). The
211 classes of compounds varied among the treated workers and queens. The abundance of linear
212 alkanes was significantly similar between workers treated with acetone and precocene, but
213 they differed from methoprene and queens groups. These compounds were higher in
214 abundance in workers treated with methoprene and queens, and they did not differ statistically
215 from each other ($W = 64$; $p = 0.59$) (Fig 3a; Table 4 and Table 5 supplementary material). The
216 methylated alkanes appeared in a higher proportion in the workers treated with precocene and
217 this compound class varied between all the workers treated groups and queens ($p \leq 0.01$),
218 except in methoprene and queens ($W = 56$; $p = 0.35$) (Fig. 3a; Table 4 and Table 5
219 supplementary material). Lastly, workers treated with methoprene were the ones who had the
220 higher proportion of alkenes. We found in our samples CHCs that have already been
221 highlighted in the literature as queen pheromones in other social wasps (*n*-C₂₇, 3MeC₂₇, *n*-C₂₈,
222 *n*-C₂₉, *n*-C₃₁) (Van Oystaeyen et al. 2014; Oi et al. 2016). We observed that these linear
223 alkanes and 3-MeC 27 appeared in higher proportions in workers treated with methoprene,
224 which were statistically different when compared to other treated groups. (Fig 3b; Table 2 and
225 3). Also, these compounds in workers treated with methoprene were statistically different in
226 relation to the queen group, except the *n*-C₂₉ (Fig 3b; Table 5).

227

228 3.2 –Reproductive traits

229 Workers belonging to different groups expressed different levels of ovary activation
230 (Treatment - Anova Chisq = 22.42, $p < 0.001$ ***; Nest - Anova Chisq = 4.11, $p = 0.042$ *).
231 Workers treated with acetone expressed mostly non-activated ovaries (76% versus 24% of
232 activated ovaries). The same trend was detected in workers treated with precocene, in which
233 88% of them expressed non-activated ovaries, while only 12% had activated ovaries. On the

234 other hand, workers treated with methoprene exhibited a higher rate of ovary activation (65%
235 of them), whereas only 35% had non-activated ovaries. The pairwise comparisons among the
236 treatments were significant between workers treated with acetone and methoprene (estimate =
237 2.47, z ratio = 3.26, $p = 0.001^{**}$) and workers treated with methoprene and precocene
238 (estimate = -2.98, z ratio = -3.73, $p = 0.0005^{***}$) (Fig. 5). However, workers treated with
239 acetone and precocene did not differ from each other (estimate = -0.51, z ratio = -0.62, $p =$
240 0.529) (Fig. 5).

241

242 **4- Discussion**

243 Our results demonstrate that the JH analogue (methoprene) and the JH inhibitor
244 (precocene) influenced the chemical profiles in treated workers of *P. occidentalis*. However,
245 only the methoprene treatment influenced ovarian activation. These results are in line with
246 previously published data for other social wasp species (Robinson and Vargo 1997; Oliveira
247 et al. 2017; Oi et al. 2021a, b; Ferreira et al., 2022).

248 Workers of *P. occidentalis* treated with methoprene chemically resembled queens but
249 were consistently different to other two groups of workers. The most evident sign of chemical
250 modulation in workers treated with methoprene is the increased proportion of some linear
251 alkanes. In contrast, workers treated with precocene suffered a decrease in their proportion of
252 linear alkanes and increased the level of methyl-alkane compounds when compared with the
253 queen and other treated workers. The variation that we detected in the chemical profile of
254 workers treated with methoprene and precocene probably occurred due to the treatments
255 affecting their JH titers. Previous studies have shown that the topical administration of JH
256 analogue is linked directly with the production of some long-chain hydrocarbons in social
257 wasps (Oliveira et al., 2017; Oi et al., 2020; Oi et al., 2021b). Our results show an increase in
258 this type of hydrocarbon. The linear alkanes (between C₂₇ to C₃₁) were the most important
259 compounds responsible to separate workers treated with methoprene and queens from the
260 acetone group. The four linear alkanes (*n*-C₂₇, *n*-C₂₈, *n*-C₂₉, and *n*-C₃₁) and one alkene (*z*-C₂₉)
261 in higher quantity may indicate putative fertility cue in *P. occidentalis* wasps and could
262 potentially be candidates for queen pheromones in this species. Those compounds were
263 previously reported for other wasps' species (Sledge et al. 2004 – *P. dominulus*; van Zweden
264 et al. 2013 - *Dolichovespula saxonica*; Van Oystaeyen et al. 2014 – *V. vulgaris*; Oi et al.
265 2015a – *V. vulgaris*). Altogether, our results support the idea that queens and fertile females
266 overproduce specific saturated hydrocarbons, which is shown in several lineages of
267 Hymenoptera (Van Oystaeyen et al. 2014; Kather and Martin, 2015). However, the difference

268 in the chemical profiles encountered among the workers and queens also can be related to
269 their different colonies of origin, since the CHCs have been reported to vary in individuals of
270 the same species belonging to different nests (Dapporto et al., 2006; Tannure-Nascimento et
271 al., 2007; Martin et al., 2008; Lorenzi and d'Ettoire, 2020). On the other hand, queens and
272 methoprene treated workers had differences only in the alkenes class, which suggests that
273 they are more chemically similar to each other. Nevertheless, our results need to be
274 interpreted carefully as only two nests were used in the experiments, meaning that larger
275 sample size may be more representative of the species.

276 Workers treated with methoprene increased ovarian activation when compared to
277 precocene and acetone treatments; this result is not surprising as JH has been shown to have a
278 gonadotropic effect in other social insects (Bloch et al. 2000; Shorter and Tibbetts 2009;
279 Tibbetts et al. 2013; Kelstrup et al. 2014b; Amsalem et al. 2014a; Oliveira et al. 2017; Walton
280 et al., 2020; Oi et al. 2020; Oi et al. 2021a, b; Prato et al. 2021; Ferreira et al., 2022).
281 However, in the species *A. mellifera* and *P. micans*, JH does not influence ovarian activation
282 but does control the chemical signaling and age polyethism (Fluri et al. 1981; Robinson 1992;
283 Hartfelder, 2000; Kelstrup et al. 2014a). It is possible that the JH acts as a pleiotropic
284 hormone in *P. occidentalis* affecting multiple traits, such as behavioral maturation, fertility,
285 and chemical signaling. However, we cannot discard the hypothesis that in our study the
286 CHCs changed as a by-product of ovary activation that was caused by hormonal treatments.

287 In summary, our results provide experimental evidence that the reproduction and the
288 CHCs production are under JH hormonal control in *P. occidentalis*. These changes in CHCs
289 may act not only in the chemical communication among workers linked to age polyethism
290 (Prato et al., 2021) but also in the communication between queens and workers. This study
291 helps to unveil functions associated with JH in the evolution of the *Polybia* lineage. The
292 presence of some queen-characteristic linear alkanes, that were also detected to be
293 upregulated by reproductive workers (treated with methoprene) indicate the existence of
294 fertility cues in this species. Future studies conducting bioassays with the queen characteristic
295 compounds are necessary to test their role in the division of reproductive labor in colonies of
296 *P. occidentalis*.

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Author contributions

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Declaration of competing interest

No competing interests declared.

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584 **Table captions**

585

586 **Table 1** Results from comparison of the compound classes among the treated females that were
587 important in the separation of the treatment groups (AC - acetone, MT - methoprene, and PC -
588 precocene). * Significant p-values ($p < 0.05$)

589

590 **Table 2** Comparisons of the main linear alkanes among the treated females that were important in the
591 separation of the treatment groups (AC - acetone, MT - methoprene, and PC - precocene). *
592 Significant p-values ($p < 0.05$)

593

594 **Table 3** Comparisons of the methyl alkanes among treated females that were important in the
595 separation of the treatment groups. * Significant p-values ($p < 0.05$). n- compound is not present in
596 these groups.

597

598 **Table 4** Comparisons of important compounds classes in separating treatment groups and queens (AC
599 - acetone, MT - methoprene, PC - precocene, and Q - queen) * Significant p-values ($p < 0.05$)

600

601 **Table 5** Comparisons of the queens' chemical compounds reported in the literature among treated
602 workers and collected queens (Van Oystaeyen et al. 2014; Oi et al. 2016). * Significant p-values ($p <$
603 0.05)

604

605

606 **Figures captions**

607

608

609 **Fig. 1** Ovarian patterns found in *Polybia occidentalis* workers treated and queens. a- queen activated
610 ovary; b - worker activated ovary and c- worker non-activated ovaries

611

612 **Fig. 2** Principal components analysis (PCA) based on the relative abundances of the chemical
613 compounds present in their cuticle of *Polybia occidentalis* females". Red - acetone; Blue - precocene;
614 Green - methoprene; Yellow and Purple - queens. Highlighted compounds: 3MeC₂₇, n-C₂₇, n-C₂₉,
615 MeC₂₉ = 15-;13-;11-;9-7MeC₂₉, MeC₃₁ = 15-;13-;11-;9MeC₃₁ and MeC₃₉ = 15-;13MeC₃₉

616

617 **Fig. 3** a) Relative abundance of different compound classes (linear alkanes, methyl alkanes, and
618 alkenes) found in *Polybia occidentalis* according to the group of treated workers (acetone,
619 methoprene, and precocene) and queen. b) Specific compounds of linear alkanes and methyl alkane n-
620 C₂₇, 3MeC₂₇, n-C₂₈, n-C₂₉, n-C₃₁ found in *Polybia occidentalis* according to the group of treated
621 workers (acetone, methoprene, and precocene) and queen. The compounds represented in "b"
622 correspond to the ones that were previously demonstrated to act as queen pheromones in other social
623 wasps (Van Oystaeyen et al. 2014; Oi et al. 2016). Different letters indicate statistical differences
624 between groups of a given compound

625

626 **Fig. 4** Abundance of different methylated alkanes found in *Polybia occidentalis* workers according to
627 the treatment groups. **1-** 13-;11-;9MeC₂₇; **2-** 7MeC₂₇; **3-** 5MeC₂₇; **4-** 3MeC₂₇; **5-** 5.13diMeC₂₇; **6-**
628 3.13-;3.11diMeC₂₇; **7-** 3.7diMe₂₇; **8-** 4MeC₂₈; **9-** 4.14-;4.12diMeC₂₈; **10-** 9.13diMeC₂₉; **11-** 5.15-
629 ;5.13-;5.11diMeC₂₉; **12-** 3.15-;3.13-;3.11diMeC₂₉) (The data are presented as mean and standard
630 deviations). Different letters indicate statistical differences between treatments of a given compound

631

632 **Fig. 5** The bars correspond to the percentages of females that expressed ovary activation among
633 different treatments methoprene, precocene and acetone group in *Polybia occidentalis* workers.
634 Significance level (** = $p < 0.01$; *** = $p < 0.001$)

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636