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This is the author's manuscript

Original Citation:

Availability:

This version is available http://hdl.handle.net/2318/1884144 since 2023-05-29T10:11:46Z

Published version:

DOI:10.1007/s10340-022-01576-3

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08 November 2026

No adverse effects of symbiotic control on the parasitism of *Halyomorpha halys* by egg parasitoids

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Abstract

The brown marmorated stink bug *Halyomorpha halys* is a polyphagous insect, which has a devastating impact on agricultural production in many countries. The alteration of symbiont vertical transmission, by removing symbionts from stink bug eggs (symbiotic control), has been recently introduced in control programmes against this insect. A major advantage of this strategy is the compatibility with natural enemies, since it allows an insecticide-free approach that is not harmful to other agroecosystem components. However, the effect of anti-symbiont products on parasitism by egg parasitoids is still unexplored. Here, we investigated the impact on parasitism by native (*Anastatus bifasciatus*, *Ooencyrtus telenomicida* and *Trissolcus kozlovi*) and exotic (*Trissolcus japonicus* and *Trissolcus mitsukurii*) parasitoids that attack *H. halys* eggs, after treatment with the micronutrient biocomplex Dentamet®, used for symbiotic control. The native wasp species were tested in no-choice bioassays, showing that treatment of the egg masses did not affect emergence percentages, but the non-reproductive effects were often reduced by the biocomplex. The exotic species *T. japonicus* and *T. mitsukurii* were used in no-choice and paired choice bioassays, showing

an opposite influence of Dentamet® on emergence percentage and preference in the two species. No-choice tests indicated the highest successful parasitoid emergence on biocomplex-treated egg masses for *T. japonicus*, while no preference in the paired comparison with eggs treated with water or untreated. In contrast, *T. mitsukurii* displayed the lowest parasitism after Dentamet® treatment in no-choice tests, and preferred egg masses without Dentamet® in paired choice tests. We did not record any natural symbiont acquisition by the parasitoids emerged from *H. halys* egg masses, indicating that the wasp fitness is very unlikely to be altered by dysbiotic effects resulting from treatments. Therefore, our results support a further implementation of symbiotic control in different crops in combination with biological control, as sustainable options for *H. halys* integrated pest management.

Keywords symbiont-targeted control, biological control, brown marmorated stink bug, *Trissolcus japonicus*, *Trissolcus mitsukurii*, integrated pest management

1. Introduction

The brown marmorated stink bug *Halyomorpha halys* (Stål) (Hemiptera: Pentatomidae) is one of the most studied pentatomid species due to its invasive potential and high polyphagy (Leskey et al. 2012a; Rice et al. 2014; Leskey and Nielsen 2018). This insect pest is native to Asia, and it was unintentionally introduced in North America in the 1990s and later in Europe in 2004 (Haye and Weber 2017; Leskey and Nielsen 2018). *Halyomorpha halys* attacks more than 300 species of many important agricultural crops and wild plants (Lee et al. 2013). In fruits, feeding activity induces symptoms such as deformation and discolouration, and seed abortion with early attacks (Rice et al. 2014; Bariselli et al. 2016; Bergmann et al. 2016; Bosco et al. 2018). Moreover, it is also an important household nuisance pest due to its aggregative overwintering behaviour inside buildings (Inkley 2012). Because of its invading potential, the population dynamics of *H. halys* have been studied under

different environmental conditions, providing important information for predicting the potential spread. A survival reduction was observed in the presence of high temperature and/or low humidity, suggesting that the insect may preferentially spread in areas where heatwaves are not frequent (Scaccini et al. 2019; Fisher et al. 2020; Khadka et al. 2020).

Currently, the control of *H. halys* mainly relies on the use of chemical pesticides, with an important environmental impact and a huge effect on human health (Nielsen et al. 2008; Leskey et al. 2012b). Moreover, most insecticides used against this pest have broad-spectrum activity, which negatively affects natural enemies and pollinators (Leskey et al. 2012a, b; Cira et al. 2017; Kuhar and Kamminga 2017). Several methods have been proposed to reduce chemical treatments within Integrated Pest Management Crop Perimeter Restructuring programmes, e.g. border row or alternate row insecticide applications (Blaauw et al. 2015; Akotsen-Mensah et al. 2020; Ludwick et al. 2020).

The main alternative to insecticides against *H. halys* is biological control. Egg parasitoids are the most studied and specialized natural enemies both in the native (Lee 2015) and in the invaded areas (Abram et al. 2017; Conti et al. 2021). In the native area, the most common egg parasitoids belong to Scelionidae and Eupelmidae (Lee et al. 2013). In particular, the samurai wasp *Trissolcus japonicus* (Ashmead) (Hymenoptera: Scelionidae) is reported as the predominant species in China and in Japan (Yang et al. 2009; Yang et al. 2015; Zhang et al. 2017; Kamiyama et al. 2022). In Europe, *Anastatus bifasciatus* (Geoffroy) (Hymenoptera: Eupelmidae) is the main native species found emerging from field-laid and sentinel *H. halys* egg masses in Italy and Switzerland (Haye et al. 2015; Costi et al. 2019; Moraglio et al. 2020; Zapponi et al. 2020). Furthermore, the generalist *Ooencyrtus telenomicida* (Vassiliev) (Hymenoptera: Encyrtidae) was obtained from frozen sentinel *H. halys* eggs in Central Italy (Roversi et al. 2016), while *Trissolcus kozlovi* Rjachovskij (Hymenoptera: Scelionidae) was found emerging from field-laid *H. halys* eggs in North Italy (Moraglio et al. 2020; Scaccini et al. 2020). Both *A. bifasciatus* and *T. kozlovi* were tested in augmentative release field trials, and proved to attack *H. halys* freshly laid eggs, although neither was able to effectively contain the bug population in the experimental conditions (Stahl et al. 2018, 2019a, 2019b; Moraglio et al.

2021a). Therefore, *T. japonicus* was considered as the most promising candidate for *H. halys* biological control, and host range studies were started in support of the application for authorization to the field release of the exotic parasitoid, which must be submitted and approved by regulatory agencies (Haye et al. 2020; Sabbatini-Peverieri et al. 2021). In Italy, the field release of *T. japonicus* has been recently authorized, and started in the summer of 2020. Meanwhile, adventive populations of *T. japonicus* have been detected in Switzerland and in Italy in 2017 and in 2018, respectively (Sabbatini-Peverieri et al. 2018; Stahl et al. 2019c; Moraglio et al. 2020). Furthermore, also adventive populations of *Trissolcus mitsukurii* (Ashmead) (Hymenoptera: Scelionidae), another Asian parasitoid of *H. halys* eggs in Japan (Arakawa and Namura 2002, Kamiyama et al. 2022), have been found in Europe, in Italy and in France in 2016 and 2021, respectively (Scaccini et al. 2020; Bout et al. 2021). This species has been considered as another promising candidate for biological control (Sabbatini-Peverieri et al. 2020; Caron et al. 2021; Giovannini et al. 2022). However, biological control is seriously hindered by the use of insecticides (Lowenstein et al. 2019).

Recently, symbiotic control protocols were designed to prevent symbiont acquisition by nymphs by treating *H. halys* eggs with antibacterial substances; this approach was proposed as a suitable option for pentatomid containment (Gonella et al. 2019, 2020). In this insect family, gut symbiotic bacteria are transmitted through maternal secretions, which are smeared on egg masses during oviposition and orally acquired by the nymphs after egg hatching (Prado et al. 2006; Otero-Bravo and Sabree 2015). Since these obligate bacteria are essential for growth, development and survival of the insect, symbiont-deprived stink bugs usually display reduced survival or fitness (Taylor et al. 2014). Different active substances have been tested to eliminate the primary symbiont of *H. halys*, i.e. "*Candidatus* Pantoea carbekii" (*P. carbekii*), from the egg surface, showing high mortality percentages soon after nymph emergence (Mathews and Barry 2014; Taylor et al. 2017; Gonella et al. 2019). These nymphs tested negative for *P. carbekii* through specific qPCR analyses, indicating that the observed effects resulted from missed symbiont acquisition (Gonella et al. 2019). In particular, the application of the zinc, copper and citric acid biocomplex Dentamet® (Diachem, Italy)

on egg masses caused more than 90% first instar nymph mortality. Therefore, in Italy foliar applications with Dentamet® were introduced in control programmes targeting field-laid egg masses of *H. halys* in several crops, as the product was provisionally authorized to control *H. halys* by the Italian Ministry of Agricultural, Food and Forestry Policies in 2021. Since this product does not show a direct insecticidal activity, it is expected not to have any effects on the survival of beneficial insects, including egg parasitoids; however, its effects on trophic interactions involving natural enemies — such as host preference and exploitation efficiency — have not yet been investigated.

Here, we assessed the impact of Dentamet® on parasitism by native and exotic egg parasitoids. Specifically, we measured parasitoid emergence percentages on treated and untreated egg masses under laboratory conditions. We conducted no-choice tests with three native (*A. bifasciatus*, *O. telenomicida*, *T. kozlovi*) and two exotic (*T. japonicus* and *T. mitsukurii*) egg parasitoids species, to assess the suitability for parasitic wasps of egg masses exposed to the biocomplex. Moreover, paired choice tests were performed using the species that showed the highest emergence percentage in no-choice tests, to evaluate their actual preference when treated egg masses were compared to untreated ones. Since several studies have shown the horizontal transmission of symbionts between parasitoids and their hosts, in some cases offering a beneficial effect to the new host (Vavre et al. 1999; Gualtieri et al. 2017; Qi et al. 2019), we also evaluated the possible acquisition of *P. carbekii* from *H. halys* egg masses in the co-evolved parasitoids, i.e. the Asian *Trissolcus* species.

2. Materials and methods

2.1 Insect rearing

Overwintered *H. halys* adults were collected in 2021 from wild and cultivated host plants in several sites in the Piedmont region, NW Italy. Insects were maintained in climatic chambers at 25 ± 1 °C, 65 ± 5% RH and 16:8 h L:D photoperiod, in net cages (930 × 475 × 475 mm) containing broad bean (*Vicia faba* L.) seedlings, apples (*Malus domestica* Borkh.), shelled hazelnuts (*Corylus avellana* L.)

127 and green beans (*Phaseolus vulgaris* L.). *Halyomorpha halys* rearing was inspected daily to collect
128 freshly laid egg masses (less than 24 hours old) to be used in this study.
129 A colony of *T. kozlovi* had already been established in laboratory since 2017 (Moraglio et al. 2021a),
130 while the other native (*A. bifasciatus* and *O. telenomicida*), and exotic egg parasitoids (*T. japonicus*
131 and *T. mitsukurii*) were obtained from field-collected parasitized egg masses of *H. halys*, sampled
132 from several sites in Piedmont in 2021. All wasp species were maintained separately from stink bug
133 rearing, in a climatic chamber at the same conditions indicated above, on fresh or frozen *H. halys* egg
134 masses, in plastic containers (100 mm diameter, 50 mm height) with a mesh on the lid, wet cotton
135 and a honey drop (applied with a needle) were provided on the lid and weekly replaced.

136 **2.2 No-choice tests**

137 No-choice tests were conducted by exposing three types of *H. halys* egg masses to the five egg
138 parasitoid species: (a) untreated, (b) treated with ultrapure water and (c) treated with Dentamet® 1%
139 v/v (dissolved in ultrapure water). Prior to the beginning of experiments, the egg masses were
140 randomly assigned to different treatments; the number of eggs per mass was recorded, then the eggs
141 were not further manipulated in order to avoid any disturbance to parasitism. The egg masses were
142 individually placed in plastic Petri dishes (60 mm diameter) and treated with the biocomplex or water
143 by a 200 mL hand sprayer under a fume hood, according to Gonella et al. (2019). A single spray (651
144 ± 7.42 µl) was applied about 20 cm away from the Petri dish. After 30 minutes from spray application,
145 when the treated egg mass was dry, a single 5-10-day-old mated female of each parasitoid species
146 was placed into a Petri dish with a single *H. halys* fresh egg mass of one of the three types and
147 maintained for 48 h. The 48 h period was selected to allow the parasitization even by the less efficient
148 egg parasitoids. Every female was used only once. All wasps were fed with a small drop of honey,
149 placed at the top of Petri dish. Egg mass visit an oviposition were recorded according to Haye et al.
150 (2015). In addition, an equal number of egg masses were treated as explained above, but they were
151 not exposed to any parasitoid species, to evaluate natural host mortality. Each trial (one egg mass per

152 treatment offered to each of the five parasitoid species, plus one egg mass per treatment unexposed
153 to any wasp) was replicated 10 times (for a total of 180 egg masses). All trials were conducted in a
154 climatic chamber at 26 ± 1 °C, 65 ± 5% RH. After the experiment, the egg parasitoids were removed,
155 and all egg masses were individually reared until the emergence of stink bug nymphs and/or wasp
156 adults. The number of unhatched *H. halys* eggs, and the number of emerged parasitoids were recorded
157 for each egg mass. The two species showing the highest parasitoid efficiency in no-choice tests were
158 used to perform a paired choice test.

159 **2.3 Paired choice tests**

160 After no-choice tests, paired choice tests were conducted with the two most efficient wasp species,
161 namely *T. japonicus* and *T. mitsukurii*, to check for a possible interference with host preference
162 caused by treatments. All wasps and egg masses were collected from the laboratory colonies as
163 described above. A single-mated 5-10-day-old wasp female of each species was placed in an arena
164 (h = 70 mm; diameter = 95 mm) with two egg masses; females were used only once. Each arena was
165 composed of the following pairs of egg masses: (A) untreated *vs* treated with water, (B) treated with
166 water *vs* treated with Dentamet® 1% v/v, and (C) untreated *vs* treated with Dentamet® 1% v/v. Each
167 comparison was replicated 10 times, and a total of 120 egg masses were used for this experiment. A
168 small drop of honey was provided in the arena as a food source for the wasp. The female was left in
169 the arena for 3 h according to the observations of Haye et al. (2020), who suggested a limited test
170 duration to reduce the number of wasps parasitizing both egg masses. The first egg mass visited by
171 the parasitoid was visually checked and recorded. At the end of the experiment, wasp females were
172 removed, and the egg masses were reared separately at 26 ± 1 °C, 65 ± 5% RH and 16:8 L:D
173 photoperiod until the emergence of all *H. halys* nymphs and/or parasitoid adults. The numbers of
174 unhatched eggs and emerged parasitoids were recorded for each egg mass.

175 **2.4 Molecular diagnosis for *P. carbekii***

To evaluate the occurrence of a host-to-parasitoid horizontal transmission of *P. carbekii*, quantitative Real Time PCR (qPCR) analysis was used to determine the presence of the *H. halys* symbiont in exotic wasps emerged from no-choice tests. Adults of *T. japonicus* and of *T. mitsukurii* were collected after the emergence from untreated (field-collected) egg masses and stored at -80 °C in RNAlater™ (Sigma-Aldrich, MO, USA). Three specimens emerging from the same egg mass were sampled from 10 egg masses parasitized by each species; hence a total of 30 individuals from each exotic species were used. RNA extraction was performed with the "SV Total RNA Isolation System" (Promega, WI, USA) according to the manufacturer instructions. RNA quality and concentration were assessed with a ND-1000 spectrophotometer (NanoDrop, DE, USA). First-strand cDNA was synthesized by "Reverse Transcription System" (Promega) and Random Primers. cDNA was used as a template for qPCR analyses with the *P. carbekii*-specific primers PcarQF/PcarQR as described by Gonella et al. (2019). Reactions were performed on a CFX ConnectTM Real-Time PCR Detection System (Bio-Rad, CA, USA) in 25 µl volume containing: 12.5 µl of SsoAdvancedTM Universal SYBR® Green Supermix (Bio-Rad), 0.1 µl of 100 µM forward and reverse primer, 11.3 µl of sterile water, and 1 µl of cDNA template. Standard curves were constructed with cloned PCR-amplified 16S rRNA gene of *P. carbekii*. Standard clones were obtained using the pGEM T-easy Vector Cloning Kit (Promega, WI, USA). The detection limit was calculated as the lowest concentration of cloned amplicons used for determining the standard curves that were successfully amplified, corresponding to 4.60 gene copies / sample. An additional qPCR targeting the insect's 18S rRNA gene (MqFw / MqRv) was performed, to verify if parasitoids emerged from treated egg masses were truly devoid of *P. carbekii* or whether the absence of symbiont was due to sample quality. Primers were used according to Marzachì and Bosco (2005), under the conditions described by Gonella et al. (2015).

2.5 Statistical analysis

In no-choice tests, parasitoid emergence (mean number of successfully parasitized eggs per egg mass) and egg mortality (mean number of unhatched eggs per egg mass) were evaluated for each parasitoid

201 species considering: (i) only egg masses in which at least one parasitoid successfully emerged, or (ii)
202 all offered egg masses (N=10) per treatment. Separately for each parasitoid species, means were then
203 compared among treatments using a generalized linear model (GLM) with a binomial probability
204 distribution and a logit link function, followed by a pairwise Bonferroni post hoc test ($P \leq 0.05$). For
205 binomial probability distribution, success / fail conditions were considered as the numbers of
206 parasitized / non-parasitized eggs and unhatched eggs / eggs with any emergence.
207 In paired choice tests, the percentages of wasps parasitizing only one exposed egg mass were
208 compared with a Pearson's Chi-square test. The number of emerged parasitoids in parasitized egg
209 masses and the number of unhatched egg masses were then evaluated using a GLM with a binomial
210 probability distribution followed by a sequential Bonferroni post hoc test ($P \leq 0.05$). Replicates in
211 which both egg masses were parasitized were not included in the analysis.
212 Statistical analyses were carried out with SPSS Statistics 27 (IBM Corp. Released 2020, Armonk,
213 NY, USA).

214 ## 3. Results

215 ### 3.1 No-choice tests

216 Females of all egg parasitoid species were alive at the end of the tests, with no apparent fitness
217 perturbation; moreover, they all were able to oviposit on *H. halys* eggs. However, only exotic species
218 were able to successfully parasitize eggs in all the 10 replicates, whereas native species proved to
219 successfully parasitize eggs in 5 up to 9 of the 10 replicates (Table 1). For this reason, the results
220 from native wasps were submitted to two separate statistical analyses, shown in Table 1 (considering
221 only parasitized egg masses), and Fig. 1 and Table S1 (considering all replicates). The emergence
222 percentage from *H. halys* eggs was generally higher in the exotic species than in the others (Table 1).
223 Specific responses observed for each parasitoid species are described below.
224 *Anastatus bifasciatus*: when only successfully parasitized egg masses were considered, emergence of
225 *A. bifasciatu*s was significantly different between treatments ($P < 0.001$), with the lowest percentage

of wasps emerging from egg masses treated with Dentamet® (Table 1). Moreover, higher percentages of unhatched eggs were found in untreated and treated with water egg masses (Table 1). However, when considering all replicates, there were no significant differences among treatments in parasitoid emergence, while Dentamet® still showed the lowest percentage of unhatched eggs compared to the other two groups (Fig. 1, Table S1).

Ooencyrtus telenomicida: according to both statistical evaluations (considering only parasitized egg masses or all replicates), females of this native species did not show significant differences in emergence among treatments (Table 1, Fig. 1, Table S1). Treatment with Dentamet® produced a significant reduction of unhatched eggs when only parasitized egg masses were analysed ($P < 0.001$) (Table 1).

Trissolcus kozlovi: among the tested native species, the highest emergence percentages were observed for this species, without significant differences among the three treatments in both statistical evaluations (only parasitized or all egg masses) (Table 1, Fig. 1, Table S1). Egg masses treated with Dentamet® showed significantly lower percentages of unhatched eggs than the other two groups (Fig. 1, Table 1).

Trissolcus japonicus: emergence was significantly higher in egg masses treated with Dentamet® than in the two other groups ($P < 0.001$) (Table 1). Statistical analyses confirmed a significant reduction of egg mortality in the Dentamet® treatment with respect to the other groups (Table 1).

Trissolcus mitsukurii: despite the percentage of parasitoid emergence being higher than 80%, egg masses treated with Dentamet® showed significantly lower emergence ($P < 0.001$) than the other groups, and significantly higher percentages of unhatched eggs only in comparison to the group treated with water (Table 1).

Unexposed egg masses: the percentage of eggs that successfully hatched was very high (average 84.26%), and no significant differences were found in egg mortality in different treatments (Table 1).

3.2 Paired choice tests

Due to their higher efficiency, *T. japonicus* and *T. mitsukurii* were selected for paired choice tests. Trials revealed that some *T. japonicus* and *T. mitsukurii* females were able to parasitize both egg masses within 3 h (Fig. 2, Table 2). For these replicates, the first choice was assessed by visual observation, and selection responses were compared with a Pearson's Chi-square test (Table 2). The first egg masses that *T. japonicus* females visited always corresponded to the parasitized egg masses. In cases where both egg masses were parasitized, the first visit was recorded on the egg mass that showed the higher number of emerged parasitoids. The same behaviour was observed for *T. mitsukurii*, except for two cases in the untreated *vs* treated with water pairs. In these replicates, both egg masses were parasitized but the higher number of parasitoids emerged from the second visited egg mass. The number of females that parasitized both egg masses was higher in *T. mitsukurii* than in *T. japonicus*; moreover, all the examined females parasitized at least one egg mass (Fig. 2, Table 2).

Trissolcus japonicus females did not show any preference among treatments, choosing indifferently between all paired egg masses (Fig. 2, Table 2). Percentages of emerging parasitoids and egg mortality were not significantly different within each pair examined (Table 2, Table S2).

Trissolcus mitsukurii females chose untreated egg masses significantly more times when compared with egg masses treated with water or treated with Dentamet®. Egg masses treated with Dentamet® were chosen significantly less times when compared with egg masses untreated or treated with water (Fig. 3, Table 2). However, the percentages of parasitized eggs were always above 75% (Table 2).

3.3 Evaluation of *P. carbekii* horizontal transmission

Molecular analysis was performed on the two exotic parasitoids emerging from untreated egg masses, to verify if the primary symbiont of *H. halys* may be horizontally transferred from the egg mass surface to the wasp. However, the presence of live *P. carbekii* cells was not found in either analysed

species through Real time PCR, as only standard samples showed successful amplification. All wasp samples tested positive when submitted to insect-targeted qPCR reactions, confirming their suitable DNA quality.

4. Discussion

This work aimed at assessing the potential interference between the application of Dentamet® on *H. halys* egg masses and parasitism by both indigenous and exotic egg parasitoids. Our experimental observations firstly confirmed the absence of direct harmful effects on parasitoid females resulting from their contact with egg masses treated with biocomplex Dentamet®, as expected since this product does not contain any insecticidal molecule. The need for pest management strategies that involve a substantial reduction of insecticide use has been widely demanded to reduce the impact on beneficial insects (Leskey et al. 2020; Ludwick et al. 2020); therefore, finding a novel technique that is conservative for parasitoid wasps can considerably support the implementation of IPM programmes.

In no-choice tests using native parasitoids, the recorded parasitoid emergence percentages were consistent with previous observations (Roversi et al. 2016; Stahl et al. 2018; Moraglio et al. 2021a, b), with only *A. bifasciatus* being significantly affected by treatment with Dentamet®. However, a reduction of unhatched eggs was reported in replicates subjected to Dentamet® application for all native species. Since the percentages of unhatched eggs were higher in egg masses exposed to native parasitoids than in unexposed ones, a considerable contribution of non-reproductive effects is suggested for these species. These effects include host feeding and oviposition damage (i.e., oviposition followed by host killing and failure of wasp emergence) (Abram et al. 2016; Stahl et al. 2019b; Moraglio et al. 2021b). Treatment with Dentamet® may deter from host feeding more than from oviposition; establishing the actual contribution of the mechanisms causing non-reproductive effects under different conditions may deserve further investigations. Overall, no evident negative impact on the fitness of these wasp species was found in short-term, since the progeny number was

not affected (considering all performed replicates). However, our data suggest that a long-term evaluation may be required to estimate if the total longevity and fecundity of native wasp females exposed to treated eggs may be hampered due to limitation of host feeding. On a food web perspective, the outcome of field treatments with Dentamet® will fit into the already complex dynamics that are being created by the introduction of the exotic stink bug host and parasitoids; this intricate web deserves further study.

As expected, no-choice tests with exotic wasps generally showed higher parasitization levels, as *T. japonicus* and *T. mitsukurii* were the sole species able to parasitize all the exposed egg masses. Emergence percentages were similar to those previously reported (Sabbatini-Peverieri et al. 2020), and they were always above 74%. On the other hand, the percentage of unhatched eggs after exposure to these wasp species was generally lower than for native parasitoids. This may be caused by (i) a higher efficiency of adult emergence after oviposition, and (ii) a lower host feeding behaviour (Abram et al. 2014, 2016, 2019; Kaser et al. 2018). *Trissolcus japonicus* most efficiently parasitized egg masses treated with Dentamet® in comparison to the other two types. Several substances (e.g. host cuticular hydrocarbons and plant volatiles) have been shown to attract *T. japonicus* (Akotsen-Mensah et al. 2021; Arif et al. 2021). We cannot exclude that some of them may share components with those included in the odour blend resulting after our treatment, even though the volatile compounds associated with egg masses treated with Dentamet® are currently unknown. The higher wasp emergence percentage in the group treated with Dentamet® was also the major cause for the reduced percentage of unhatched eggs, whereas a little contribution of non-reproductive effects is expected for this species.

In contrast, *T. mitsukurii* showed significantly lower emergence percentages in egg masses treated with Dentamet®, despite the high proportion of parasitized eggs (> 81%) for all treatments. Since the exposure to *H. halys* egg masses in the absence of the host plant was reported to be inefficient in attracting this wasp (Rondoni et al. 2022), the odour blend emitted after treatment with Dentamet® may further impair the wasp recognition of the host eggs, at least in our experimental conditions,

where no plant was offered to the wasp. The elimination of *P. carbekii* may interfere with the attraction exerted by some symbiont-derived volatiles; the interference with attraction for volatiles may be exacerbated by the general non-preference of *T. mitsukurii* for wet eggs, as evidenced by the untreated *vs* treated with water paired choice test. Strikingly, we observed a possible interaction of treatment with non-reproductive effects, as egg masses treated with Dentamet® showed the highest percentages of unhatched eggs, with a statistical significance when compared with the water treated ones. Future work is required to clarify whether the recorded lower attraction by *T. mitsukurii* is associated or not with unsuitability of Dentamet®- treated eggs; however, as these eggs are less attractive, the possible unsuitability does not seem to be of concern. Furthemore, all percentages of unhatched eggs after exposure to *T. mitsukurii* were similar to the natural proportion observed for egg masses that were not exposed to any parasitoid, indicating that the alteration of non-reproductive effects after treatment with Dentamet® is still modest for this wasp.

The results recorded in no-choice experiments using the exotic parasitoids *T. japonicus* and *T. mitsukurii* were confirmed also in paired choice tests. Haye et al. (2020) suggested that a period of less than 12 h is preferable for paired choice tests involving egg parasitoids of Pentatomidae, to limit the number of females using both egg masses for oviposition, which results in hampered interpretation of host preference. Therefore, we provided the females with egg masses for as little as 3 h. Even in this short period, all females of both species were able to parasitize at least one egg mass, and some of them could still parasitize both egg masses. A diverging behaviour was observed for the two wasp species: both species tended to attack only one of the two masses (more than 50% of parasitism events), but only *T. mitsukurii* showed a significant preference, mostly discarding egg mass treated with Dentamet®. After the egg mass was chosen, also the number of parasitized eggs was different for the two species: emergence percentage for *T. japonicus* was not affected by treatments, whereas we recorded a larger offspring on untreated or water treated egg masses for *T. mitsukurii*, consistently with the results of no-choice tests. The mechanisms (e.g. altered volatile

351 composition) causing such differential behaviour in the host selection process in *T. japonicus* and *T.*
352 *mitsukurii* after egg mass exposure to Dentamet® requires further investigations.
353 Besides the diverging outcome of the short-term preference exhibited by the two *Trissolcus* species,
354 we assessed the occurrence of possible long-term effects on the coevolved host-parasitoid interaction
355 occurring between these exotic wasps and their host *H. halys*. Specifically, we assessed the possible
356 occurrence of a natural horizontal transmission of *P. carbekii* from stink bug egg masses (in untreated
357 specimens) to the adult parasitoids, to figure out if a beneficial relation may have established between
358 the symbiont and wasps over time. The symbiont is vertically transmitted through the egg mass
359 surface (Prado et al., 2006), and this route may have exposed egg parasitoids to acquire the bacterium
360 during emergence from the egg. Horizontal transmission of bacterial symbionts from host to
361 parasitoids have been documented in a number of insects (Dicke et al. 2020), and it may be required
362 for wasp long-term performance, resulting in impaired competition of symbiont-deprived parasitoids.
363 However, qPCR screening of cDNA from untreated samples of both exotic egg parasitoids suggested
364 a lack of *P. carbekii* acquisition from the host, allowing us to exclude any detrimental effects due to
365 possible changes in the microbial community of wasps emerged from treated egg masses.
366 Taken together, our results support the combination of biological control, especially when performed
367 by exotic egg parasitoids, and symbiotic control (at least using Dentamet® or other products with a
368 similar mode of action). Such an integrated approach offers a potential control effectiveness close to
369 100% with total avoidance of insecticidal molecules. In fact, the efficiency recorded for *T. japonicus*
370 and *T. mitsukurii* was always above 75%; furthermore, as reported by Gonella et al. (2019), more
371 than 90% of *H. halys* nymphs emerging from egg masses treated with Dentamet® are deprived of *P.*
372 *carbekii* and die before reaching the second instar, highlighting the powerful potential of a combined
373 effect. Nonetheless, further work will be needed to clarify the containment of *H. halys* using
374 symbiotic control under field conditions. Indeed, possible failure in reaching the egg masses through
375 field application of the anti-symbiont formulate may reduce the final pest mortality; similarly,
376 parasitoid populations may be unable to reach all the egg masses. On the other hand, the combined

effect offered by a combined approach may become even synergistic in the field, with special regard to the diverging response observed for *T. japonicus* and *T. mitsukurii* in terms of egg mass selection. In the field, the unavoidable concurrent presence of egg masses reached by foliar treatments and egg masses that escape the sprays, may offer a choice for *T. japonicus* and *T. mitsukurii*. Egg mass selection may be differentially driven by the treatment for the two wasp species, resulting in a broader range of attacked masses. While *T. japonicus* can parasitize egg masses regardless of the presence of Dentamet® on their surface (providing an increment of mortality from treated egg masses), *T. mitsukurii* may seek for egg masses that were not reached by the treatment. It is worth dedicating future work to analyse such a scenario, since it would increase the final suppression effect, also limiting the competition between the two wasp species.

Declarations

Ethical Approval

All applicable international, national, and/or institutional guidelines for the care and use of animals were followed while conducting this research.

Conflict of interest

The authors declare that there are no conflicts of interest.

Author Contributions

BO, STM, EG, AA and LT conceived the ideas and designed the methodology; BO, STM and FT collected the data; BO and STM analysed the data; BO, STM and EG wrote the manuscript; FT, LT and AA critically reviewed the manuscripts; AA and LT acquired funds. All authors contributed critically to the drafts and gave final approval for publication.

Acknowledgements

The authors thank Sara Scovero and Sara Morello for their essential help in insect rearing and support during experimental work.

Funding

Research was partially funded by Regione Piemonte.

Data availability

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request.

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Table 1 Outcomes of no-choice tests of single egg parasitoid female when exposed to different treatments (U, untreated; W, treated with water; D, treated with Dentamet®). Statistical analysis was performed for parasitized egg masses (i.e. egg masses in which at least one parasitoid successfully emerged out of the exposed egg masses). Asterisks indicate significant differences between treatments (within the same parasitoid species) according to binomial GLM (2 degrees of freedom), with $P < 0.050$ (*), $P < 0.005$ (**) or $P < 0.001$ (***). Different letters refer to significantly different values according to sequential Bonferroni test; n.s. = not significant.

Species	Treatment	Parasitized egg masses on the exposed egg masses	Mean no. eggs per egg mass (±SE)	% (mean eggs ±SE per parasitized egg mass)[a]			
				Parasitoid emergence	χ^2; P value	Unhatched eggs	χ^2; P value
Anastatus bifasciatus	U	7/10	22.9 ± 1.4	16.9 ± 6.0 b		34.4 ± 7.5 a	
	W	5/10	21.4 ± 2.1	19.7 ± 5.4 a	105.952; <0.001 (***)	34.4 ± 9.6 a	10.396; 0.006 (*)
	D	5/10	25.0 ± 1.0	14.2 ± 2.4 c		20.9 ± 6.5 b	
Ooencyrtus telenomicida	U	8/10	20.9 ± 2.0	13.7 ± 3.6		66.1 ± 7.1	
	W	7/10	19.8 ± 2.0	17.1 ± 2.7	4.267; 0.118 (n.s.)	59.1 ± 9.1	4.021; 0.134 (n.s.)
	D	7/10	23.4 ± 1.9	8.4 ± 1.2		49.4 ± 10.8	
Trissolcus kozlovi	U	9/10	25.6 ± 1.0	33.2 ± 9.5		47.8 ± 8.3 a	
	W	8/10	24.4 ± 1.4	28.0 ± 8.1	1.687; 0.430 (n.s.)	44.1 ± 4.2 a	13.799; 0.001 (**)
	D	8/10	26.5 ± 0.5	34.4 ± 6.8		32.5 ± 3.8 b	
Trissolcus japonicus	U	10/10	25.7 ± 0.8	83.5 ± 3.9 b		16.5 ± 3.9 a	
	W	10/10	24.3 ± 2.3	74.3 ± 7.6 b	22.258; <0.001 (***)	24.5 ± 6.9 a	21.328; <0.001 (***)
	D	10/10	24.9 ± 1.8	92.0 ± 3.3 a		7.6 ± 3.2 b	
Trissolcus mitsukurii	U	10/10	26.3 ± 1.0	89.5 ± 4.2 a		8.9 ± 3.2 ab	
	W	10/10	22.9 ± 1.5	94.9 ± 1.7 a	22.65; <0.001 (***)	3.9 ± 1.3 b	14.364; 0.001 (**)
	D	10/10	24.4 ± 1.1	81.3 ± 5.3 b		14.5 ± 3.9 a	
Unexposed egg masses	U	-	27.0 ± 1.2	-		12.9 ± 2.6	
	W	-	24.4 ± 1.3	-		15.2 ± 4.1	2.811; 0.245 (n.s.)
	D	-	25.8 ± 0.6	-		18.1 ± 4.3	

Table 2 Outcomes of paired choice tests of single *T. japonicus* and *T. mitsukurii* females when concurrently exposed to two differently treated egg masses of *H. halys* (U, untreated; W, treated with water; D, treated with Dentamet®). Replicates in which both egg masses were parasitized were not included in statistical analysis. Asterisks indicate significant differences between the number of females that chose one or the other egg mass according to the Pearson Chi-square test, with $P < 0.050$ (*) or $P < 0.005$ (**); n.s. = not significant according to the Pearson Chi-square test (No. of females parasitizing) or binomial GLM (parasitoid emergence and egg mortality rates).

Species	Treatment	Mean no. eggs per egg mass (± SE)	No. of parasitized egg masses	No. of females parasitizing					% (mean eggs ±SE per parasitized egg mass)[a]	
				Both egg masses	Only untreated control (%)	Only treated with water (%)	Only Dentamet® (%)	P value[a]	Emerged parasitoids (sig)[b]	Unhatched eggs (sig)[b]
Trissolcus japonicus	U	20.7 ± 2.2	7	2	5 (71.4)			0.317 (n.s.)	85.7 ± 9.3 (n.s.)	14.3 ± 9.3 (n.s.)
	W	23.1 ± 2.1	5			3 (60.0)			88.7 ± 9.6 (n.s.)	11.3 ± 9.6 (n.s.)
	W	23.9 ± 1.7	6	2		4 (66.7)		1.000 (n.s.)	57.4 ± 18.4 (n.s.)	24.1 ± 7.6 (n.s.)
	D	24.4 ± 1.6	6				4 (66.7)		50.4 ± 14.8 (n.s.)	24.2 ± 10.3 (n.s.)
	U	24.5 ± 1.2	5	0	5 (100.0)			1.000 (n.s.)	89.6 ± 2.0 (n.s.)	9.6 ± 1.5 (n.s.)
	D	26.3 ± 1.0	5				5 (100.0)		84.6 ± 15.4 (n.s.)	15.4 ± 15.4 (n.s.)
Trissolcus mitsukurii	U	27.4 ± 0.8	8	2	6 (75.0)			0.003 (**)	94.2 ± 2.3	4.7 ± 1.7
	W	25.4 ± 0.9	3			1 (33.3)			96.3 ± 0.0	3.7 ± 0.0
	W	20.8 ± 2.3	10	5		5 (50.0)		0.002 (**)	80.0 ± 8.8	15.4 ± 6.9
	D	20.2 ± 2.3	5				0		-	-
	U	25.0 ± 0.9	9	4	5 (55.5)			0.021 (*)	91.7 ± 5.4	5.0 ± 3.0
	D	26.4 ± 0.9	5				1 (20.0)		75.0 ± 0.0	21.4 ± 0.0

[a]Replicates with both parasitized egg masses were not included

[b]*Trissolcus mitsukurii* was not included in GLM tests since it parasitized both egg masses in most cases

649 **Table S1** Results of statistical analyses applied in no-choice tests with native parasitoids, referring to
650 all replicates. Binomial GLM was used (2 degrees of freedom). Results referred to *T. japonicus* and
651 *T. mitsukurii* are the same as indicated in Table 1.

Species	**Measured value**	**Test result**	**P value**
A. bifasciatus	% of parasitoids emerged	χ^2=2.063	0.356
A. bifasciatus	% of unhatched eggs	χ^2=39.363	<0.001
O. telenomicida	% of parasitoids emerged	χ^2=3.250	0.197
O. telenomicida	% of unhatched eggs	χ^2=4.021	<0.001
T. kozlovi	% of parasitoids emerged	χ^2=0.798	0.671
T. kozlovi	% of unhatched eggs	χ^2=13.977	0.001

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654 **Table S2** Results of statistical analyses applied to emergence rate and egg mortality rate in paired
655 choice tests using *T. japonicus*. Binomial GLM was used (1 degree of freedom); replicates with both
656 egg masses parasitized were not included.

Treatments pair	**Measured value**	**Test result**	**P value**
Untreated *vs* Treated with water	% of parasitoids emerged	χ^2=0.472	0.492
Untreated *vs* Treated with water	% of unhatched eggs	χ^2=0.472	0.492
Treated with water *vs* Dentamet®	% of parasitoids emerged	χ^2=0.132	0.716
Treated with water *vs* Dentamet®	% of unhatched eggs	χ^2=0.413	0.520
Untreated *vs* Dentamet®	% of parasitoids emerged	χ^2=1.369	0.242
Untreated *vs* Dentamet®	% of unhatched eggs	χ^2=1.008	0.315

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Figures captions

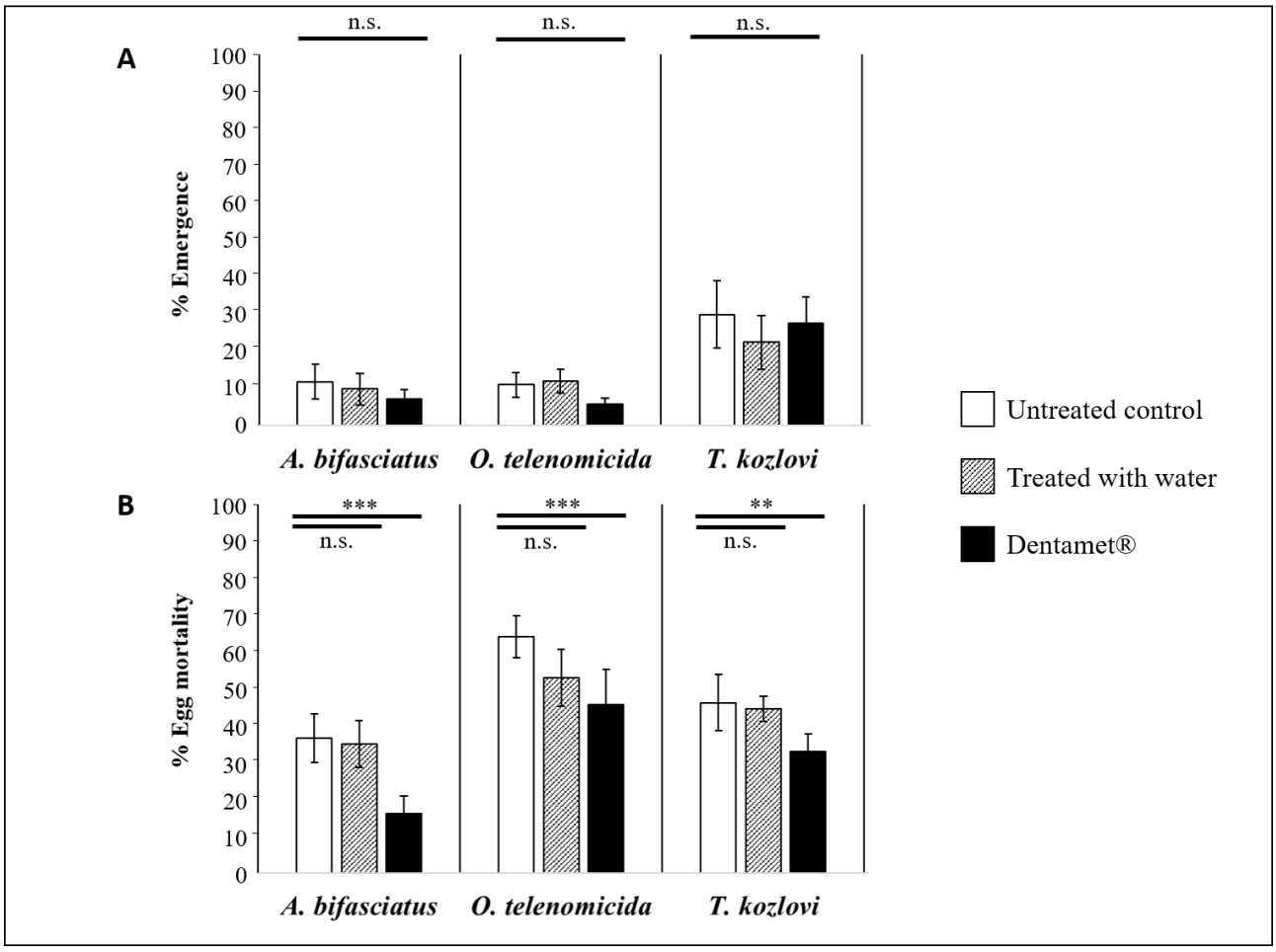


Fig. 1 Results of no-choice tests with native parasitoid species, considering all replicates. Mean of emergence rates (A), and mean of egg mortality rates (B) recorded for *H. halys* egg masses exposed to different egg parasitoids and treatments. Bars indicate standard errors; asterisks indicate significant differences between treatments according to binomial GLM, with $P < 0.005$ (**) or $P < 0.001$ (***); n.s. = not significant.

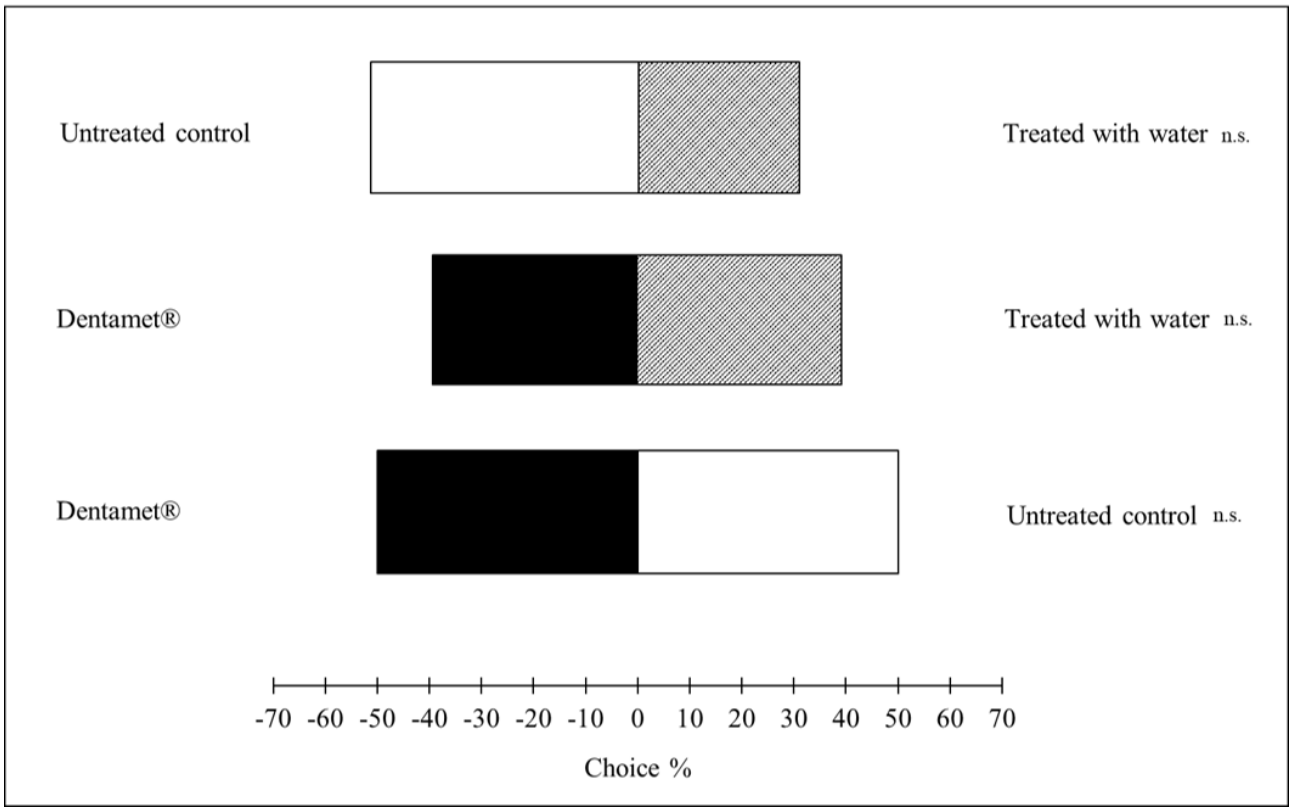
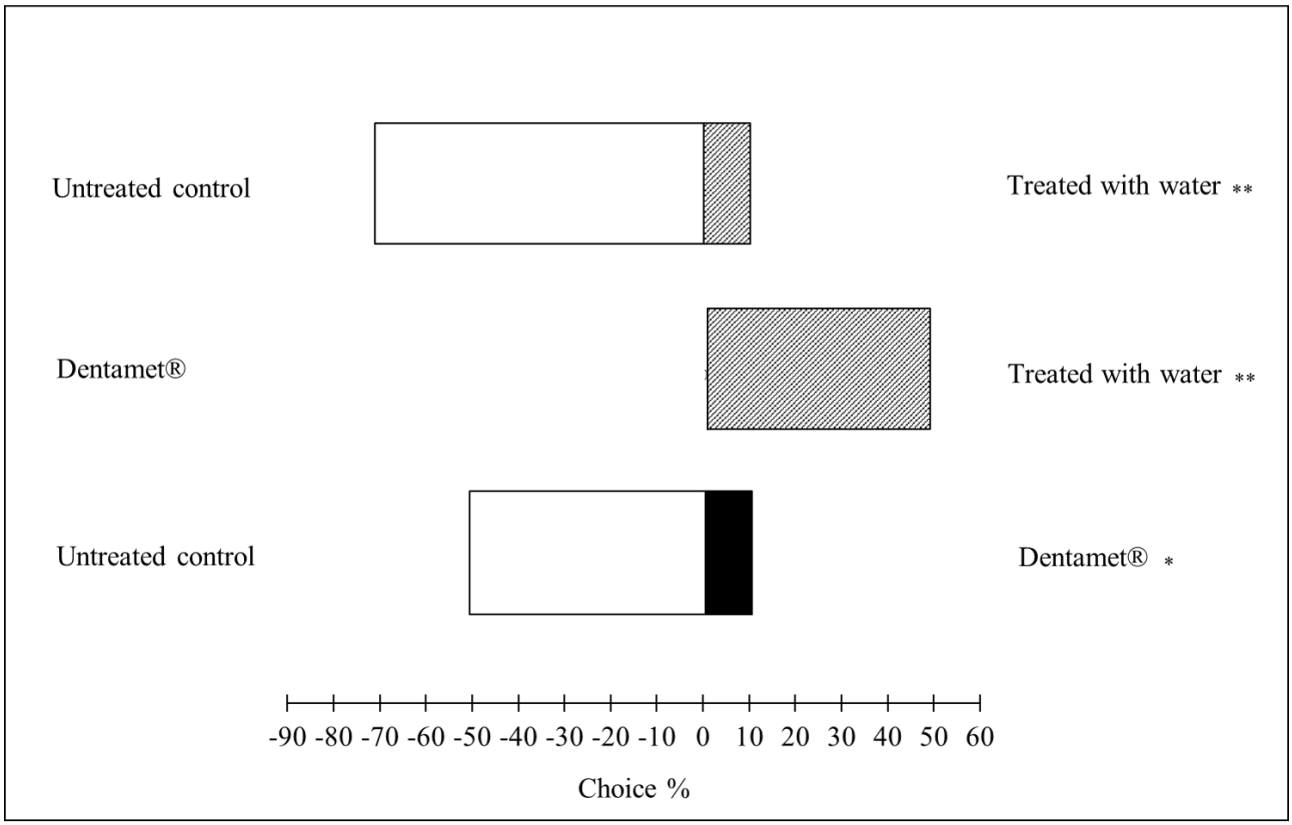


Fig. 2 Results of paired choice tests performed with *T. japonicus* females exposed to egg masses of the following groups: untreated (white boxes), treated with water (grey boxes), treated with Dentamet® (black boxes). n.s.: not significant.



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671 **Fig. 3** Results of paired choice tests performed with *T. mitsukurii* females exposed to egg masses of
672 the following groups: untreated (white boxes), treated with water (grey boxes), treated with
673 Dentamet® (black boxes). Asterisks indicate significant differences between the number of egg
674 parasitoids that chose one or the other egg mass according to the Pearson's Chi-square test, with $P < 0.050$ (*) or $P < 0.005$ (**); n.s.: not significant.

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