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The dual effects of root-cap exudates on nematodes: from quiescence in plant-parasitic nematodes to frenzy in entomopathogenic nematodes

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Abstract

To defend themselves against herbivores and pathogens, plants produce numerous secondary metabolites, either constitutively or de novo in response to attacks. An intriguing constitutive example is the exudate produced by certain root-cap cells that can induce a state of reversible quiescence in plant-parasitic nematodes, thereby providing protection against these antagonists. The effect of such root exudates on beneficial entomopathogenic nematodes (EPNs) remains unclear, but could potentially impair their use in pest management programmes. We therefore tested how the exudates secreted by green pea (Pisum sativum) root caps affect four commercial EPN species. The exudates induced reversible quiescence in all EPN species tested. Quiescence levels varied with the green pea cultivars tested. Notably, after storage in root exudate, EPN performance traits were maintained over time, whereas performances of EPNs stored in water rapidly declined. In sharp contrast to high concentrations, lower concentrations of the exudate resulted in a significant increase in EPN activity and infectiousness, but still reduced the activity of two plant-parasitic nematode species. Our study suggests a finely tuned dual bioactivity of the exudate from green pea root caps. Appropriately formulated, it can favour long-term storage of EPNs and boost their infectiousness, while it may also be used to protect plants from plant-parasitic nematodes.

Key words: Below-ground food web, biological control, entomopathogenic nematode (EPN), Heterodera glycines, Heterorhabditis bacteriophora, Meloidogyne incognita, plant defence, plant-parasitic nematode, root-cap exudate.

Introduction

The estimated production of over 200,000 different compounds (Barber and Martin, 1976) ranks plant’s secondary metabolites among the most diverse functional chemicals in the biosphere. Thousands of papers have been devoted to the description of ecological and evolutionary processes mediated by plant secondary metabolites (Dudareva et al., 2013). Stahl (1888) was one of the first to suggest that these metabolites are involved in plant defences and they have since been proposed to contribute as an important selecting force in the interaction between plants and herbivores (Ehrlich and Raven, 1964). Historically, work on chemical plant defences has focused on foliar parts of plants (Hartmann, 2007) and roots have been mostly ignored (Hunter, 2001). Yet up to 20% of the photosynthetic fixed carbon is exuded via below-ground tissues as secondary metabolites (Barber and Martin, 1976) and these exudates shape rhizospheric interactions (e.g. Hiltpold and Turlings, 2012; Rasmann et al., 2012a and references therein; Erb et al., 2013).

In the context of plant and plant-parasitic nematode interactions, root exudates are often beneficial to the plant antagonists, as they may use plant secondary metabolites to locate host plants (Prot, 1980; Rolfe et al., 2000; Curtis et al., 2009;...
Reynolds et al., 2011). In addition to attracting nematodes, root exudates also trigger egg hatching in certain plant-parasitic nematode species (e.g. Perry and Clarke, 1981; Denijis and Lock, 1992; Perry and Gaur, 1996; Gaur et al., 2000; Wesemael et al., 2006; Pudasaini et al., 2008; Khokon et al., 2009; Oka and Mizukubo, 2009). Yet root exudates can also protect roots against plant-parasitic nematodes. For instance, metabolites exuded from the root-cap cells of legumes and maize (Zea mays) slow down movement in plant-parasitic nematodes, sometimes resulting in a state of quiescence, reducing the ability of the nematodes to infect the plant (Zhao et al., 2000). This apparent defence mechanism has been found in several plant species and is usually reversible (Hubbard et al., 2005).

In addition to plant-parasitic nematodes, root exudates have also been reported to affect entomopathogenic nematodes (EPNs), plant protagonists in the rhizosphere. Rasmann et al. (2005) found that insect herbivory on maize roots induces the emission of volatile secondary metabolites that can attract the EPN Heterorhabditis megidis Poinar, Jackson and Klein (Rhabditida: Heterorhabditidae), Steinernema feltiae Filipjev (Rhabditida: Steinernematidae), and S. carpocapsae Weiser (Rhabditida: Steinernematidae), following the methodology by Hubbard et al. (2005), H. bacteriophora, S. feltiae, and S. carpocapsae were provided by Landi-Reba AG (Switzerland), and H. megidis by Becker Underwood (UK).

Briefly, ~1000 active infective juveniles of each EPN species were taken from batches freshly hatched from Galleria mellonella L. (Lepidoptera: Galleridae). EPNs were suspended in water and aliquots of 30 infective juveniles in 50 μl of water were pipetted into 36 wells of a 96-well plate (BD Biosciences, CA, USA) and 175 μl of root cap exudate was added to each EPN suspension. After 12 h, using a dissecting microscope, the number of quiescent EPN-infective juveniles was evaluated by direct counting of individuals exhibiting, or not exhibiting, an active sinusoidal form and movement. Instead of root-cap exudate, 175 μl of water was pipetted into the wells of the control plates. The experiment was repeated 10 times for each EPN species.

To evaluate EPN recovery from exudate-induced quiescence, 125 μl was pipetted out of the wells, leaving settled EPN in the bottom, and replaced with the same volume of water. After 12 h, resumption of EPN activity was evaluated using a dissecting microscope. To test if recovered EPNs were still infectious, five times 10 individuals per species per replicate were sampled and applied on top of G. mellonella larvae that had been individually placed in wells of a 24-well plate. The baits were then covered with 10% moist sand (white sand, Migros, Switzerland) and stored at 25°C in the dark. EPN infection was assessed after 48 h.

**Materials and methods**

**Plant material**

*Pisum sativum* L. cv. Lancet seeds (Wyss Samen und Planzen AG, Switzerland) were sterilized in 95% ethanol for 5 min. Seeds were then rinsed and immersed in pasteurized distilled water for 12 h. Imbibed seeds were germinated for 3 days at 25°C in the dark in plastic boxes (15×13.5×5 cm) on a 1.5 cm layer of 1.0% PhytoAgar (Duchefa Biochemie, The Netherlands).

**Root-cap exudate collection**

Exudates were collected from 15 pea germinates by arranging them on a Teflon plate at the periphery of a 1 ml drop of ultra-pure water, with only the tips (~5 mm) submerged. After 2 min of immersion, the liquid was gently agitated in order to disperse the border cells and the associated exudate. The solution obtained was then centrifuged at 4000 g for 10 min in a 1.5 ml centrifuge tube (Vaudaux-Eppendorf AG, Switzerland), resulting in a pellet of the suspended border cells. The supernatant, defined as root-cap exudate, was collected. Following this technique, exudates from several hundred of pea germinates were collected, and pooled in one solution. To facilitate the experimentation, this last solution was split into 7 ml amber vials (Supelco, Sigma-Aldrich Chemie GmbH, Switzerland) before being stored at -20°C.

**Exudate-induced quiescence and recovery of EPNs**

The pea root-cap exudate was tested on the following EPN species: *Heterorhabditis bacteriophora* Poinar (Rhabditida: Heterorhabditidae), *Heterorhabditis megidis* Poinar, Jackson and Klein (Rhabditida: Heterorhabditidae), *Steinernema feltiae* Filipjev (Rhabditida: Steinernematidae), and *S. carpocapsae* Weiser (Rhabditida: Steinernematidae), following the methodology by Hubbard et al. (2005), *H. bacteriophora*, *S. feltiae*, and *S. carpocapsae* were provided by Landi-Reba AG (Switzerland), and *H. megidis* by Becker Underwood (UK).

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**Quiescence induction by the root exudates of various green-pea cultivars**

Following the procedures described above, eight additional cultivars (cv. Kelvin, cv. Kelvin Sprinter, cv. Merveilles précoce, and cv. Sprint, Wyss Samen und Planzen AG, Switzerland; cv. Exquis, cv. Surab, cv. Picolo, and cv. Carmini, Landi-Reba AG, Switzerland) were tested for their ability to induce quiescence in *H. bacteriophora*. Water was used as a control and the EPNs were kept in the respective solutions for 12 h before the number of quiescent nematodes was recorded. Each test was repeated 10 times.

**Effect of root-cap exudate on the quality of EPNs over time**

Either 41.25 ml of root-cap exudate from green pea cv. Lancet or the same amount of water (control) were poured into 50 ml culture flasks (BD Biosciences, CA, USA), and 7.5 ml of water containing 20 000 *H. bacteriophora* infective juveniles were added. The flasks were then stored at room temperature (25°C) in the dark. Every third day, starting 12 h after the induction of quiescence, the infectiousness, mobility, and lipid content of the nematodes from both solutions were assessed for 18 days.
**EPN infectiousness**  EPN infectiousness was measured using the five-on-one sand well bioassay (modified after Grewal, 2005). In a 24-well tissue plate (BD Biosciences, CA, USA), 10 G. mellonella larvae were individually placed in wells and covered with 10% moist sand (white sand, Migros, Switzerland). Five infective juveniles were pipetted from either root-cap exudate or water solution and added to each well. 24-well plates were stored at 25°C for 3 days and G. mellonella larvae were dissected. The number of infected insect larvae was recorded.  

**EPN mobility**  EPN mobility was measured using a method modified after Tomalak, 2005. Into a 10-cm diameter Petri dish we pipetted a 10-μl drop of 0.1% caffeine (99% pure, Alfa Aesar GmbH, Germany) on top of the 10 ml of 2% solidified agar. About 100 EPNs from either root-cap exudate or water formulations were slowly pipetted onto a 5-mm diameter filter paper (cut out of bigger filter papers; Schleicher & Schuell GmbH, Germany). To facilitate removal of excessive water, the filter paper was laid on a paper towel. The filter was transferred immediately onto the drop of caffeine, with the EPNs on the upper side of the disc. The Petri dish was covered and stored in the dark at room temperature for 30 min. As caffeine strongly repels EPNs, motile EPNs would quickly crawl away from this region onto the agar plate. After incubation, the disc was rinsed in a 1.5-ml centrifuge tube (Vaudaux-Eppendorf AG, Switzerland) containing water and the total number of sessile nematodes, recovered from the disc, was counted and the percentage of mobile EPNs calculated. This experiment was repeated five times per treatment.  

**Estimation of the neutral lipid content**  Estimation of the neutral lipid content was performed using a method modified after Patel et al., 1997). About 50 infective juveniles were sampled from each formulation described above. They were poured into a glass vial (7 ml amber glass vial; Suppelco, Sigma-Aldrich Chemie GmbH, Switzerland) and flooded with 70% ethanol saturated with Oil Red O (Sigma-Aldrich Chemie GmbH, Switzerland). The glass vials were incubated at 60°C for 20 min. The excess of staining solution was removed before the addition of 5 ml of glycerol:water (50:50 v/v) solution. EPNs were left to settle overnight at room temperature. After placing the EPNs on a slide, the lipid content of the stained nematodes was estimated by comparison with a neutral lipid index scale [from 8 (fully stained) to 1 (no staining); Patel et al., 1997].  

**Heat and cold stability of the green-pea exudate**  The stability of green-pea exudate exposed to a temperature gradient was assessed. Aliquots of the exudate were either frozen (–80°C and –20°C) for 12 h or heated up (20, 30, 40, 60, 80, and 100°C) for 10 min. Controls consisted of water only, subjected to the same temperature treatments. For all treatments, induction of EPN quiescence was tested as described above and repeated seven times with different individuals from three different batches of nematodes.  

**Dilution of root cap exudate and the effect on bioactivity**  Following the procedure described in the previous sections, different concentrations of pea root-cap exudate were tested. Exudate from green pea cv. Lancet was collected and freeze dried. The resulting material was diluted in 1 ml (original concentration), 1.25, 1.5, 1.75, 2, 3, and 4 ml of water. Quiescence induction in H. bacteriophora was tested for each concentration; controls consisted of water only. The number of quiescent nematodes was recorded after 12 h of induction. Each test was repeated 15 times.  

Increasing activity of EPNs was observed with the reduction of the exudate concentration. Therefore, we assessed EPN activity more precisely by counting the number of oscillations per minute for individual H. bacteriophora after 12 h exposure to a low concentration of pea root-cap exudate (in 1.5 ml of water). Head oscillations of EPNs exposed to the original concentration of the exudate (in 1 ml of water) and to water only were also recorded. This test was repeated with 10 different individuals from three different batches of nematodes.  

Following the five-on-one methodology (see above and Grewal, 2005), the infectiousness of highly active EPNs in low concentration green pea exudate was compared to those in water only. In 24-well plates, 12 G. mellonella larvae were covered with 10% moist sand previously mixed with either 1.5× diluted green-pea exudate or water (9.1, v/v). Five infective juveniles of the EPN H. bacteriophora were individually transferred in each well. Plates were wrapped in Parafilm and incubated at 25°C in the dark. The number of EPN-infected larvae was recorded every 12 h for 96 h, starting 12 h after EPN inoculation. This experiment was replicated four times with three different batches of EPNs.  

The effect of the exudate concentrations was also tested on plant-parasitic nematodes. Following the procedure described above, the head oscillation counts per minute of the soybean cyst nematode *Heterodera glycine* Ichinohe (Tylenchida: Heteroderinae) and the root-knot nematode *Meloidogyne incognita* Kofoid and White (Tylenchida: Heteroderinae) were recorded in 1.5×-diluted exudate and in water.  

**Statistical analyses**  Statistical tests were conducted in SigmaPlot 12.1.0.15 (Systat Software GmbH Germany).  

**Quiescence induction by the root exudates of various green pea cultivars**  To determine differences in quiescence induced by the different cultivars we analysed the data using a one-way ANOVA procedure. Statistical differences within groups were evaluated using a Tukey post-hoc test.  

**Effect of root-cap exudate on the quality of EPNs over time**  EPN infectiousness, mobility, and neutral lipid content were tested using an RM two-way ANOVA procedure with treatment and time as factors. Statistical differences within groups were evaluated using a Tukey post-hoc test.  

**Heat and cold stability of the green pea exudate**  Differences in induced quiescence after temperature treatments were measured using a one-way ANOVA on ranks procedure. Statistical differences within groups were evaluated using a Tukey post-hoc test.  

**Dilution of root-cap exudate and the effect on bioactivity**  The effect of dilution on the induced quiescence to EPNs was tested using an one-way ANOVA on ranks procedure. Statistical differences within groups were evaluated using a Tukey post-hoc test.  

**Differences between H. glycine and M. incognita**  Head oscillations per minute recorded in water or in both concentrations of exudate were tested with a one-way ANOVA comparing the EPN head oscillations when exposed to both concentrations of green pea exudate or water. Infectiousness of EPNs exposed to either water or low-concentration exudate was tested using an RM two-way ANOVA procedure with treatment and time as factors. Statistical differences within groups were evaluated using a Tukey post-hoc test.  

**Results**  

**Exudate-induced quiescence and recovery of EPNs**  

Each EPN species that experienced contact with the green pea root exudate exhibited a state of quiescence (Table 1). Recovery was complete for *S. carpopcapsae*, *H. megidis*, and *H. bacteriophora*. A large majority of *S. feltiae* larvae was recovered from quiescence. Each EPN species was still infectious after recovering from induced quiescence (Table 1).  

**Quiescence induction by the exudates of various green-pea cultivars**  

The induction of quiescence in *H. bacteriophora* was highly dependent on the green-pea genotypes (Fig. 1, ANOVA, $F_{5,99} = 26.686, P ≤ 0.001$). Exudate from cv. Kelvin Sprinter had the lowest induction
rate of quiescence (~25%), whereas the exudate of some cultivars resulted in almost 100% quiescence (i.e. cv. Picolo) (Fig. 1).

**Effect of root-cap exudate on the quality of EPNs over time**

Induced quiescence in *H. bacteriophora* helped to maintain its performance over time (Fig. 2A, B, C). Overall, infectiousness of *H. bacteriophora* that had been stored in exudate was higher than for EPNs that had been stored in water (Fig. 2A; two-way RM ANOVA, $F_{1,119} = 12.05$, $P < 0.001$). After recovering from quiescence, EPNs that had been stored in exudate were significantly more mobile than EPNs that had been stored in water (Fig. 2A; two-way RM ANOVA, $F_{1,119} = 66.839$, $P < 0.001$) and this remained significantly higher over the 18 days of the experiment (Fig. 2B; two-way RM ANOVA, $F_{6,119} = 3.845$, $P = 0.003$). Lipid reserves in quiescent nematodes were significantly higher than in EPNs stored in water (Fig. 2C; two-way RM ANOVA, $F_{1,119} = 53.129$, $P < 0.001$) and remained so over the course of the experiment (Fig. 2C; two-way RM ANOVA, $F_{6,119} = 2.414$, $P = 0.03$).

**Heat and cold stability of the green-pea exudate**

Exposing the exudate to different temperatures did not impair its bioactivity, and quiescence induction levels were similar to those measured in the control exudate. After heating or freezing, the exudate still induced quiescence in exposed *H. bacteriophora*, which was not the case for the water control [Supplementary Figure S1 (heat: ANOVA on Recovery (%), $H = 45.419$, $P < 0.001$); Supplementary Figure S2 (cold: ANOVA on ranks, $H = 33.289$, $P < 0.001$)].

**Dilution of root-cap exudate and the effect on bioactivity**

Reducing the exudate concentration quickly reached a threshold beyond which induction of quiescence in *H. bacteriophora* failed (Fig. 3; ANOVA on ranks, $H_{1,15} = 52.42$, $P < 0.001$), but instead resulted in increased movement (Fig. 4; ANOVA, $F_{2,89} = 491.427$, $P < 0.001$). At a low concentration, the exudate had a positive impact on the infectiousness of *H. bacteriophora* (Fig. 5; RM two-way ANOVA, $F_{1,107} = 20.991$, $P = 0.006$), and EPN exposed to low concentration exudates were more effective at killing *G. mellonella* over time as compared to EPN exposed to water only (Fig. 5; RM two-way ANOVA, $F_{8,107} = 5.88$, $P < 0.001$). Contrary to the effect on EPNs, the same low concentration of exudate significantly reduced the mobility of the tested plant-parasitic nematode species *H. glycine* (Fig. 4B; ANOVA, $F_{2,89} = 225.023$, $P < 0.001$) and *M. incognita* (Fig. 4C; ANOVA on ranks, $H = 67.876$, $P < 0.001$).

**Discussion**

Earlier observations had revealed that green pea root-cap exudate causes a loss of mobility and induces a state of quiescence in several plant-parasitic nematode species (Zhao et al., 2000; Hubbard et al., 2005), as well as in the bacteria-feeding nematode *Cenorhabditis elegans* Maupas (Zhao et al., 2000; Hubbard et al., 2005); and the EPN *S. feltiae* (Hubbard et al., 2005). The present study demonstrates that the activity of other EPN species also drops dramatically when exposed to the green pea root-cap exudate (Table 1). Quiescence is usually a

| Table 1. Exudate-induced quiescence, recovery, and subsequent infectiousness of three EPN species |
|-------------------------------------------------|-----------------|-----------------|--------------|
| Species                          | Quiescence (%) | Recovery (%)   | Infectiousness |
| Steinernema feltia              | 91 ± 6          | 57 ± 12        | + a           |
| S. carpocapsae                 | 93 ± 4          | 100            | +            |
| Heterorhabditis megalis       | 89 ± 9          | 97 ± 2         | +            |
| H. bacteriophora               | 95 ± 2          | 100            | +            |

aSome individuals remained quiescent after removal of the exudate. Nonetheless, larvae treated with recovered *S. feltiae* were infected.

![Figure 1](https://academic.oup.com/jxb/article-abstract/66/2/603/2893249) Fig. 1. Induced quiescence in *Heterorhabditis bacteriophora* varies among green pea cultivars. Induced EPN quiescence was significantly influenced by the cultivar the root-cap exudates were collected from (grey bars). Three cultivars induced quiescence close or equal to 100% whereas some barely reached 30%, but were still different from the percentage of inactive nematodes recorded in water (white bar). One cultivar (Kelvin Sprinter) was not different from water. Letters indicate statistical differences. Bars represent SEM.
reversible response in nematodes to toxic or unfavourable environmental conditions (Evans and Perry, 1976). As plant-parasitic nematodes, as well as other pathogens, preferably choose the elongation zone right behind the root tip to penetrate root tissues (Prot, 1980; Curl and Truelove, 1986; Gunawardena and Hawes, 2002), the induction of quiescence in root pathogens around this susceptible region has been interpreted as a defensive mechanism (Hawes et al., 2000; Zhao et al., 2000; Wuyts et al., 2006). Recent findings showed that plant histone-linked extracellular DNA (exDNA) might be involved in this putative root tip defence (Wen et al., 2009; Hawes et al., 2011; Hawes et al., 2012). Functioning as an extracellular trap attracting and immobilizing pathogens (Hawes et al., 2012), exDNA has been shown to be a key component in plant resistance to infection by pathogenic fungi (Wen et al., 2009). By reducing or inhibiting the motility of these pathogens, roots can grow away from the threat, avoiding damage to their apical tips, while offering less vulnerable regions to herbivores. Interestingly, this is in contradiction to the common dogma that plants cannot escape a threat by moving away. Indeed, in addition to immobilizing pathogens (Hawes et al., 2000), roots can potentially grow through a cumulative total of 6 m

Fig. 2. Induced quiescence enhances over time storage of the EPN Heterorhabditis bacteriophora. Quality traits were better conserved over time when EPNs were stored in a solution with root-cap exudates from green pea cv. Lancet (grey circle) as compared to EPNs stored in water (white circle). (A) EPN infectiousness remained significantly higher in root-cap exudates as compared to water during most of the experiment. (B) When stored in root-cap exudates, EPNs stayed mobile for longer compared to the EPNs in water over most of the duration of the experiment. (C) Because EPNs stored in root-cap exudate were quiescent, their lipid reserves were maintained at a significantly higher level over time compared to lipid reserves measured in EPNs stored in water. (D–F) After being stained with Oil Red O, lipids appear in red inside the nematode body. A visual assessment allows an evaluation of the lipid content according to the 1–8 lipid index (Patel et al., 1997). According to this index, (D) fully stained EPNs were given an 8, (E) intermediately stained EPNs received a 4, and (F) individuals without staining were recorded as 1. Level of significance between treatments on one day are indicated as not significant (ns), $P < 0.05$ (*), $P \leq 0.01$ (**), $P \leq 0.001$ (**). Statistical differences within treatments over time are noted with letters. Bars indicate SEM.
of soil within a day (Lynch and van Beem, 1993). This provides the roots with an escape strategy that is comparable to animals fleeing unfavourable biotic and abiotic conditions. To determine whether this strategy may indeed be effective against the relatively large and mobile nematodes requires additional experiments. The present study shows that the exudates involved in this potential defence mechanism not only affects the roots’ ‘foes’ but can also momentarily impair beneficial microorganisms such as EPNs (Table 1).

As with other plant defence traits, the quiescence potency of root exudates varies a lot among plant families (Hubbard et al., 2005), but we also found dissimilarities among genotypes of a particular species (Fig. 1). The results presented here provide a new example of how plant breeding can affect the concentrations of secondary metabolites that are important for plant performance (e.g. Hubbard et al., 2005; Rasmann et al., 2005; Köllner et al., 2008; Hiltpold et al., 2010c; Erb et al., 2011; Robert et al., 2012; Meihls et al., 2013).
Induced quiescence in EPNs may have interesting applications in biological control. Indeed, EPN mass production has been optimized (Ehlers, 2001), but downstream processes such as storage and transport can still be problematic. Inducing quiescence in EPNs with the root exudate significantly maintained EPN performance traits such as infectiousness, mobility, and lipid content (Fig. 2A–C). As in insect parasitoids (Casas et al., 2005; Denis et al., 2013), EPNs rely on their lipid reserves while foraging and infecting insect hosts (Patel et al., 1997). Loss of lipids was significantly lower in quiescent *H. bacteriophora* than in EPNs stored in water (Fig. 2C), possibly explaining their higher mobility and infectiousness after recovery (Fig. 2A and B). Similar to root penetration by plant-parasitic nematodes (Hubbard et al., 2005), infectiousness and mobility of EPNs recovering from quiescence tended to increase (Fig. 2A and B), suggesting a general increase of nematode activity after quiescence recovery. Whereas the timescale used in this study is not pertinent for long-term storage, these data clearly imply that the addition of the active compounds to the storage formulation will increase the EPN shelf life.

Induced quiescence in EPNs can be problematic when aiming at controlling insect pests in the rhizosphere. However, this apparent biological conflict might not occur in natural environments. Indeed, at lower concentrations of the exudate, as can be expected in the field, EPN activity was significantly boosted, while plant-parasitic nematodes remained quiescent (Fig. 4). Further tests under natural conditions must be performed, but the enhanced infectiousness of ‘frenetic’ EPNs (Fig. 5) suggests a dual effect of the tested exudate: root defences can simultaneously subdue ‘foes’, and invigorate ‘friends’.

The present study reinforces the idea that plant exudates can help shape the dynamics of interactions not only in the rhizosphere, but also at more distant regions in the soil (Rasmann et al., 2012b; Erb et al., 2013; Hiltpold et al., 2013). The strategy of plants to protect their root tips with exudates is in apparent conflict with an alternative strategy, the attraction of natural enemies upon herbivorous attack. Yet, because exudate bioactivity seems tuned to have opposite effects on plant mutualists and antagonists, the two strategies might effectively complement each other. The complexity of biotic and abiotic interactions in the rhizosphere is huge and difficult to fully understand, but each new finding leads to a better and broader understanding of this very particular ecosystem. Therefore the prospect of harnessing root exudation and exploiting it for enhanced pest control (Hiltpold and Turlings, 2012) may indeed be realistic (Degenhardt et al., 2009; Hiltpold et al., 2010a; b; Ali et al., 2012; Hiltpold et al., 2012).

**Supplementary material**

Supplementary data can be found at *JXB* online.

**Supplementary Figure S1.** Effect of heat on the bioactivity of the pea root-cap exudate and the induction of quiescence in *H. bacteriophora*.

**Supplementary Figure S2.** Induction of quiescence in *H. bacteriophora* after the exudate was frozen.

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