Response of the Pacific Coast wireworm, Limonius canus, and the dusky wireworm, Agriotes obscurus (Coleoptera: Elateridae), to insecticide-treated wheat seeds in a soil bioassay

Réponse du taupin du Pacifique, Limonius canus, et du taupin obscur, Agriotes obscurus (Coleoptera : Elateridae), à des semences de blé traitées à l’insecticide dans un test biologique dans le sol

Willem G. van Herk, Robert S. Vernon, Chandra Moffat et Chantelle Harding

Des larves du taupin obscur, Agriotes obscurus, et du taupin du Pacifique, Limonius canus, ont été exposées à des semences de blé germées et traitées à l’insecticide dans un dispositif-fenêtre rempli de terre en 2005 et en 2006. La position des larves ainsi que leur comportement envers les semences (contact ou répulsion) ont été observés toutes les 5 min durant 3 ou 5 h. La santé des larves a été observée durant 70 ou 126 j après l’exposition. Les semences ont été traitées avec les fongicides Dividend XLRTA (difénoconazole, méfénoxame) ou Raxil MD (tébuconazole, métalaxyle), ou encore avec les insecticides Vitavax Dual (lindane), Poncho 600F (clothianidine), Cruiser 350FS (thiaméthoxame), Admire 240FS (imidaclopride), Gaucho 480FL (imidaclopride), Tefluthrin 20CS (tefluthrine), ou une combinaison Tefluthrin-Cruiser. La plupart des vers fil-de-fer (> 80 %) sont entrés en contact avec les semences dans tous les traitements. Ils sont généralement demeurés en contact avec les semences durant toute la période d’observation dans les traitements témoins (Dividend, Raxil, semences non traitées). Les larves ont été repoussées après un bref contact (< 20 min) dans tous les traitements de Tefluthrin, sauf si elles étaient moribondes. La majorité des vers fil-de-fer se sont remis de la morbidité induite par le contact avec les semences à l’intérieur de 21 h et n’ont pas fait de rechute, à l’exception des larves de L. canus exposées au Cruiser et de quelques larves de A. obscurus exposées aux insecticides Gaucho et Admire. Le taux de mortalité était bas (< 50 %) dans tous les traitements sauf pour les larves de L. canus exposées au Cruiser à 15 et 30 g.m.a. 100 kg-1 graine (60 et 75 %, respectivement). Le taux de mortalité était significativement plus bas lorsque les larves de L. canus étaient exposées à des combinaisons Tefluthrin-Cruiser que lorsqu’elles étaient exposées seulement au Cruiser. Ces résultats suggèrent que pour évaluer l’efficacité des insecticides à lutter contre les populations de vers fil-de-fer, une observation directe de leur comportement ainsi qu’une évaluation à long terme de leur état de santé sont nécessaires. L’impact de la répulsion et de la morbidité causées par les insecticides sur la lutte aux vers fil-de-fer dans les champs est également abordé.
Larvae of the dusky wireworm, Agriotes obscurus, and the Pacific Coast wireworm, Limonius canus, were exposed to germinated, insecticide-treated wheat seeds in a soil-filled bioassay in 2005 and 2006. Position in the bioassay and contact and/or repellence behaviour towards the seeds were recorded every 5 min for 3 or 5 h. Wireworm health was recorded for 70 or 126 d after exposure. Seeds were treated with the fungicides Dividend XLRTA (difenoconazole, mefenoxam) or Raxil MD (tebuconazole, metalaxyl), and/or the insecticides Vitavax Dual (lindane), Poncho 600F (clothianidine), Cruiser 350FS (thiamethoxam), Admire 240FS (imidacloprid), Gaucho 480FL (imidacloprid), Tefluthrin 20CS (tefluthrin), or Tefluthrin-Cruiser combinations. Most wireworms (> 80%) came into contact with the seeds in all treatments. Wireworms generally remained in contact throughout the observation period in the control treatments (Dividend, Raxil, untreated seeds). Unless moribund, wireworms were repelled after brief (< 20 min) contact in all Tefluthrin treatments. Most wireworms recovered from contact-induced morbidity within 21 d and did not relapse, except L. canus exposed to Cruiser and some A. obscurus exposed to Gauch and Admire. Wireworm mortality was low (< 50%) in all treatments except L. canus exposed to Cruiser at 15 and 30 g a.i. 100 kg⁻¹ seed (60 and 75%, respectively). Mortality was significantly less important when L. canus larvae were exposed to Tefluthrin-Cruiser combinations than when exposed to Cruiser alone. We suggest that efficacy assessments of insecticides for wireworm control require direct observation of their behaviour and long-term post-exposure health assessments, and discuss the impact of repellence and/or morbidity elicited by insecticides on wireworm control in the field.

Keywords: Agriotes obscurus, insecticide, Limonius canus, repellence, seed treatments, wireworm.
INTRODUCTION

Wireworms are important pests of wheat, vegetable and soft fruits in both North America and Europe (Vernon 2005; Parker and Howard 2001), and are likely to become an even more serious problem in some countries as effective organochlorine (OC), organophosphate (OP) and carbamate insecticides are being removed without the promise of suitable replacements (Grove et al. 2000; Vernon et al. 2001). The search for lower-risk replacement chemicals has so far focused on pyrethroid (i.e. tefluthrin, bifenthrin), chloronicotinoid (i.e. imidacloprid, acetamiprid), thianicotinoid (i.e. clothianidin, thiamethoxam) and phenyl pyrazole (i.e. fipronil) insecticides (Parker and Howard 2001), some of which have demonstrated their effectiveness against wireworms and are being used for wireworm management (Kuhar et al. 2003; Parker and Howard 2001). Recent reports suggest that imidacloprid and thiamethoxam are effective against Agriotes sp. and Melanotus sp. in corn and sugarbeet (Andersch and Schwarz 2003; Maienfisch et al. 2001; Pons and Albajes 2002), and that clothianidin is more effective against Melanotus sp. in corn than either chlorpyrifos or tefluthrin (Andersch and Schwarz 2003). However, these assessments are based on plant establishment and/or yield and do not consider the direct effects of these chemicals on wireworm health and behaviour.

The direct effects of insecticides on wireworm health and behaviour are of interest and importance for several reasons. Recent toxicology work has shown that wireworms can recover from long-term (> 150 d) morbidity induced by dermal exposure to imidacloprid, clothianidin and thiamethoxam (van Herk et al. 2008a; Vernon et al. 2008). In addition, certain insecticides are suspected to have repellent as well as toxic effects on wireworms, including aldrin and lindane (Long and Lilly 1958; Toba et al. 1988), terbufos (Belcher and Tenne 1987), and chlorpyrifos and fonofos (Horne and Horne 1991; Missionnier and Brunel 1979). Recent work by van Herk et al. (2008b) showed that the dusky wireworm, Agriotes obscurus L., was repelled by droplets of tefluthrin, chlorpyrifos, lindane and imidaclorpid in a soil-less bioassay.

To determine if some of the neonicotinoid and synthetic pyrethroid insecticides listed above elicit repellence in wireworms when incorporated into soil, van Herk and Vernon (2007a) developed a bioassay that makes it possible to observe wireworm orientation, contact and repellence behaviours in response to insecticide-treated wheat seeds in soil. Repellence, generally defined as movement away from a stimulus (Dethier et al. 1960), is here said to occur when wireworms retreat > 5 cm from the seeds within 30 min of contact (van Herk and Vernon 2007a).

Preliminary studies using this assay indicated that larvae of both A. obscurus and the Pacific Coast wireworm, Limonius canus LeConte, were repelled by Tefluthrin-treated wheat seeds in soil (van Herk and Vernon 2007a). However, this report did not include wireworm response to seeds treated with commercial formulations of thiamethoxam, clothianidin and imidacloprid, or the effect(s) of repeated contact with insecticide-treated seeds, i.e. whether it elicited a change in contact and foraging behaviour or resulted in aversion learning. These questions are of interest as some neonicotinoid insecticides may elicit repellence in wireworms (as discussed above), and because behavioural resistance to insecticides resulting from aversion learning is a common and important resistance mechanism (Bernays and Chapman 1987; Lockwood et al. 1984; Sparks et al. 1989).

Three observation studies were conducted using larvae of A. obscurus (in 2005) and L. canus (in 2005 and 2006) to determine wireworm response to seeds treated with thiamethoxam, clothianidin and imidacloprid, to determine whether exposure to both tefluthrin and thiamethoxam resulted in greater mortality than exposure to seeds treated with either chemical singly (as suggested by preliminary work by van Herk and Vernon; unpublished), and to determine if repeated contact with tefluthrin-treated seeds elicited a behavioural change. In the 2005 studies, both species were exposed to wheat seeds treated with lindane, tefluthrin or clothianidin, and A. obscurus larvae were also exposed to seeds treated with imidacloprid, thiamethoxam, and thiamethoxam and tefluthrin combined. In the 2006 study, L. canus larvae were exposed to seeds treated with lindane and different concentrations of tefluthrin, thiamethoxam, and tefluthrin plus thiamethoxam. In this paper we describe the effects of these insecticide treatments on wireworm behaviour, morbidity and mortality.

MATERIALS AND METHODS

Wireworms

Agriotes obscurus larvae used in the 2005 study were collected by hand-sifting soil taken from pastureland in 2005 at the Pacific Agri-Food Research Centre (PARC) in Agassiz, BC. Collected larvae represented all instars, but were considered as belonging to one population since they were collected within 100 m of each other. Limonius canus larvae used in the 2005 study were collected using flour baits placed in a fallowed field at an organic farm in Kelowna, BC, in June 2005, while L. canus larvae used in the 2006 study were collected similarly from the same location in June 2006. Virtually all collected larvae were late instar (i.e. > 15 mm long) and were considered as belonging to one population since they were collected within 50 m of each other. All wireworms were stored at 4°C in 40 L Rubbermaid (Rubbermaid, Atlanta, GA) tubs filled with sandy-clay loam soil.
collected at PARC, until needed. To reduce variability, only late-instar feeding wireworms were used in the bioassays. Feeding wireworms were obtained by placing potato slices in the storage tubs. Wireworms assembled at the potato baits were then isolated in soil without food following methods described by van Herk and Vernon (2007a), and were used in bioassays within 3 d.

Bioassay
Wireworms were exposed to germinated wheat seeds in soil-filled circular bioassays as described by van Herk and Vernon (2007a). Using circular bioassays ensures that wireworms moving along the edge of the arena (a common pre-orienting behaviour) remain within 13 cm from the seeds and do not become trapped in corners. Bioassay arenas consisted of three separate 30 cm x 30 cm sections of transparent, 4 mm thick Plexiglas® connected by small carriage bolts. A 25 cm diam hole machined into the centre section created a circular chamber that could be filled with soil to a depth of 4 mm. A transparent plastic grid overlaying both the top and bottom sections divided the chamber into 113 equally-sized cells that were grouped into eight concentric rings and four quadrants. Rings and cells were numbered from the centre outwards, with ring 1 consisting of a single cell (cell 1) touching all four quadrants; other cells were restricted to individual quadrants. Five wheat seeds (cv. AC Superb) were placed in cell 1 after the bioassay chamber had carefully been filled with an even 4 mm layer of screened, sandy-clay loam soil. Soil used was adjusted to contain 20% moisture by weight. Wheat seeds were pre-germinated for 44-48 h at 25 ± 1°C on moist paper towels, ensuring approximately 15 mm shoot length prior to bioassays. Seedlings were placed in ring 1; seedling shoots extended partly into ring 2 (cells 2-5).

After the seeds were placed, the top section of the bioassay was put into place and fastened. Arenas were positioned horizontally on a raised wooden frame to make observations possible through both the top and bottom sections. Seedlings were allowed to grow in the soil in the assembled bioassay for 30 min to establish CO2 gradients (Doane et al. 1975), after which wireworms (one per arena) were introduced head first into the bioassay chamber through a 5 mm hole in the top centre of either cell 89, 96, 103 or 110 (the centre cells of ring 8 in quadrants I, II, III and IV, respectively). The wireworm introduction hole was sealed with pressure-sensitive tape (VWR International Ltd., Delta, BC) throughout the seed incubation and wireworm observation periods, and opened only for wireworm insertion (< 1 min).

Wireworm position in the bioassay, contact and/or repellence behaviour towards the seed, and health (if abnormal) were recorded every 5 min for 3 h in 2005 and for 5 h in 2006. Wireworms were considered to have come into contact with the seeds when they were within rings 1 or 2 of the bioassay (van Herk and Vernon 2007a). The duration of contact was estimated by multiplying the number of observed contact events by 5 min, i.e. the interval between observations. For the 2006 study, wireworms were observed long enough to assess two or more contact periods and to measure the duration between these contact periods (hereafter referred to as ‘inter-contact’ periods). All observations were conducted at room temperature (21 ± 1°C) under low intensity red light (0.75 µE s-1 m-2, measured with a Li-188B integrating quantum radiometer/photometer; Li-Cor, Lincoln, NB).

Post-exposure wireworm health was assessed immediately after bioassays using methods and criteria developed by Vernon et al. (2008). Wireworms that could move out of an 8 cm diam circle drawn on 12.5 cm filter paper in Petri dish arenas within 2 min were designated as ‘alive’. Wireworms that were incapable of directed movement but were capable of body movements obvious to the naked eye were designated as ‘writhing’. Wireworms that made no visible body movements with or without prodding were inspected under a dissecting microscope to determine if they exhibited leg and/or mouthpart movements. These wireworms were designated as ‘appendage movement’. Wireworms exhibiting no spontaneous or elicited writhing or leg/mouthpart movements were temporarily classified as ‘probably dead’, but were not recorded as ‘dead’ until they showed signs of decomposition (Vernon et al. 2008). Wireworms were stored individually into 150 mL plastic containers (Fisher Scientific, Whitby, ON) with screened soil (as described above) for 70 d after exposure (DAE) in the 2005 studies, and for 126 DAE in 2006. Wireworm health was assessed 1 and 7 DAE, and weekly thereafter. For the 2006 study, wireworm health was also assessed 3 DAE. Mortality was compared among treatments at 56 DAE in the 2005 studies and at 70 DAE in the 2006 study, after which time no further mortality was observed.

Insecticide treatments
Wheat seeds used in the 2005 studies were treated with Vitavax Dual (containing 50 g lindane and 54 g carbathiin) at 124 g a.i. 100 kg-1 seed, Gaucho 480FL (imidacloprid) at 15 g a.i. 100 kg-1 seed, Admire 240FS (thiamethoxam) at 10 and 30 g a.i. 100 kg-1 seed, Tefluthrin 20CS (tefluthrin) at 10 g a.i. 100 kg-1 seed, Cruiser 350FS (clothianidin) at 25 g a.i. 100 kg-1 seed, Tefluthrin 20CS (both at 10 g a.i. 100 kg-1 seed) and a combination of Cruiser 350FS and Tefluthrin 20CS (both at 10 g a.i. 100 kg-1 seed) (Table 1). Wheat seeds treated with Poncho and Gaucho were treated by Gustafson Inc. (now Bayer CropScience Canada, Toronto, ON), and were also treated with the fungicide Raxil MD (1.5 g a.i. tebuconazole and 2.0 g a.i. metalaxyl 100 kg-1 seed). Seeds treated with Vitavax Dual, Tefluthrin and/or Cruiser were treated by Syngenta Crop Protection Canada Inc. (Guelph, ON) and, except for Vitavax Dual, were also treated with the fungicide Dividend XLRTA (containing 3.21% difenoconazole and 0.27% mfenoxam) at 13 g a.i. 100 kg-1 seed. These insecticide/fungicide combinations reflect treatments under evaluation for wireworm control by Bayer and Syngenta. Admire 240FS was applied to untreated seeds by the authors. In addition, untreated wheat seeds and seeds treated with Raxil MD and Dividend XLRTA alone were tested as control treatments. Due to limited wireworm availability, not all treatments were tested on both wireworm species in 2005 (Table 1).
Wheat seeds used in the 2006 study were treated by Syngenta Crop Protection Canada Inc. with Vitavax Dual at 124 g a.i. 100 kg⁻¹ seed, Tefluthrin 20CS, Cruiser 350FS, or both Tefluthrin 20CS and Cruiser 350FS. Tefluthrin 20CS and Cruiser 350FS were tested individually at 5, 10, 15, 20 and 30 g a.i. 100 kg⁻¹ seed to determine if the concentration of either chemical affected wireworm behaviour. The combined Tefluthrin 20CS and Cruiser 350FS treatments were tested at 5, 10, 15, 20 and 20 g a.i. 100 kg⁻¹ seed of each insecticide to determine if using both chemicals on the seeds had an enhanced effect on wireworm behaviour and health. Seeds treated with Tefluthrin and/or Cruiser were also treated with the fungicide Dividend XLRTA at 13 g a.i. 100 kg⁻¹ seed.

Between 20 and 30 Agriotes obscurus larvae were exposed to each treatment in 2005. Due to a shortage of L. canus in 2005, only 10 wireworms were exposed to the Rayil and Dividend treatments. Depending on the availability of feeding wireworms, 20 to 60 L. canus larvae were exposed to each treatment in the 2006 study. Each treatment was conducted over several weeks, with several, randomly-chosen treatments assayed per day. Ten to twenty wireworms were observed concurrently on each observation day. As wireworms have long larval periods (approximately 6 mo per instar), can be maintained in storage for extensive periods (> 2 yr), and were handled similarly in all bioassays (i.e. similar storage conditions, selection and handling, bioassay preparation, observation methods), observation date was not considered to have an impact on wireworm behaviour.

These insecticides were chosen for study as they are currently being evaluated for wireworm management (see above) and as some (e.g. Poncho, Gauch) appear to provide stand protection in wheat without reducing wireworm populations (R.S. Vernon, unpublished data). Of these insecticides, clothianidin (Poncho), thiamethoxam (Cruiser) and tefluthrin (Force 3.0G) are currently registered in Canada for wireworm management in corn; Vitavax Dual was also included as it has historically been used for wireworm management and is considered repellent to wireworms (Long and Lilly 1958; Toba et al. 1988).

### Table 1. Contact duration (min) of A. obscurus and L. canus larvae exposed for 180 min to wheat seeds treated with pesticides, proportion of moribund larvae at the end of the bioassay, and proportion of dead larvae in 2005

<table>
<thead>
<tr>
<th>Treatment¹</th>
<th>C (N)²</th>
<th>Mean (SEM) duration of first contact (min)</th>
<th>Mean (SEM) duration of total contact (min)</th>
<th>Proportion (of C) moribund after bioassays</th>
<th>Proportion (of C) dead at 56 DAE³</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Agriotes obscurus</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Untreated 26 (29)</td>
<td>122.7 (11.4) a</td>
<td>128.1 (10.3) a</td>
<td>0 a</td>
<td>0 a</td>
<td></td>
</tr>
<tr>
<td>Dividend 19 (20)</td>
<td>112.4 (15.0) ab</td>
<td>127.6 (10.8) a</td>
<td>0 a</td>
<td>0 a</td>
<td></td>
</tr>
<tr>
<td>Rayil 19 (22)</td>
<td>90.0 (14.8) abc</td>
<td>105.3 (12.9) ab</td>
<td>0 a</td>
<td>0 a</td>
<td></td>
</tr>
<tr>
<td>Cruiser (10) 20 (20)</td>
<td>104.5 (9.9) ab</td>
<td>124.8 (9.0) a</td>
<td>0.90 cd</td>
<td>0.15 ab</td>
<td></td>
</tr>
<tr>
<td>Cruiser (30) 20 (20)</td>
<td>64.3 (9.9) abc</td>
<td>88.0 (8.9) abc</td>
<td>0.40 bc</td>
<td>0 a</td>
<td></td>
</tr>
<tr>
<td>Tefluthrin (10) 20 (24)</td>
<td>15.5 (1.9) d</td>
<td>48.0 (7.8) cd</td>
<td>0.45 bc</td>
<td>0.05 a</td>
<td></td>
</tr>
<tr>
<td>Cruiser (10) + Tefluthrin (10) 20 (25)</td>
<td>13.8 (2.1) d</td>
<td>34.5 (5.6) d</td>
<td>0.35 b</td>
<td>0.25 ab</td>
<td></td>
</tr>
<tr>
<td>Gauch (15) 18 (20)</td>
<td>79.4 (16.2) abc</td>
<td>100.6 (13.0) ab</td>
<td>0.83 bcd</td>
<td>0.22 ab</td>
<td></td>
</tr>
<tr>
<td>Admire (10) 20 (22)</td>
<td>71.4 (13.3) abc</td>
<td>79.5 (12.5) abcd</td>
<td>0.75 bcd</td>
<td>0.45 b</td>
<td></td>
</tr>
<tr>
<td>Admire (30) 18 (19)</td>
<td>71.1 (14.3) abc</td>
<td>74.7 (13.8) bcd</td>
<td>0.72 bcd</td>
<td>0.39 b</td>
<td></td>
</tr>
<tr>
<td>Poncho (25) 18 (20)</td>
<td>50.3 (10.3) cd</td>
<td>72.8 (9.2) bcd</td>
<td>0.72 bcd</td>
<td>0.11 ab</td>
<td></td>
</tr>
<tr>
<td>Vitavax (124) 20 (20)</td>
<td>113.8 (13.0) ab</td>
<td>115.3 (12.4) ab</td>
<td>1.0 d</td>
<td>0.10 ab</td>
<td></td>
</tr>
<tr>
<td><strong>Limonius canus</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Untreated 17 (20)</td>
<td>117.4 (16.8) a</td>
<td>136.2 (13.4) a</td>
<td>0.0 a</td>
<td>0 a</td>
<td></td>
</tr>
<tr>
<td>Dividend 10 (10)</td>
<td>83.0 (19.7) ab</td>
<td>124.5 (11.3) ab</td>
<td>0.0 a</td>
<td>0 a</td>
<td></td>
</tr>
<tr>
<td>Rayil 10 (10)</td>
<td>111.5 (20.0) a</td>
<td>135.5 (14.8) a</td>
<td>0.0 a</td>
<td>0 a</td>
<td></td>
</tr>
<tr>
<td>Tefluthrin (10) 20 (20)</td>
<td>21.3 (3.0) c</td>
<td>36.8 (3.9) c</td>
<td>0.4 b</td>
<td>0 a</td>
<td></td>
</tr>
<tr>
<td>Poncho (25) 17 (19)</td>
<td>45.9 (7.6) bc</td>
<td>63.5 (10.0) bc</td>
<td>0.53 b</td>
<td>0.11 a</td>
<td></td>
</tr>
<tr>
<td>Vitavax (124) 19 (20)</td>
<td>56.1 (14.4) bc</td>
<td>91.8 (12.8) ab</td>
<td>0.68 b</td>
<td>0.15 a</td>
<td></td>
</tr>
</tbody>
</table>

¹ Dividend = Dividend XLRTA, Rayil = Rayil MD, Cruiser = Cruiser 350FS, Tefluthrin = Tefluthrin 20CS, Gauch = Gauch 480FL, Admire = Admire 240FS, Poncho = Poncho 600F, Vitavax = Vitavax Dual. Numbers in parentheses indicate grams a.i. 100 kg⁻¹ seed.
² C = number of wireworms that came into contact with the seeds; N = number of wireworms that were exposed.
³ Values followed by the same letter within a column are not significantly different at α = 0.05. Analyses were conducted separately per species.
⁴ DAE = days after exposure.
RESULTS

Duration of first contact

For all three studies (A. obscurus in 2005, L. canus in 2005, and L. canus in 2006), a high percentage (> 80%) of wireworms introduced into bioassay arenas came into contact with the seeds during the observation period (Tables 1 and 2), suggesting that insecticide treatments did not prevent wireworm movement towards, or contact with, the seeds. However, in each study, first contact duration differed significantly among treatments (F_{11,227} = 9.77, P < 0.0001, Table 1; F_{5,87} = 8.61, P < 0.0001, Table 1; F_{6,476} = 22.64, P < 0.0001, Table 2), as the insecticides affected normal wireworm behaviour. In the 2006 L. canus study, poorer seed germination in untreated seeds and, to a lesser extent, in the Cruiser 20 g a.i. treatment also affected first contact duration. Seedlings in these two treatments had shorter roots (approx. 5-10 mm, instead of 15 mm in the other treatments), likely because seeds were mistakenly given less moisture during pre-germination (Cruiser 20 g) or were not treated with a fungicide (untreated seeds). All seeds used in the 2006 study came from the same lot and were treated at the same time, and the poorer germination seen in the above treatments does not suggest poorer germination in the field, though the absence of fungicide in the untreated seeds may cause a slight delay in emergence. However, poorer seed germination will affect seedling CO2 production, which may affect wireworm contact with the seeds since CO2 stimulates feeding in wireworms (Doane et al. 1975).

Control treatments

Agriotes obscurus larvae exposed to control treatments (untreated seeds or seeds treated with Dividend XLRTA or Raxil MD) remained in contact in an apparently healthy state for 122.7 (SEM = 11.4), 112.4 (15.0) and 90.0 (14.8) min, respectively, with an apparently healthy state for 122.7 (SEM = 11.4), 112.4 (15.0) and 90.0 (14.8) min, respectively, with seedlings in these two treatments had shorter roots (approx. 5-10 mm, instead of 15 mm in the other treatments), likely because seeds were mistakenly given less moisture during pre-germination (Cruiser 20 g) or were not treated with a fungicide (untreated seeds). All seeds used in the 2006 study came from the same lot and were treated at the same time, and the poorer germination seen in the above treatments does not suggest poorer germination in the field, though the absence of fungicide in the untreated seeds may cause a slight delay in emergence. However, poorer seed germination will affect seedling CO2 production, which may affect wireworm contact with the seeds since CO2 stimulates feeding in wireworms (Doane et al. 1975).

Table 2. Contact duration (min) of L. canus larvae exposed for 300 min to wheat seeds treated with insecticides, proportion of moribund larvae at the end of the bioassay, and proportion of dead larvae after 70 DAE

<table>
<thead>
<tr>
<th>Treatment1</th>
<th>C1 (N)2</th>
<th>Mean (SEM) duration of first contact (min)</th>
<th>Mean (SEM) duration of second contact (min)</th>
<th>Mean (SEM) duration of total contact (min)</th>
<th>Proportion (of C1) moribund after bioassays</th>
<th>Proportion (of C1) dead at 70 DAE3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Untreated</td>
<td>33 (40)</td>
<td>118.0 (19.1) bc1</td>
<td>70.3 (16.4) bc1</td>
<td>183.6 (14.8) ab1</td>
<td>0 a1</td>
<td>0.06 ab1</td>
</tr>
<tr>
<td>Dividend</td>
<td>40 (40)</td>
<td>182.4 (18.0) a</td>
<td>126.8 (28.2) a</td>
<td>232.1 (11.2) a</td>
<td>0 a</td>
<td>0.05 ab</td>
</tr>
<tr>
<td>Cruiser (5)</td>
<td>20 (20)</td>
<td>171.3 (21.1) ab</td>
<td>75.9 (24.3) b</td>
<td>230.0 (11.8) a</td>
<td>0 a</td>
<td>0.25 bc</td>
</tr>
<tr>
<td>Cruiser (10)</td>
<td>37 (40)</td>
<td>141.2 (13.8) abc</td>
<td>62.1 (16.7) bcd</td>
<td>176.6 (12.8) ab</td>
<td>0.05 ab</td>
<td>0.35 cd</td>
</tr>
<tr>
<td>Cruiser (15)</td>
<td>20 (20)</td>
<td>156.5 (17.2) abc</td>
<td>37.7 (13.1) bcd</td>
<td>203.5 (15.8) ab</td>
<td>0.35 bcd</td>
<td>0.60 de</td>
</tr>
<tr>
<td>Cruiser (20)</td>
<td>33 (40)</td>
<td>106.5 (14.5) c</td>
<td>63.0 (12.0) bcd</td>
<td>165.8 (14.6) b</td>
<td>n/a</td>
<td>n/a</td>
</tr>
<tr>
<td>Cruiser (30)</td>
<td>20 (20)</td>
<td>98.3 (13.4) c</td>
<td>54.7 (12.5) bcd</td>
<td>157.0 (15.4) b</td>
<td>0.35 bcd</td>
<td>0.75 e</td>
</tr>
<tr>
<td>Tefluthrin (5)</td>
<td>54 (60)</td>
<td>25.3 (2.5) d</td>
<td>25.0 (4.3) bcd</td>
<td>62.5 (6.6) c</td>
<td>0.19 abc</td>
<td>0.04 a</td>
</tr>
<tr>
<td>Tefluthrin (10)</td>
<td>53 (59)</td>
<td>22.5 (1.6) d</td>
<td>20.5 (2.8) cd</td>
<td>61.9 (5.8) c</td>
<td>0.19 abc</td>
<td>0.08 ab</td>
</tr>
<tr>
<td>Tefluthrin (15)</td>
<td>18 (18)</td>
<td>27.2 (4.5) d</td>
<td>31.4 (11.0) bcd</td>
<td>82.8 (14.6) c</td>
<td>0.28 abc</td>
<td>0 a</td>
</tr>
<tr>
<td>Tefluthrin (20)</td>
<td>17 (19)</td>
<td>21.8 (6.5) d</td>
<td>15.8 (2.9) d</td>
<td>39.1 (5.9) c</td>
<td>0.41 cd</td>
<td>0 a</td>
</tr>
<tr>
<td>Tefluthrin (30)</td>
<td>20 (20)</td>
<td>20.9 (2.7) d</td>
<td>20.5 (9.3) cd</td>
<td>64.5 (9.4) c</td>
<td>0.20 abc</td>
<td>0.15 abc</td>
</tr>
<tr>
<td>Cruiser (5) + Tefluthrin (5)</td>
<td>20 (20)</td>
<td>27.5 (3.9) d</td>
<td>22.4 (4.6) cd</td>
<td>73.3 (9.1) c</td>
<td>0.30 abc</td>
<td>0.05 ab</td>
</tr>
<tr>
<td>Cruiser (10) + Tefluthrin (10)</td>
<td>20 (20)</td>
<td>23.5 (3.5) d</td>
<td>18.0 (2.0) d</td>
<td>68.3 (10.4) c</td>
<td>0.15 abc</td>
<td>0.05 ab</td>
</tr>
<tr>
<td>Cruiser (15) + Tefluthrin (15)</td>
<td>20 (20)</td>
<td>27.5 (4.1) d</td>
<td>17.8 (3.1) d</td>
<td>64.8 (5.5) c</td>
<td>0.30 abc</td>
<td>0 a</td>
</tr>
<tr>
<td>Cruiser (20) + Tefluthrin (20)</td>
<td>17 (19)</td>
<td>31.5 (4.0) d</td>
<td>15.7 (3.4) d</td>
<td>59.1 (6.6) c</td>
<td>0 a</td>
<td>0 a</td>
</tr>
<tr>
<td>Vitavax (124)</td>
<td>53 (60)</td>
<td>98.6 (12.3) c</td>
<td>46.9 (6.9) bcd</td>
<td>147.5 (11.0) b</td>
<td>0.74 d</td>
<td>0 a</td>
</tr>
</tbody>
</table>

1 Dividend = Dividend XLRTA, Cruiser = Cruiser 350FS, Tefluthrin = Tefluthrin 20CS, Vitavax = Vitavax Dual. Numbers in parentheses indicate grams a.i. 100 kg^-1 seed.
2 C1 = number of wireworms that came into contact with the seeds at least once; N = number of wireworms exposed.
3 Values followed by the same letter within a column are not significantly different at α = 0.05.
4 C2 = number of wireworms that came into contact with the seeds twice.
5 DAE = days after exposure.
most wireworms still in contact at the end of the observation period (Table 1). Similarly, larvae of *L. canus* exposed to control treatments in 2005 remained in contact on average for 80 min or more, with most larvae still in contact at the end of the observation period (Table 1). In the 2005 studies, the initial contact duration of *A. obscurus* and *L. canus* exposed to untreated seeds and to seeds treated with either Dividend or Raxil was similar (*t* = 0.31, *P* = 0.76; *t* = 0.83, *P* = 0.41; *t* = 0.82, *P* = 0.42, respectively). First contact duration of *L. canus* with Dividend-treated seeds was longer in 2006 than in 2005 as bioassays were continued longer in 2006 than in 2005 and as normal wireworm feeding behaviour is to remain in contact with a suitable host for extensive periods (van Herk and Vernon 2007a). However, due to the poor seed germination in untreated seeds in the 2006 study (as discussed above), the first contact duration of *L. canus* exposed to untreated seeds was similar in the 2005 and 2006 studies.

**Tefluthrin 20SC**
The first contact duration of *A. obscurus* was shortest in the Tefluthrin treatments (with and without Cruiser), with wireworms moving away from seeds before showing signs of morbidity, often after < 10 min contact (Table 1). Similarly, in 2005, *L. canus* remained in contact with Tefluthrin-treated seeds for a significantly shorter period than with seeds in the control treatments (Table 1). In the 2006 study, the first contact duration of *L. canus* was briefest in all Tefluthrin (with or without Cruiser) treatments, with no significant difference (*P* > 0.05) in contact duration among the treatments containing Tefluthrin (Table 2). First contact duration of *A. obscurus* and *L. canus* in the 2005 studies, and of *L. canus* exposed to seeds treated with Tefluthrin at 10 g a.i. in 2005 and 2006, did not differ significantly (*t* = 0.78, *P* = 0.33; *t* = 0.34, *P* = 0.74, respectively).

**Cruiser 350FS**
The first contact duration of *Agriotes obscurus* larvae exposed to Cruiser at 10 g a.i. (104.5 min, SEM = 9.9) was the same as that of larvae exposed to control seeds (Table 1), partly because a high percentage (50%) of wireworms became moribund and therefore stopped moving. Contact duration of *Agriotes obscurus* larvae exposed to Cruiser at 30 g a.i. was more brief (64.3 min, SEM = 9.9) than that of larvae exposed to Cruiser at 10 g a.i., and fewer larvae (10%) became moribund in situ. Exposure to seeds treated with both Tefluthrin and Cruiser resulted in significantly shorter contact duration than exposure to Cruiser alone (Table 1).

In the 2006 *L. canus* study, there was no significant difference (*P* > 0.05) between first contact duration of larvae exposed to seeds treated with Cruiser at 5, 10 and 15 g a.i., and control seeds (Table 2). As with *A. obscurus*, the first contact duration of *L. canus* larvae exposed to high rates of Cruiser (20-30 g a.i.) was significantly shorter than when the larvae were exposed to Dividend XLRTA (Table 2). In addition, contact duration of *L. canus* with seeds treated with Cruiser alone was significantly longer than with seeds treated with the same rates of both Cruiser and Tefluthrin (Table 2), indicating that the presence of Tefluthrin decreased the duration of contact in the combined treatments.

**Vitavax Dual**
The first contact duration of *Agriotes obscurus* larvae exposed to Vitavax Dual (113.8 min, SEM = 13.0) was the same as that of larvae exposed to control seeds (Table 1), partly because most wireworms (90%) became moribund in situ. In contrast, first contact duration of *L. canus* larvae exposed to Vitavax Dual in 2005 was shorter than that of *A. obscurus* exposed to the same treatment (*t* = 2.98, *P* = 0.005), as fewer *L. canus* larvae (47%) became moribund in situ. In the 2006 study, the first contact duration of *L. canus* was significantly shorter in the Vitavax Dual than in the Dividend treatments, as some wireworms (30%) moved away from the seeds shortly before becoming moribund. While first contact in the Vitavax Dual treatment was significantly longer in 2006 than in 2005 (*t* = 2.23, *P* = 0.03), this was due to the immobility of moribund wireworms and the longer observation periods in 2006 than in 2005.

**Other treatments**
Most *A. obscurus* that came into contact with seeds treated with Gaucho 480FL and Admire 240FS (at both 10 and 30 g a.i.) for the first time remained in contact for periods numerically but not significantly shorter than with control seeds (Table 1), and some wireworms became moribund in situ (33, 53 and 37%, respectively).

Contact duration of *A. obscurus* larvae exposed to Poncho 600F 25 g a.i. seeds was more brief (50.3 min, SEM = 10.3) than that of larvae exposed to control seeds (Table 1), and a small percentage (25%) of wireworms became moribund in situ. First contact duration of *A. obscurus* and *L. canus* (45.9 min, SEM = 7.6) exposed to Poncho treatments was similar (*t* = 0.40, *P* = 0.69) (Table 1).

**Duration of subsequent contacts**
Some wireworms that had moved away from the seeds after the first contact subsequently re-contacted the seeds. In the 2005 studies, most *A. obscurus* (95%) and *L. canus* (80%) came into contact with seeds treated with Tefluthrin (alone) more than once. However, there was no significant difference between the duration of the first and second contacts (*A. obscurus*: 15.0 (SEM = 2.0), 15.3 (2.5) min, respectively; *L. canus*: 20.3 (3.5), 13.8 (3.5) min, respectively; *t* = 1.21, *P* = 0.24). Similarly, most (85%) *A. obscurus* re-contacted seeds treated with the Tefluthrin and Cruiser combination treatment, but there was no significant difference between the duration of the first and second contacts (14.7 (2.3), 15.6 (2.4) min, respectively; *t* = 0.31, *P* = 0.76). Not enough repeated contacts were observed in the other treatments to permit analysis (data not shown).

The longer observation periods in the 2006 study made it possible to observe repeated contacts with seeds in all treatments (Table 2). Duration of the second contact differed significantly among treatments (*F*16,329 = 6.91, *P* < 0.0001), with contact with Dividend-treated seeds being significantly longer than with all other treatments, including untreated seeds (Table 2). As with the first contact, duration of
the second contact was shortest in Tefluthrin treatments, and no significant difference in second contact duration was observed among Tefluthrin (plus Cruiser) treatments (Table 2).

Within-treatment comparisons between mean first and second contact durations (Table 2) indicated no significant difference ($P > 0.05$) when wireworms were exposed to Dividend or untreated seeds, or to treatments containing Tefluthrin, except for the Tefluthrin and Cruiser combination treatment at 20 g a.i. in which second contact duration was significantly shorter than the first ($t = 3.01, P = 0.005$). Second contact duration was briefer than the first in the Vitavax Dual ($t = 3.67, P = 0.0004$), Cruiser at 5 g a.i. ($t = 2.91, P = 0.01$), 10 g a.i. ($t = 3.58, P = 0.0007$), 15 g a.i. ($t = 5.51, P < 0.0001$), 20 g a.i. ($t = 2.31, P = 0.02$) and 30 g a.i. ($t = 2.38, P = 0.02$) treatments, suggesting that these insecticides had an effect on $L. canus$ contact behaviour.

To further explore this sublethal effect, first and second contact durations were compared within treatments for only those larvae that came into contact twice (data for the first contact period not shown for this subgroup). This comparison revealed no significant difference ($P > 0.05$) when wireworms were first exposed to Dividend, Vitavax Dual or Cruiser at 5, 10 and 30 g a.i. However, second contact was significantly shorter than first contact when wireworms were exposed to Cruiser at 15 g a.i. [first contact = 141.9 (15.6) min, second contact = 37.7 (13.1); $t = 3.60, P = 0.001$] and 20 g a.i. [first contact = 90.0 (14.1), second contact = 63.0 (12.0); $t = 2.46, P = 0.02$], indicating that a single exposure to Cruiser could affect subsequent behaviour.

The effect of repeated contacts with Tefluthrin was assessed by within-treatment comparisons of the mean duration of the first, second and third contacts for $L. canus$ larvae that made three or more contacts with seeds in the 2006 study (Table 3). This comparison indicated no significant difference ($P > 0.05$) in contact duration among treatments. Similarly, comparisons among different treatments for all first, all second and all third contact durations indicated no significant difference among treatments ($P > 0.05$). However, in all treatments (except Tefluthrin at 10 g a.i.), the second inter-contact interval was considerably longer than the first inter-contact interval (Table 3). While these differences were not statistically significant ($P > 0.05$), it may indicate that repeated contact with Tefluthrin had an effect on wireworm behaviour. Previous work has shown that even very brief (1 min) contact with Tefluthrin-treated seeds will induce temporary morbidity in $L. canus$ (van Herk and Vernon 2007b).

### Total contact duration

Due to the repellence elicited by Tefluthrin, total contact duration in $L. canus$ in the 2006 study differed significantly among treatments ($F_{16,478} = 32.56, P < 0.0001$; Table 2). Contact in the Tefluthrin treatments was consistently shorter than in all other treatments, and the total contact duration in Cruiser (alone) treatments decreased as concentration increased, suggesting that when exposed to Cruiser, wireworm deterrence from contact increases with the concentration (Table 2).

Similarly, despite repeated contacts with seeds containing Tefluthrin (1-5 contacts), the total contact duration of $A. obscurus$ remained lowest among these treatments over the 180-min observation period ($F_{13,227} = 8.56, P < 0.0001$; Table 1), and it was significantly lower than in the control, Cruiser at 10 g a.i., Gaucho and Vitavax Dual treatments. In 2005, total contact duration of $L. canus$ over the observation period differed significantly among treatments ($F_{5,87} = 14.18, P < 0.0001$; Table 1), with total contact duration in the Tefluthrin and Poncho treatments being significantly shorter than in the control treatments (Table 1).

### Post-contact wireworm health and mortality

There were significant differences among treatments in the proportion of dead larvae of $A. obscurus$, $L. canus$ in 2005 and $L. canus$ in 2006 that were moribund at the end of the observation period ($\chi^2 = 119.7, df = 11, P < 0.0001$, Table 1; $\chi^2 = 32.87, df = 5, P < 0.0001$, Table 1; $\chi^2 = 128.81, df = 15, P < 0.0001$, Table 2). There were significant differences among treatments in the proportion of dead larvae at 56 DAE in $A. obscurus$ ($\chi^2 = 42.72, df = 11, P < 0.0001$), but not in the 2005 $L. canus$ study ($\chi^2 = 8.65, df = 5, P = 0.12$).

### Table 3. Contact and inter-contact durations (min) of $L. canus$ larvae exposed to wheat seeds in the 2006 study for wireworms that came into contact with seeds treated with Tefluthrin 20SC (and Cruiser 350FS) three times during a 300-min observation period

<table>
<thead>
<tr>
<th>Insecticide¹</th>
<th>C²</th>
<th>Mean (SEM) first contact</th>
<th>Mean (SEM) second contact</th>
<th>Mean (SEM) third contact</th>
<th>Mean (SEM) first inter-contact period</th>
<th>Mean (SEM) second inter-contact period</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tefluthrin (5)</td>
<td>23</td>
<td>20.4 (2.5)</td>
<td>20.4 (3.1)</td>
<td>22.4 (4.2)</td>
<td>47.2 (7.8)</td>
<td>57.2 (9.7)</td>
</tr>
<tr>
<td>Tefluthrin (10)</td>
<td>32</td>
<td>22.0 (2.2)</td>
<td>20.8 (3.6)</td>
<td>19.2 (4.7)</td>
<td>51.4 (10.0)</td>
<td>48.8 (9.1)</td>
</tr>
<tr>
<td>Tefluthrin (15)</td>
<td>13</td>
<td>24.2 (4.0)</td>
<td>25.0 (10.7)</td>
<td>15.0 (1.7)</td>
<td>32.7 (10.1)</td>
<td>59.2 (12.4)</td>
</tr>
<tr>
<td>Tefluthrin (30)</td>
<td>13</td>
<td>20.5 (2.2)</td>
<td>24.6 (13.5)</td>
<td>20.2 (2.2)</td>
<td>35.5 (10.7)</td>
<td>69.2 (13.5)</td>
</tr>
<tr>
<td>Cruiser (5) + Tefluthrin (5)</td>
<td>12</td>
<td>29.6 (4.8)</td>
<td>18.3 (3.7)</td>
<td>29.2 (8.2)</td>
<td>40.0 (11.4)</td>
<td>66.7 (12.9)</td>
</tr>
<tr>
<td>Cruiser (10) + Tefluthrin (10)</td>
<td>15</td>
<td>25.7 (4.4)</td>
<td>18.0 (2.2)</td>
<td>27.7 (7.3)</td>
<td>48.0 (9.6)</td>
<td>71.3 (14.8)</td>
</tr>
<tr>
<td>Cruiser (15) + Tefluthrin (15)</td>
<td>13</td>
<td>20.8 (2.3)</td>
<td>15.4 (3.1)</td>
<td>17.3 (2.8)</td>
<td>40.4 (9.2)</td>
<td>71.9 (18.9)</td>
</tr>
</tbody>
</table>

¹ Cruiser = Cruiser 350FS, Tefluthrin = Tefluthrin 20CS. Numbers in parentheses indicate grams a.i. 100 kg⁻¹ seed.
² C = number of wireworms that made three contacts.
There were also significant differences in the proportion of *L. canus* wireworms dead at 70 DAE ($X^2 = 157.04$, df = 15, $P < 0.0001$) in 2006.

**Control treatments**

Morbidity and mortality were not observed in *A. obscurus* or *L. canus* in the 2005 studies when larvae were exposed to control treatments (Table 1). Similarly, no morbidity and very low mortality was observed in *L. canus* larvae exposed to control treatments in the 2006 study (Table 2).

**Tefluthrin 20SC**

Morbidity of both *A. obscurus* and *L. canus* in the 2005 studies was low (< 50%) after bioassays with Tefluthrin and Tefluthrin plus Cruiser (Table 1), and all moribund wireworms had fully recovered by 7 DAE. Recovered wireworms did not subsequently relapse into morbidity, and no mortality occurred (Table 1, Figs. 1 and 2). Similarly, in the 2006 study, nearly all *L. canus* larvae that were moribund at the end of the observation period after bioassays with Tefluthrin or Tefluthrin plus Cruiser had fully recovered by 3 DAE (Fig. 3), only causing low (or zero) mortality in all treatments containing Tefluthrin (Table 2). Wireworms in all treatments containing Tefluthrin were often observed to be ‘writhing’ after contact at some point during the observation period, but turned out to be ‘alive’ when checked at the end of the observation period (data not shown).

**Cruiser 350FS**

Considerably more *A. obscurus* larvae were moribund after bioassays with Cruiser at 10 g a.i. than at 30 g a.i. (Table 1), but nearly all wireworms made a full recovery by 14 DAE (Fig. 1). Despite the high per-
A high percentage (90%) of moribund larvae, mortality was low in the 10 g a.i. treatment, while there was no mortality in the 30 g a.i. treatment (Table 1). A smaller proportion of *A. obscurus* larvae were moribund, but a slightly greater proportion died after bioassays with the Cruiser and Tefluthrin combination than after bioassays with either chemical at 10 g a.i. singly (Table 1).

In the 2006 study, no *L. canus* larvae were moribund after exposure to Cruiser at 5 g a.i. or Cruiser plus Tefluthrin at 20 g a.i., and only a low percentage (< 40%) of wireworms were moribund in all other Cruiser (plus Tefluthrin) treatments (Table 2). Wireworms that were moribund after exposure to Cruiser (all concentrations) had fully recovered by 14 DAE, but a considerable percentage (25-75%) relapsed thereafter and had died by 70 DAE (Fig. 3; Table 2).

Mortality in Cruiser treatments increased with concentration (data for Cruiser at 20 g a.i. were excluded from the analysis due to concerns regarding seed appetite, as discussed above). Mortality was significantly lower in bioassays with Tefluthrin plus Cruiser than in bioassays with Cruiser alone at the same rates.

**Vitavax Dual**

A high percentage of larvae of *A. obscurus* (100%) and *L. canus* (68%) in the 2005 studies were moribund after bioassays with Vitavax Dual (Table 1), but nearly all moribund wireworms had fully recovered by 28 DAE (Figs. 1 and 2). Similarly, a high percentage (74%) of *L. canus* larvae were moribund after bioassays with Vitavax Dual (Table 2), but all had fully recovered by 14 DAE and did not relapse subsequently, causing no mortality in this treatment (Table 2, Fig. 3).

**Other treatments**

A high percentage (> 70%) of *A. obscurus* larvae were moribund after bioassays with Admire 240FS, Gaucho 480FL and Poncho 600F (Table 1), but nearly all moribund wireworms in these treatments had fully recovered by 14 DAE (Fig. 1). However, some wireworms exposed to Admire and Gaucho relapsed and died 14 DAE (Fig. 1). Wireworm morbidity and mortality after contact with Poncho was similar for *L. canus* and *A. obscurus* in the 2005 studies (Table 1, Figs. 1 and 2).
DISCUSSION

Evaluation of insecticide treatments

Control treatments
Contact with untreated seeds or seeds treated with Dividend or Raxil had no significant impact on *A. obscurus* or *L. canus* health or mortality. In all three treatments, most (>50%) wireworms remained in contact with the seeds until the end of the observation period. In addition, the low morbidity rate in these control treatments suggests that the presence of fungicides in the other treatments likely did not affect wireworm health and behaviour, and will likely not cause wireworm mortality in the field.

**Tefluthrin 20SC**
Both *A. obscurus* and *L. canus* were strongly repelled by Tefluthrin. At all concentrations tested, wireworms only came into contact with treated seeds briefly before moving away. While brief contact induced morbidity in some wireworms, nearly all recovered fully and there was little mortality in either species. Both the low mortality and rapid recovery (without relapse) of *A. obscurus* and *L. canus* following Tefluthrin-induced morbidity are similar to previous observations in which *L. canus* larvae had been exposed to treated wheat seeds in Eppendorf tubes (van Herk and Vernon 2007b).

The similarity between first and second contact durations in both *A. obscurus* and *L. canus* exposed to Tefluthrin suggests that duration of the second

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**Figure 3. Health of Limonius canus larvae 0, 1, 3, 7, 14, 28, 42 and 56 d after exposure (DAE) to insecticide-treated wheat seeds during 300-min observation periods in a soil-window bioassay; 2006 study.** See text for an explanation of wireworm health categories.
contact was not affected by the previous contact (i.e. that wireworms were not repelled from the seeds more quickly), and that the larvae did not learn to avoid seeds treated with Tefluthrin. Since wireworms can become moribund within 5 min of contact with Tefluthrin and subsequently recover within 1-2 h (van Herk and Vernon 2007b), it is likely that wireworms observed to be ‘writhing’ after contact during bioassays with Tefluthrin were temporarily moribund from contact but had recovered before the end of the observation period.

Together these results indicate that Tefluthrin may be ineffective for reducing wireworm populations in the field, but may give stand protection in the field due to its repellence effect. Contrary to expectations, combining Tefluthrin with Cruiser reduced the efficacy of Cruiser by significantly reducing the duration of contact with seeds. Comparisons of post-contact health profiles between the Cruiser, Tefluthrin, and Tefluthrin plus Cruiser treatments (Fig. 3) indicate that wireworms exposed to both Tefluthrin and Cruiser responded in a similar way as those exposed to Tefluthrin alone.

**Cruiser 350FS**

Larvae of *A. obscurus* and *L. canus* were not repelled by Cruiser 350FS. While *L. canus* in the 2006 study moved away from seeds treated with Cruiser at 20 and 30 g a.i. more quickly than from seeds treated with Dividend, only a low proportion of larvae were moribund at the end of the observation period. The considerable mortality (increasing with concentration) in *L. canus* observed at all concentrations of Cruiser suggests that a 2-3 h contact suffices to kill wireworms. *Agriotes obscurus* exposed to Cruiser at 10 g a.i. remained in contact for a shorter period, became moribund more quickly, and experienced less mortality than *L. canus* exposed to the same rate. This result, along with the briefer contact period and lower morbidity of *A. obscurus* at 30 than at 10 g a.i., suggests that increasing the contact duration with Cruiser may increase wireworm mortality, and that increasing the concentration of Cruiser on wheat seeds may reduce contact duration in *A. obscurus* (possibly by accelerating the onset of morbidity). It also suggests that the optimum concentration of Cruiser for wireworm management may vary with wireworm species.

The difference in post-contact health and mortality between the Cruiser and Cruiser plus Tefluthrin treatments suggests that the reduced contact duration with seeds containing Tefluthrin reduced the amount of Cruiser absorbed either orally and/or dermally by the wireworms, and it therefore indicates that placing both insecticides on wheat seeds may be less effective for wireworm control than exposing wireworms to Cruiser alone.

**Vitavax Dual**

Lindane is thought to elicit repellence in some wireworm species, including in *L. californicus*, a species closely related to *L. canus* (Toba et al. 1988). However, while first contact duration for *L. canus* in the 2005 and 2006 studies was considerably shorter in the Vitavax Dual treatments than in the control treatments, neither *A. obscurus* nor *L. canus* were repelled by Vitavax Dual. Wireworms often remained in contact for 1 h or more, and most wireworms of both species became moribund in situ. Wireworms that moved away without becoming moribund often returned for subsequent contact(s). While most *A. obscurus* and *L. canus* were moribund by the end of the observation period, mortality was low both in 2005 and 2006, suggesting that wireworms may become moribund before ingesting and/or making contact with lethal doses of lindane. The difference in duration of the first contact between *L. canus* and *A. obscurus* exposed to Vitavax Dual may indicate that wireworm response to lindane differs among species.

**Other treatments**

*Agrionet obscurus* larvae were not repelled by Gaucheo 480FL or Admire 240FS in the 2005 study, indicating that neither formulation of imidacloprid elicits repellence at the concentrations tested. In contrast, previous studies in open-air bioassays suggested that imidacloprid at high concentrations (> 1% a.i. in water or acetone) was slightly repellent to *A. obscurus* (van Herk et al. 2008b). There were no significant differences in the proportion of moribund wireworms at the end of the observation period or dead at 56 DAE among the Admire and Gaucheo treatments, suggesting that both formulations affected wireworm health similarly. As in the Cruiser 350FS treatments, the relapse and death after temporary recovery of *A. obscurus* exposed to Gaucheo and Admire (Fig. 1) stress the importance of long-term post-contact health assessments.

Larvae of *A. obscurus* and *L. canus* were not repelled by Poncho 600F but contact with seeds was shorter than in all other insecticide treatments, except those containing Tefluthrin. Wireworms appeared to move away from treated seeds due to the onset of morbidity, and at the end of the observation period most wireworms that had come into contact with treated seeds were moribund. The subsequent low mortality in both wireworm species suggests that wireworms may become moribund before they can absorb lethal doses of insecticide. The similar response of *A. obscurus* to Poncho 600F at 25 g a.i. and Cruiser 350FS at 30 g a.i. is of interest as insects metabolize the active ingredient of Cruiser (thiamethoxam) into clothianidin (the active ingredient in Poncho) (Nauen et al. 2003).

**Impacts of wireworm repellence and morbidity on insecticide effectiveness**

The results presented here indicate that contact with insecticide-treated seeds in the soil may cause morbidity in wireworms without causing subsequent mortality and that, consequently, long-term post-contact wireworm health checks are important for assessing an insecticide’s efficacy. These results also indicate that wireworms will move towards seeds treated with insecticides but will often move away after contact. As an insect’s internal state (e.g. concentration of nutrients or immune peptides in hemolymph) affects its behaviour (Miller and Strickler 1984; Pompilio et al. 2006; Riddell and Mallon 2006), it is probable that the onset of morbidity is the stimulus that elicits repellence in wireworms. Wireworms that moved away from insecticide-treated seeds often
showed signs of morbidity later on, even if contact was very brief (i.e. in the tefluthrin treatments). Insecticides that cause a rapid induction of morbidity (e.g. tefluthrin) may not be effective for wireworm population control, as wireworms may be repelled before they can absorb toxic doses. While wireworms that are repelled may return for subsequent contact(s), repeated induction of morbidity by some insecticides (e.g. Tefluthrin) increases their ability to recover from morbidity (van Herk and Vernon 2007b). Furthermore, wireworms that become moribund after coming into contact with seeds treated with Poncho, Gaucho, Admire or Vitavax Dual may do so before ingesting enough insecticide to die. Since recovery from morbidity induced by these chemicals may continue for months (van Herk et al. 2008a; Vernon et al. 2008), wireworms returning to the same insecticide-treated plants that induced morbidity initially may not be affected once the plants are established (i.e. wheat and corn). Finally, while an insecticide that elicits repellence may permit crop establishment and provide stand protection, it may be ineffective for wireworm management, particularly if the crop matures in the soil (e.g. potatoes) and if wireworms return after the chemical’s repellent effects have dissipated.

Evaluation of bioassays
The rapid orientation and high proportion of wireworms that came into contact with wheat seeds in these studies confirm that, contrary to what Chaton et al. (2008) suggested, wireworm host finding is non-random. Like many subterranean insect larvae, wireworms follow CO2 gradients to find their hosts in the soil (Doane et al. 1975; Guerenstein and Hildebrand 2008) and are able to detect the presence of a single germinating wheat seed in the soil from a 20 cm distance (Doane and Klinger 1978; Westcott et al. 1980). These characteristics have been used to develop effective bioassays for studying wireworm behaviour (e.g. Doane et al. 1975; Horton and Landolt 2002; van Herk and Vernon 2007a).

The results presented here also confirm that observing wireworm position and behaviour every 5 min is sufficient to assess contact and feeding behaviour (van Herk and Vernon 2007a), and they indicate that different insecticides elicit different behaviours in wireworms. However, as these observations were conducted under laboratory conditions over limited observation periods, the results presented here may not accurately reflect what occurs in the field. Increasing the number of seeds in the bioassay or the duration of observation periods will likely increase wireworm mortality in treatments in which wireworms became moribund in situ or continued to come into repeated contact with treated seeds.

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