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Effects, but no interactions, of ubiquitous pesticide and parasite stressors on honey bee (*Apis mellifera*) lifespan and behaviour in a colony environment

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Summary

Interactions between pesticides and parasites are believed to be responsible for increased mortality of honey bee (*Apis mellifera*) colonies in the northern hemisphere. Previous efforts have employed experimental approaches using small groups under laboratory conditions to investigate influence of these stressors on honey bee physiology and behaviour, although both the colony level and field conditions play a key role for eusocial honey bees. Here, we challenged honey bee workers under *in vivo* colony conditions with sublethal doses of the neonicotinoid thiacloprid, the miticide tau-fluvalinate and the endoparasite *Nosema ceranae*, to investigate potential effects on longevity and behaviour using observation hives. In contrast to previous laboratory studies, our results do not suggest interactions among stressors, but rather lone effects of pesticides and the parasite on mortality and behaviour, respectively. These effects appear to be weak due to different outcomes at the two study sites, thereby suggesting that the

role of thiacloprid, tau-fluvalinate and *N. ceranae* and interactions among them may have been overemphasized. In the future, investigations into the effects of honey bee stressors should prioritize the use of colonies maintained under a variety of environmental conditions in order to obtain more biologically relevant data.

Introduction

All living organisms are exposed to a broad array of environmental stressors, including pests, parasites and contaminants. Mortality represents the strongest and most defined index of effect (i.e. death); however, sublethal impacts affecting behaviour and physiology can also be measured (e.g. Marcogliese and Pietrock, 2011; Pettis *et al.*, 2012; Schneider *et al.*, 2012b). To obtain a thorough understanding of the effects of a particular stressor or combination of stressors, it is therefore crucial to examine multiple potential indices of effect.

The Western honey bee (*Apis mellifera*; hereafter honey bee) is a eusocial insect that can be used to investigate the environmental effects of parasites and pesticides due to its well-described natural history and ease of maintenance in an experimental setting. Additional interest in honey bee health has been stimulated by severe colony mortalities reported recently (Neumann and Carreck, 2010). The widely distributed ectoparasitic mite *Varroa destructor* has been identified as one important driver for colony losses (Genersch *et al.*, 2010; Le Conte *et al.*, 2010; Dietemann *et al.*, 2012); however, it appears that concurrent assaults by multiple other stressors likely have a large influence on colony survival (Potts *et al.*, 2010). While the detrimental consequences of stressor-driven mortality are apparent, the dimensions of the impact of sublethal effects on honey bee colonies are often less visible. Sublethal effects can comprise various parameters ranging from anatomical and physiological impairments to more complex processes such as orientation or foraging behaviour (e.g. Desneux *et al.*, 2007). The functioning of the colony superorganism as a unit depends heavily on the social behaviours among the individuals in the hive because the coordination of fundamental tasks in a colony (e.g. brood care, cleaning, foraging,

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attending etc.) requires the transfer of relevant information among the members of the colony (Moritz and Southwick, 1992). Even though social in-hive behaviours are key for colony functioning, few studies have investigated potential stressor effects on social behaviour, despite data suggesting that stressors can influence other behaviours (e.g. foraging) (Schneider *et al.*, 2012a; Dussaubat *et al.*, 2013a).

The microsporidian *Nosema ceranae* is an obligatory intracellular midgut parasite that host-switched from the Eastern honey bee (*Apis cerana*) to the Western honey bee more than a decade ago (Paxton *et al.*, 2007). It has since developed a nearly ubiquitous distribution worldwide (e.g. Klee *et al.*, 2007; Williams *et al.*, 2008; Giersch *et al.*, 2009; Higes *et al.*, 2009a; Invernizzi *et al.*, 2009; Yoshiyama and Kimura, 2011). Despite numerous investigations of the impact of the parasite, its role in honey bee mortalities is highly debated (Fries, 2010; Higes *et al.*, 2013). Whereas some studies did not detect increased individual bee or colony mortality (e.g. Invernizzi *et al.*, 2009; Genersch *et al.*, 2010; Williams *et al.*, 2011; Dainat *et al.*, 2012; Martin *et al.*, 2013), others have reported lethal effects in the laboratory (Higes *et al.*, 2007) as well as colony deaths (Martín-Hernández *et al.*, 2007; Higes *et al.*, 2008; 2009b). Observed sublethal effects of *N. ceranae* on individuals include host immune suppression (Antúnez *et al.*, 2009), energetic stress (Mayack and Naug, 2009; 2010; Naug and Gibbs, 2009), as well as altered flight behaviour (Kralj and Fuchs, 2010; Dussaubat *et al.*, 2013a) and pheromone production (Dussaubat *et al.*, 2010). Numerous studies have demonstrated that parasites can alter the behaviour of infested honey bees (e.g. Wang and Mofller, 1970; Delfinado-Baker *et al.*, 1992); however, none have investigated if *N. ceranae* affects social behaviour within a colony setting.

Pesticides, acting singly or in combination, can also affect non-target organisms such as solitary bees (Sandrock *et al.*, 2014a), bumble bees (Fauser-Misslin *et al.*, 2014) and honey bees (Bortolotti *et al.*, 2003; Desneux *et al.*, 2007; Aliouane *et al.*, 2009; Wu *et al.*, 2011; Henry *et al.*, 2012; Sandrock *et al.*, 2014b). Doses of pesticides that exceed a certain threshold level (depending on substance and type of exposure) affect the survival of exposed honey bees, while sublethal doses of pesticides can exhibit various effects on individual honey bees, including development, learning performance and orientation (Desneux *et al.*, 2007; Blacquière *et al.*, 2012). While many studies have investigated this kind of pesticide effects on honey bees, similar to *N. ceranae*, little is known about the potential impact of pesticides on honey bee social behaviour at the colony level. The neonicotinoid crop protection insecticide thiacloprid and the pyrethroid tau-fluvalinate are

two pesticides widely applied to combat pest insects (Elbert *et al.*, 2008) and *V. destructor* (Tsigouri *et al.*, 2001), respectively. Residues of both substances are common in bee hive matrices; thiacloprid in honey (Tanner and Czerwenka, 2011), bee bread (Genersch *et al.*, 2010), nectar and pollen (Pohorecka *et al.*, 2012), and tau-fluvalinate in beeswax and pollen (Chauzat and Faucon, 2007; Mullin *et al.*, 2010). Thiacloprid is of relatively low toxicity to bees (oral LD₅₀ = 17.32 µg bee⁻¹) versus other neonicotinoids, and can act synergistically with *N. ceranae* to kill honey bees in the laboratory (Vidau *et al.*, 2011; Retschnig *et al.*, 2014a). Tau-fluvalinate has an acute contact toxicity of 0.2 g µg bee⁻¹, but was reported to have no lethal effect at daily oral doses of 5 or 10 µg bee⁻¹ (Decourtye *et al.*, 2005). However, it was shown to promote honey bee mortality in the presence of the miticide coumaphos (Johnson *et al.*, 2009) as well as influence honey bee locomotion (Teeters *et al.*, 2012). Although combined effects of tau-fluvalinate and any neonicotinoid have not yet been investigated in honey bees, exposure of bumble bees to a similar combination of pesticides (i.e. a neonicotinoid and a pyrethroid) increased worker mortality and impaired foraging behaviour (Gill *et al.*, 2012).

The simultaneous exposure to a combination of parasites and pesticides can lead to interactions between the stressors in the host and can cause increased host mortality or various sublethal effects (Marcogliese and Pietrock, 2011). For example, in honey bees, concurrent exposure to *N. ceranae* and certain neonicotinoid insecticides caused both lethal and sublethal effects (e.g. Alaux *et al.*, 2010; Vidau *et al.*, 2011; Aufauvre *et al.*, 2012; Pettis *et al.*, 2012). In the past, the investigation of specific mechanisms of stressor effects often took place in laboratory studies under standardized conditions (e.g. Alaux *et al.*, 2010; Forsgren and Fries, 2010; Aufauvre *et al.*, 2012), which allowed for the control of potentially interfering factors (Williams *et al.*, 2013). However, it remains unclear to what extent such findings can be extrapolated to honey bees in the field. As demonstrated in previous investigations, the study arena (laboratory versus field) can have a strong influence on the physiological development (Maleszka *et al.*, 2009) as well as measured stressor effects in individual bees, including interactive effects of pesticides on honey bee mortality (Schmuck *et al.*, 2003). Naturally, laboratory studies focus on parameters that can be tested reliably in this particular study arena, including worker longevity and parasite intensity (e.g. Alaux *et al.*, 2010; Vidau *et al.*, 2011). However, some traits that are crucial for the functioning of the honey bee colony, such as the social in-hive behaviour of the workers, have received too little attention so far. Therefore, the primary objective of this study was to look at potential stressor effects on honey bee worker

longevity, the ultimate measure of stress impact, as well as on important behaviours among workers including antennation (communication), grooming (hygiene) and trophallaxis (nutrition), as well as flight activity (nutrition and hygiene) (Moritz and Southwick, 1992). Using observation hives in two locations, we investigated the lethal and sublethal effects of the widely applied pesticides thiacloprid and tau-fluvalinate, as well as the ubiquitous parasite *N. ceranae*, on individual honey bees that faced natural conditions. Experimental individuals were allocated to one of four treatment groups (control, pesticides, *N. ceranae*, and *N. ceranae* and pesticides); pesticide and *N. ceranae* exposure occurred during development and post-emergence, respectively. Due to previous reports of the effects of *N. ceranae* and pesticides on honey bee survival and behaviour (e.g. Alaux *et al.*, 2010; Kralj and Fuchs, 2010; Aufauvre *et al.*, 2012), we expected to observe a similar impact of these stressors and anticipated to find stronger effects on individuals that were exposed to the combination of both *N. ceranae* and pesticides due to potential synergistic interactions (Alaux *et al.*, 2010; Vidau *et al.*, 2011; Aufauvre *et al.*, 2012; Pettis *et al.*, 2012).

Results

Mortality

Location A. Honey bee workers exposed to pesticides during development showed significantly higher mortality than did control individuals during the 14 day trial (Kaplan–Meier, log-rank test with Bonferroni correction, both $P_s = 0.0006$, Fig. 1). No such significant difference

Survivorship (%)

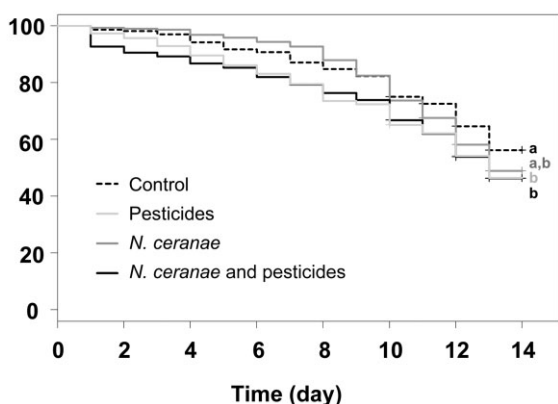


Fig. 1. Kaplan–Meier survival curve of the experimental honey bee (*Apis mellifera*) workers at location A (Switzerland). Workers that were exposed to pesticides (thiacloprid and tau-fluvalinate) during development showed significantly higher mortality than the control group (log-rank test with Bonferroni correction, both $P_s = 0.0006$). Significant differences between treatments are marked with different letters (a, b).

Survivorship (%)

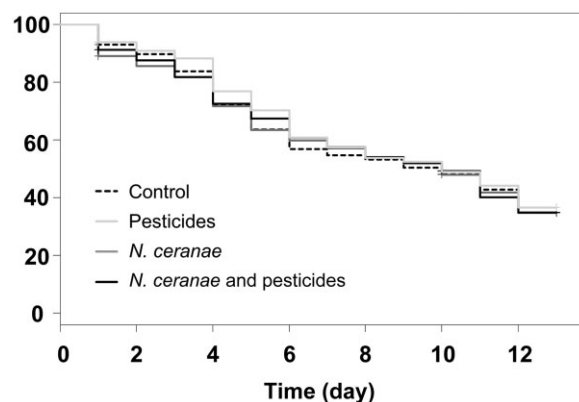


Fig. 2. Kaplan–Meier survival curve of the experimental honey bee workers at location B (Germany). No differences in mortality were observed between the investigated treatment groups (log-rank test with Bonferroni correction, all $P_s = 1$).

was observed between control workers and those belonging to the *N. ceranae*-only treatment group (Kaplan–Meier, log-rank test with Bonferroni correction, $P = 0.3$). Similarly, no significant difference in mortality occurred among the non-control treatment workers (pesticides versus *N. ceranae*, pesticides versus *N. ceranae* and pesticides, and *N. ceranae* versus *N. ceranae* and pesticides, Kaplan–Meier, log-rank test with Bonferroni correction, $P = 0.19$; 1; 0.23). Mortality, when compared using only data at termination day, was similar to survival analyses that incorporated daily mortality; workers exposed to pesticides showed significantly higher mortality compared with control individuals, and no significant difference was observed among non-control treatment individuals (binary logistic regression with Bonferroni correction, pesticides groups versus control, both $P_s < 0.012$, for all other comparisons $P_s > 0.186$).

Location B. No significant difference in mortality was observed among treatments when daily deaths (Kaplan–Meier, log-rank test with Bonferroni correction, all $P_s = 1$, Fig. 2), or total death number at experiment termination (binary logistic regression with Bonferroni correction, all $P_s = 1$) were considered.

Comparison of mortality between locations A and B. In all treatment groups, the workers showed significantly higher mortality in location B compared with location A (log-rank test, all $P_s < 0.001$, Table 1, Figs. 1 and 2).

Behaviour

Location A – In-hive behaviour. A total of 22147 individual behaviours were observed during 14 days (Fig. 3),

Table 1. Overview and comparison of the stressor impacts on honey bees (*Apis mellifera*) in locations A (Switzerland) and B (Germany).

| Treatments | Mortality | | | Behaviour | |
|----------------------------------|----------------------------------|------------|---------------------------------------|-----------------------------------|------------|
| | Location A | Location B | Location A versus B | Location A | Location B |
| Control | – | – | Higher mortality in B ($P < 0.001$) | – | – |
| Pesticides | Higher mortality ($P < 0.001$) | – | Higher mortality in B ($P < 0.001$) | – | – |
| <i>N. ceranae</i> | – | – | Higher mortality in B ($P < 0.001$) | Higher inactivity ($P < 0.001$) | – |
| <i>N. ceranae</i> and pesticides | Higher mortality ($P < 0.001$) | – | Higher mortality in B ($P < 0.001$) | Higher inactivity ($P < 0.05$) | – |

The absence of significant effects is marked as '–' in the table.

with frequency of observations of the three behavioural categories consistent for each treatment: other (including all behaviours except for social behaviours and motionlessness, such as walking, feeding, brood care, cleaning etc.) was observed most (total: 16280 events, 71.41–75.18% events per treatment), followed by motionless (total 3250 events, 13.09–16.37% events per treatment), and social (antennation total: 1458 events, 6.21–6.94% events per treatment; grooming total: 696 events, 3.0–3.29% events per treatment; trophallaxis total: 463 events, 1.99–2.26% events per treatment). For all possible combinations of behaviour comparisons ($n = 24$), only three showed significant differences (all P s < 0.05); all others had P -values greater than 0.23 (multinomial logistic regression with False discovery rate (FDR) correction, Fig. 3). Workers inoculated with *N. ceranae* (*N. ceranae*, and *N. ceranae* and Pesticides), regardless of pesticide exposure, were motionless more than control individuals (multinomial logistic regression with FDR correction, both $P < 0.016$). Additionally, workers exposed to *N. ceranae* only were motionless more than those exposed to pesticides only (multinomial logistic regression with FDR correction, $P = 0.0024$, Table 2 and Fig. 3).

Location B – Flight activity. There were no significant differences in flight activity, measured as number of flights per minute, among the three treatment groups (pesticides, *N. ceranae* or the combination of both) and the controls [analysis of variance (ANOVA) and Tukey–Kramer test, $P < 0.05$]. However, the *N. ceranae* treatment group showed significantly higher flight activity compared with the pesticides treatment group (ANOVA and Tukey–Kramer test, $P < 0.05$).

Treatment confirmation

Pesticides. Pesticide application to the donor colonies was confirmed by residue analyses of the respective chemical substances in the feeding solutions as well as of different hive matrices. Sucrose feed contained an average level of 611.5 ppb of thiacloprid in the treatment

and no detectable thiacloprid residues in the control solutions. In the pesticide-treated colonies, thiacloprid residues were detected in honey (190 ppb), wax (147 ppb) and pollen (68 ppb), whereas tau-fluvalinate was found in wax (8280 ppb) and pollen (105 ppb). In the control colonies, traces of thiacloprid (7.7 ppb in honey, 34.2 ppb in wax and 3.6 ppb in pollen), but not tau-fluvalinate, was detected.

Nosema ceranae

Location A. Workers inoculated with *N. ceranae* showed mean spore amounts of 14.32×10^6 [standard deviation (SD): 5.73×10^6] for the *N. ceranae* only and 14.56×10^6 (SD: 6.31×10^6) for the *N. ceranae* and pesticides treatment group. There was no significant difference between the spore amounts of these two treatment groups (Wilcoxon rank-sum test, $P > 0.05$). Workers that were not inoculated with *N. ceranae* showed median spore

Proportion of behaviours (%)

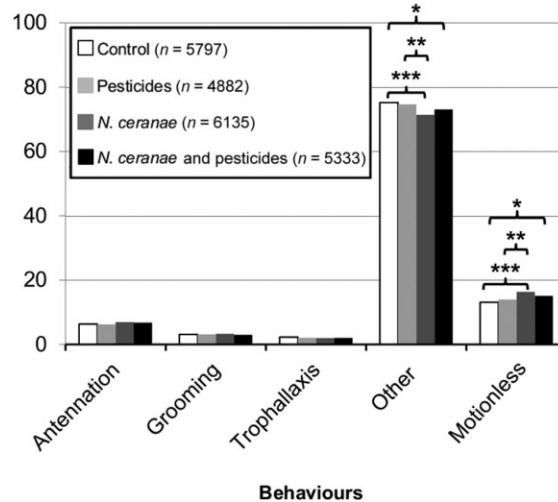


Fig. 3. Frequency of honey bee behaviours in the different treatments at location A (Switzerland). Significant differences among treatments were detected only between the behavioural categories being idle and other behaviours and are indicated with asterisks (* $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$).

Table 2. Comparisons of behaviour ratios (reference: category 'other behaviours') in honey bees (*Apis mellifera*) among pairs of treatments.

| Compared treatments | Behaviour (Reference: other behaviours) | P-value with FDR correction |
|-----------------------------------------------------------|-----------------------------------------|-----------------------------|
| Control versus pesticides | Motionless | 0.5573 |
| | Antennation | 0.9698 |
| | Grooming | 0.9698 |
| | Trophallaxis | 0.8414 |
| Control versus <i>N. ceranae</i> | Motionless | 0.0000 ^a |
| | Antennation | 0.2304 |
| | Grooming | 0.6352 |
| | Trophallaxis | 0.8238 |
| Control versus <i>N. ceranae</i> and pesticides | Motionless | 0.0152 ^b |
| | Antennation | 0.5573 |
| | Trophallaxis | 0.8238 |
| | Grooming | 0.9698 |
| Pesticides versus <i>N. ceranae</i> and pesticides | Motionless | 0.2619 |
| | Antennation | 0.5573 |
| | Grooming | 0.9698 |
| | Trophallaxis | 0.9698 |
| <i>N. ceranae</i> versus Pesticides | Motionless | 0.0024 ^c |
| | Antennation | 0.2304 |
| | Grooming | 0.7272 |
| | Trophallaxis | 0.9698 |
| <i>N. ceranae</i> versus <i>N. ceranae</i> and pesticides | Motionless | 0.2304 |
| | Antennation | 0.8238 |
| | Grooming | 0.6155 |
| | Trophallaxis | 0.9698 |

a. Motionless was more frequent in the *N. ceranae* treatment group.
b. Motionless was more frequent in the *N. ceranae* and pesticides treatment group.

c. Motionless was more frequent in the *N. ceranae* treatment group. Higher motionlessness was observed in three out of 24 treatment comparisons (in grey). Footnotes at the end of the table give more details about the direction of the outcome.

amounts (not normally distributed) of 0 spores per bee. However, 21 (35%) and 24 (40%) of the 60 workers analysed from the control and pesticides-only treatment groups were infected with *N. ceranae* at day 14. Mean spore counts in these workers were 8.73×10^6 spores per bee in the control and 10.8×10^6 spores per bee in the pesticides treatment group. Compared with inoculated workers, mean infection level in the non-inoculated individuals was significantly lower (ANOVA and Tukey–Kramer test, $P < 0.001$).

Location B. Mean spore counts of the inoculated workers were 2.93×10^6 (SD: 4.54×10^6) for the *N. ceranae* only and 2.33×10^6 (SD: 3.19×10^6) for the *N. ceranae* and pesticides treatment group. There was no significant difference between spore counts of these treatment groups (Wilcoxon rank-sum test, $P > 0.05$). Workers not inoculated with *N. ceranae* showed mean spore counts of 0.06×10^6 (SD: 0.25×10^6) spores per bee in the control and 0.014×10^6 (SD: 0.07×10^6) in the pesticides treatment group. Five control workers (8.62%) and three (5.45%) individuals of the pesticides-only treatment group

were infected with *N. ceranae* at the end of the study with mean spore counts of 0.67×10^6 and 0.26×10^6 , respectively. Compared with the *N. ceranae* inoculated workers, the mean infection level in those not fed *N. ceranae* was significantly lower (ANOVA and Tukey–Kramer test, $P < 0.01$).

Discussion

The data consistently revealed for both study locations no evidence of any interactions between parasite and pesticide stressors, as well as no effect of *N. ceranae* on worker mortality. However, overall worker mortality and the effect of pesticide exposure on mortality differed between the two locations. *Nosema ceranae* influenced in-hive activities by increasing frequency of motionless behaviour, but did not show an effect on flight activity.

The field-realistic approach of this study allowed for stressor exposure and collection of mortality and behavioural data under colony conditions. The vast majority of stressor-specific investigations are performed in the laboratory (e.g. Alaux *et al.*, 2010; Forsgren and Fries, 2010; Aufauvre *et al.*, 2012). Although this promotes a relatively controlled environment whereby potentially confounding factors can be more easily excluded (e.g. temperature, humidity, nutrition, etc.) (Williams *et al.*, 2013), results may not always reflect natural conditions because important features to honey bees, like eusociality, are not well represented (e.g. Mattila and Otis, 2006; Maleszka *et al.*, 2009; Retschnig *et al.*, 2014b). Alternatively, incidental exposure of experimental workers to *N. ceranae* and pesticides in colony-level studies is typically much greater than those used for laboratory assays. Similar to Wu and colleagues (2011), traces of pesticide residues were detected in control hives, possibly due to drifting bees or environmental contamination (e.g. Mullin *et al.*, 2010). Likewise, some control workers were infected with *N. ceranae*; this is not surprising as contaminated hive materials are believed to be major sources of *N. ceranae* infection (Higes *et al.*, 2008; Giersch *et al.*, 2009). The mean *N. ceranae* spore amounts of the respective treatment groups were in line with other studies that applied similar methods (e.g. Paxton *et al.*, 2007; Alaux *et al.*, 2010; Pettis *et al.*, 2012).

For both parasite and pesticide stressors, the effects on worker mortality were not consistent at the two study locations. Strong effects on honey bee health are usually highly reproducible, such as the considerable damage due to *V. destructor* parasitism (e.g. Liebig, 2001; Fries *et al.*, 2003; Rosenkranz *et al.*, 2010; Schäfer *et al.*, 2010). Inconsistencies of stressor effects in both locations suggest that they are rather weak. Regardless, pesticide exposure of immature workers increased mortality at the

adult stage in one study location, thereby supporting previous work that showed increased mortality of adults can occur when larvae are exposed to pesticides (Wu *et al.*, 2011; Pettis *et al.*, 2012; Berry *et al.*, 2013).

Sublethal application of either tau-fluvalinate (Berry *et al.*, 2013) or thiacloprid (Siede *et al.*, 2014) on honey bee colonies did not reveal measurable effects on the population dynamics of bees or brood. Here we present the first approach to measure the combined application of these two pesticides at the colony level. In bumblebees, the combination of a neonicotinoid and pyrethroid was demonstrated to increase worker mortality (Gill *et al.*, 2012); our study also observed this effect in honey bees in one location. In contrast to mortality, the data showed no evidence for an impact of the pesticides on the observed behaviours as has been shown in bumblebees (Gill *et al.*, 2012). This could be explained because previous studies that demonstrated sublethal pesticide effects typically applied similar doses of pesticides that have a comparatively higher toxicity, such as clothianidin or imidacloprid (Schneider *et al.*, 2012a; Teeters *et al.*, 2012; Yang *et al.*, 2012).

Nosema ceranae showed no effect on honey bee mortality at both locations. This is in line with a growing number of studies (e.g. Invernizzi *et al.*, 2009; Genersch *et al.*, 2010; Williams *et al.*, 2011; Martin *et al.*, 2013), but contrary to others (e.g. Higes *et al.*, 2007; Martín-Hernández *et al.*, 2007; Higes *et al.*, 2008; Higes *et al.*, 2009b; Williams *et al.*, 2014). This may be explained by variable strains of *N. ceranae* exhibiting a different virulence or differential susceptibility of bees in different geographic regions (Dussaubat *et al.*, 2013b). A further reason for the different outcomes may be that the effect of *N. ceranae* on individual honey bee mortality has so far been tested in laboratory studies only, where the bees might have been influenced by more stressful conditions compared with a natural colony environment (e.g. Retschnig *et al.*, 2014b). Although *N. ceranae* appeared to not influence flight activity at one location, the parasite reduced the overall activity of bees at the other location. This might be explained by the energetic stress caused by *N. ceranae* (Mayack and Naug, 2009; 2010; Naug and Gibbs, 2009).

In contrast to previously reported synergistic effects between neonicotinoid pesticides and *N. ceranae* (Alaux *et al.*, 2010; Vidau *et al.*, 2011; Aufauvre *et al.*, 2012; Pettis *et al.*, 2012), our data provided no such evidence. A potential explanation for this difference may be that previous studies were carried out under laboratory conditions. It is known that influence of stressors may differ depending on test arena (e.g. laboratory versus field) (Schmuck *et al.*, 2003; Mattila and Otis, 2006), which may potentially be a consequence of a higher sensitivity due to the artificial conditions in the laboratory (e.g. Huang *et al.*,

2014; Retschnig *et al.*, 2014b). Experimental workers in the present study lived in a colony environment (i.e. natural hive composition including queen, workers and drones) where they could feed (pollen, bee bread, honey), socially interact and exit the hive.

The mortality of the experimental workers in the two study locations showed clear differences, and the significantly greater worker mortality at one location compared with the other was consistent for all treatments. The workers that remained geographically closer to their donor colonies showed an overall better survival. Although it is difficult to determine mechanisms for these differences due to experimental methods, potential reasons for the higher mortality in the second location may include factors such as genotype–environment interactions (e.g. Costa *et al.*, 2012) or the transportation of the bees in the pupal stage (300 km) (Oldroyd, 2007; Pettis and Delaplane, 2010; Pirk *et al.*, 2014). Such potential impacts should be considered in future studies and closely investigated to improve the investigation of honey bee stressors in natural conditions.

Experimental procedures

Study set-up

The study was performed in summer 2012 at two locations: location A: Bern, Switzerland; and location B: Stuttgart, Germany. Both locations employed experimental honey bee workers from the same donor colonies located in Bern, Switzerland. Four treatment groups: (i) control, (ii) pesticides (thiacloprid and tau-fluvalinate), (iii) *N. ceranae* and (iv) *N. ceranae* and pesticides were investigated for differences in survivorship (both locations), in-hive behaviour (location A) and flight activity (location B).

At location A, eight local European honey bee colonies (*A. mellifera*) headed by sister queens (hereafter called donor colonies for the experimental workers) were randomly assigned to either the pesticide or the non-pesticide treatment ($n=4$ per group). For the pesticide treatments, thiacloprid and tau-fluvalinate were applied for 6 weeks to encompass two complete brood cycles prior to removal of workers for the experiments. Thiacloprid was administered weekly by supplying colonies with 1 kg of 1000 ppb of 98.0% thiacloprid (Dr. Ehrenstorfer GmbH, Germany) sucrose solution (72–73% Hostettler® syrup, Hostettler Spezialzucker AG, Switzerland) using an in-hive feeder; control workers were fed with sucrose solution only. Tau-fluvalinate was applied using two Apistan® strips (Vita [Europe] LTD, UK), each 0.8 g active substance, placed in the lower brood chamber of each colony according to the manufacturer's recommendations. To confirm exposure, the thiacloprid and control solutions, as well as honey, wax and pollen samples were collected and analysed for pesticide residues at the United States Department of Agriculture (USDA) National Science Laboratory, Gastonia, USA, using routine liquid chromatographic procedures (Mullin *et al.*, 2010).

Two weeks prior to the start of data collection, three observation hives were set up using standard approaches

(Scheiner *et al.*, 2013) in both locations A and B. Briefly, each observation hive was equipped with a mated egg-laying local queen of the same year and two stacked Zander frames containing ~2000 bees: one frame contained brood in various developmental stages and the other consisted of stored honey and empty cells.

To obtain age cohorts of workers for experiments, queens from the eight donor colonies were caged on an empty brood frame for 48 h. Prior to emergence, brood frames were transferred to the laboratory and maintained in frame holders in the dark at 34.5°C and ≥ 50% relative humidity in an incubator (Williams *et al.*, 2013). For transport to location B, brood frames (1–2 per donor colony) containing age cohorts of workers within capped brood cells (i.e. pupae) were carefully added to the brood chambers of a full-size colony for the ~300 km journey by car. Frames were kept under the same conditions as described above upon arrival at the new site.

After emergence, workers (4752 in total) at both locations were randomly assigned to the appropriate treatment group, marked on the thorax using coloured number plates and paints (Marabu Brillant, Gerstaecker, Switzerland) and inoculated with either *N. ceranae* or control suspension using a group feeding approach (Fries *et al.*, 2013). For this, workers were starved for approximately 2 h in disposable plastic cages (20 individuals per cage). For the *N. ceranae* inoculum, fresh spores were obtained from naturally infected foragers that were collected at the hive entrance of local colonies in both sites the day prior. Midguts were carefully extracted from the workers using forceps, crushed in water and then purified by multiple centrifugation runs at 5000 g (Fries *et al.*, 2013). Spores were then quantified using light microscopy and a haemocytometer (Cantwell, 1970). Dilution of the suspension using 50% (w/v) sucrose solution yielded a final concentration of 2 000 000 spores per 1.5 ml, whereas the control solution consisted of only freshly prepared 50% (w/v) sucrose solution. Each disposable plastic cage was supplied with either 1.5 ml *N. ceranae* or control inoculum using a 2 ml microcentrifuge tube with a 2 mm diameter hole in bottom tip to allow feeding, thus providing each of the 20 workers per cage with ~100 000 spores. Feeding devices were filled with 50% (w/v) sucrose solution when the entire suspension was consumed during frequent checks; after 48 h, all devices were refilled completely. After the inoculation process, a total of 792 workers, 198 individuals per treatment per observation hive, were sprayed with sucrose solution and carefully inserted into the appropriate observation hive at night.

Location A

Mortality and behaviours of experimental honey bee workers was assessed by examining the observation hives twice daily, between 09h00–12h00 and 14h00–17h00, during 14 consecutive days. Order of observation hive viewing was rotated daily to avoid a potential bias of time. Observed behaviours were allocated to the following categories: (i) social interactions between adults (i.e. antennation, trophallaxis and grooming), (ii) motionless (i.e. individual is not moving) and (iii) other (i.e. performing any task not included in the previous categories, see Scheiner *et al.*,

2013). Social behaviours of experimental workers with two or more other individuals were defined as the following: antennation (contact of the moving antennae), trophallaxis (exchange of food) and grooming (cleaning manipulation using the mouthparts and antennae).

At day 14, all surviving workers were carefully collected using forceps from observation hive frames and immediately frozen at –20°C. To ensure maximum recovery of marked workers, multiple collection attempts occurred during day and night. A subsample of 20 collected workers per treatment group per observation hive was used to determine *N. ceranae* infection levels. This was achieved by homogenizing each individual in a 2 ml Eppendorf tube using a bead mill homogenizer (MM300 Retsch), one metal bead and 1 ml of nuclease-free water. *Nosema ceranae* quantification was performed according to previously mentioned techniques.

Location B

Similar to location A, mortality at location B was determined daily by recording all of the marked workers. Flight activity observations occurred between day 7 post-insertion of the marked workers until day 13, when the experiment was terminated. Departing and returning workers were viewed through a 10 cm long transparent plastic tube connecting the colony to the outdoors. Workers surviving to day 13 were collected according to previously discussed methods for location A. Similarly, a subsample of 16–28 workers, depending on number of available bees after collection, per treatment group and observation hive was used to determine *N. ceranae* infection levels. This was achieved by pressing out the midgut content by gently squeezing the abdomen of each individual. The gut suspension was viewed using light microscopy and a haemocytometer according to Cantwell (1970).

Statistics

Differences in survival of experimental workers during the study were tested using Kaplan–Meier survival statistics with the log-rank test (Mantel–Haenszel test) and Bonferroni correction, whereas survival at experiment termination was tested using binary logistic regression using tests that are based on the standard normal z-statistic (Wald statistic). For these analyses, workers collected at the end of the experiment were considered censored, as were those observed but not collected on the final day. Furthermore, workers that disappeared during the experiment were considered dead on the last day they were seen. Differences in survival of the workers between the two locations were analysed using the log-rank test. Comparison of social interactions between adults, motionless and other behaviours among treatments were performed using multinomial logistic regression with *P*-values deduced from Wald statistics using the category ‘other behaviours’ as a reference. Thus, the ratio of one specific behaviour versus other was compared between two treatment groups for each case. FDR correction was applied to compensate for multiple comparisons (Benjamini and Hochberg, 1995). Flight activity was compared using repeated measures ANOVA. *Nosema ceranae* data were

analysed using Kruskal–Wallis one-way ANOVA, because of non-normal data distribution, followed by the Tukey–Kramer multiple comparison tests. All statistical analyses were carried out using the programmes SYSTAT 13 (Systat Software, USA), R (version 3.0.0., The R Foundation for statistical computing platform) and NCSS (version 8, NCSS LLC, USA).

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References

- Alaux, C., Brunet, J.L., Dussaubat, C., Mondet, F., Tchamitchan, S., Cousin, M., *et al.* (2010) Interactions between *Nosema* microspores and a neonicotinoid weaken honey bees (*Apis mellifera*). *Environ Microbiol* **12**: 774–782.
- Aliouane, Y., el Hassani, A.K., Gary, V., Armengaud, C., Lambin, M., and Gauthier, M. (2009) Subchronic exposure of honeybees to sublethal doses of pesticides: effects on behavior. *Environ Toxicol Chem* **28**: 113–122.
- Antúnez, K., Martín-Hernández, R., Prieto, L., Meana, A., Zunino, P., and Higes, M. (2009) Immune suppression in the honey bee (*Apis mellifera*) following infection by *Nosema ceranae* (Microsporidia). *Environ Microbiol* **11**: 2284–2290.
- Aufauvre, J., Biron, D.G., Vidau, C., Fontbonne, R., Roudel, M., Diogon, M., *et al.* (2012) Parasite-insecticide interactions: a case study of *Nosema ceranae* and fipronil synergy on honeybee. *Sci Rep* **2**: 1–7.
- Benjamini, Y., and Hochberg, Y. (1995) Controlling the false discovery rate – a practical and powerful approach to multiple testing. *J R Stat Soc B* **57**: 289–300.
- Berry, J.A., Hood, W.M., Pietravalle, S., and Delaplane, K.S. (2013) Field-level sublethal effects of approved bee hive chemicals on honey bees (*Apis mellifera* L.). *PLoS ONE* **8**: e76536.
- Blacquière, T., Smaghe, G., van Gestel, C.A.M., and Mommaerts, V. (2012) Neonicotinoids in bees: a review on concentrations, side-effects and risk assessment. *Ecotoxicology* **21**: 973–992.
- Bortolotti, L., Montanari, R., Marcelino, J., Medrzycki, P., Maini, S., and Porrini, C. (2003) Effects of sub-lethal imidacloprid doses on the homing rate and foraging activity of honey bees. *B Insectol* **56**: 63–67.
- Cantwell, G.E. (1970) Standard methods for counting *Nosema* spores. *Am Bee J* **110**: 222–223.
- Chauzat, M.P., and Faucon, J.P. (2007) Pesticide residues in beeswax samples collected from honey bee colonies (*Apis mellifera* L.) in France. *Pest Manag Sci* **63**: 1100–1106.
- Costa, C., Buchler, R., Berg, S., Bienkowska, M., Bouga, M., Bubalo, D., *et al.* (2012) A Europe-wide experiment for assessing the impact of genotype-environment interactions on the vitality and performance of honey bee colonies: experimental design and trait evaluation. *J Apic Sci* **56**: 147–158.
- Dainat, B., Evans, J.D., Chen, Y.P., Gauthier, L., and Neumann, P. (2012) Predictive markers of honey bee colony collapse. *PLoS ONE* **7**: e32151.
- Decourtye, A., Devillers, J., Genecque, E., Le Menach, K., Budzinski, H., Cluzeau, S., and Pham-Delègue, M.H. (2005) Comparative sublethal toxicity of nine pesticides on olfactory learning performances of the honeybee *Apis mellifera*. *Arch Environ Contam Toxicol* **48**: 242–250.
- Delfinado-Baker, M., Rath, W., and Boecking, O. (1992) Phoretic bee mites and honeybee grooming behavior. *Int J Acarol* **18**: 315–322.
- Desneux, N., Decourtye, A., and Delpuech, J.M. (2007) The sublethal effects of pesticides on beneficial arthropods. *Annu Rev Entomol* **52**: 81–106.
- Dietemann, V., Pflugfelder, J., Anderson, D., Charriere, J.D., Chejanovsky, N., Dainat, B., *et al.* (2012) *Varroa destructor*: research avenues towards sustainable control. *J Apic Res* **51**: 125–132.
- Dussaubat, C., Maisonnasse, A., Alaux, C., Tchamitchan, S., Brunet, J.L., Plettner, E., *et al.* (2010) *Nosema* spp. infection alters pheromone production in honey bees (*Apis mellifera*). *J Chem Ecol* **36**: 522–525.
- Dussaubat, C., Maisonnasse, A., Crauser, D., Beslay, D., Costagliola, G., and Soubeyrand, S. (2013a) Flight behavior and pheromone changes associated to *Nosema ceranae* infection of honey bee workers (*Apis mellifera*) in field conditions. *J Invertebr Pathol* **113**: 42–51.
- Dussaubat, C., Sagastume, S., Gómez-Moracho, T., Botías, C., García-Palencia, P., Martín-Hernández, R., *et al.* (2013b) Comparative study of *Nosema ceranae* (Microsporidia) isolates from two different geographic origins. *Vet Microbiol* **162**: 670–678.
- Elbert, A., Haas, M., Springer, B., Thielert, W., and Nauen, R. (2008) Applied aspects of neonicotinoid uses in crop protection. *Pest Manag Sci* **64**: 1099–1105.
- Fausser-Misslin, A., Sadd, B.M., Neumann, P., and Sandrock, C. (2014) Influence of combined pesticide and parasite exposure on bumblebee colony traits in the laboratory. *J Appl Ecol* **51**: 450–459.
- Forsgren, E., and Fries, I. (2010) Comparative virulence of *Nosema ceranae* and *Nosema apis* in individual European honey bees. *Vet Parasitol* **170**: 212–217.
- Fries, I. (2010) *Nosema ceranae* in European honey bees (*Apis mellifera*). *J Invertebr Pathol* **103**: 73–79.
- Fries, I., Hansen, H., Imdorf, A., and Rosenkranz, P. (2003) Swarming in honey bees (*Apis mellifera*) and *Varroa destructor* population development in Sweden. *Apidologie* **34**: 389–397.
- Fries, I., Chauzat, M.P., Chen, Y.P., Doublet, V., Genersch, E., Gisder, S., *et al.* (2013) Standard methods for *Nosema* research. *J Apic Res* **52**: 52.1.14 <http://dx.doi.org/10.3896/IBRA.1.52.1.14>

- Genersch, E., von der Ohe, W., Kaatz, H., Schroeder, A., Otten, C., Buchler, R., *et al.* (2010) The German bee monitoring project: a long term study to understand periodically high winter losses of honey bee colonies. *Apidologie* **41**: 332–352.
- Giersch, T., Berg, T., Galea, F., and Hornitzky, M. (2009) *Nosema ceranae* infects honey bees (*Apis mellifera*) and contaminates honey in Australia. *Apidologie* **40**: 117–123.
- Gill, R.J., Ramos-Rodriguez, O., and Raine, N.E. (2012) Combined pesticide exposure severely affects individual- and colony-level traits in bees. *Nature* **491**: 105–119.
- Henry, M., Beguin, M., Requier, F., Rollin, O., Odoux, J.F., Aupinel, P., *et al.* (2012) A common pesticide decreases foraging success and survival in honey bees. *Science* **336**: 348–350.
- Higes, M., García-Palencia, P., Martín-Hernández, R., and Meana, A. (2007) Experimental infection of *Apis mellifera* honeybees with *Nosema ceranae* (Microsporidia). *J Invertebr Pathol* **94**: 211–217.
- Higes, M., Martín-Hernández, R., Botías, C., Bailon, E.G., Gonzalez-Porto, A.V., Barrios, L., *et al.* (2008) How natural infection by *Nosema ceranae* causes honeybee colony collapse. *Environ Microbiol* **10**: 2659–2669.
- Higes, M., Martín-Hernández, R., Garrido-Bailón, E., Botías, C., and Meana, A. (2009a) The presence of *Nosema ceranae* (Microsporidia) in North African honey bees (*Apis mellifera intermissa*). *J Apic Res* **48**: 217–219.
- Higes, M., Martín-Hernández, R., Garrido-Bailon, E., Gonzalez-Porto, A.V., Garcia-Palencia, P., Meana, A., *et al.* (2009b) Honeybee colony collapse due to *Nosema ceranae* in professional apiaries. *Environ Microbiol Rep* **1**: 110–113.
- Higes, M., Meana, A., Bartolomé, C., Botías, C., and Martín-Hernández, R. (2013) *Nosema ceranae* (Microsporidia), a controversial 21st century honey bee pathogen. *Environ Microbiol Rep* **5**: 17–29.
- Huang, S.K., Csaki, T., Doublet, V., Dussaubat, C., Evans, J.D., Gajda, A.M., *et al.* (2014) Evaluation of cages designs and feeding regimes for honey bee in vitro experiments. *J Econ Entomol* **107**: 54–62.
- Invernizzi, C., Abud, C., Tomasco, I.H., Harriet, J., Ramallo, G., Campa, J., *et al.* (2009) Presence of *Nosema ceranae* in honeybees (*Apis mellifera*) in Uruguay. *J Invertebr Pathol* **101**: 150–153.
- Johnson, R.M., Pollock, H.S., and Berenbaum, M.R. (2009) Synergistic interactions between in-hive miticides in *Apis mellifera*. *J Econ Entomol* **102**: 474–479.
- Klee, J., Besana, A.M., Genersch, E., Gisder, S., Nanetti, A., Tam, D.Q., *et al.* (2007) Widespread dispersal of the microsporidian *Nosema ceranae*, an emergent pathogen of the western honey bee, *Apis mellifera*. *J Invertebr Pathol* **96**: 1–10.
- Kralj, J., and Fuchs, S. (2010) *Nosema* sp. influences flight behavior of infected honey bee (*Apis mellifera*) foragers. *Apidologie* **41**: 21–28.
- Le Conte, Y., Ellis, M., and Ritter, W. (2010) *Varroa* mites and honey bee health: can *Varroa* explain part of the colony losses? *Apidologie* **41**: 353–363.
- Liebig, G. (2001) How many *Varroa* mites can be tolerated by a honey bee colony? *Apidologie* **32**: 482–484.
- Maleszka, J., Barron, A.B., Helliwell, P.G., and Maleszka, R. (2009) Effect of age, behaviour and social environment on honey bee brain plasticity. *J Comp Physiol [A]* **195**: 733–740.
- Marcogliese, D.J., and Pietrock, M. (2011) Combined effects of parasites and contaminants on animal health: parasites do matter. *Trends Parasitol* **27**: 123–130.
- Martín-Hernández, R., Meana, A., Prieto, L., Salvador, A.M., Garrido-Bailon, E., and Higes, M. (2007) Outcome of colonization of *Apis mellifera* by *Nosema ceranae*. *Appl Environ Microbiol* **73**: 6331–6338.
- Martin, S.J., Hardy, J., Villalobos, E., Martín-Hernández, R., Nikaido, S., and Higes, M. (2013) Do the honeybee pathogens *Nosema ceranae* and deformed wing virus act synergistically? *Environ Microbiol Rep* **5**: 506–510.
- Mattila, H.R., and Otis, G.W. (2006) Effects of pollen availability and *Nosema* infection during the spring on division of labor and survival of worker honey bees (Hymenoptera: Apidae). *Environ Entomol* **35**: 708–717.
- Mayack, C., and Naug, D. (2009) Energetic stress in the honeybee *Apis mellifera* from *Nosema ceranae* infection. *J Invertebr Pathol* **100**: 185–188.
- Mayack, C., and Naug, D. (2010) Parasitic infection leads to decline in hemolymph sugar levels in honeybee foragers. *J Insect Physiol* **56**: 1572–1575.
- Moritz, R.F.A., and Southwick, E.E. (1992) Bees as Superorganisms: An Evolutionary Reality. Berlin Heidelberg, Germany: Springer Verlag.
- Mullin, C.A., Frazier, M., Frazier, J.L., Ashcraft, S., Simonds, R., van Engelsdorp, D., and Pettis, J.S. (2010) High levels of miticides and agrochemicals in North American apiaries: implications for honey bee health. *PLoS ONE* **5**: e9754.
- Naug, D., and Gibbs, A. (2009) Behavioral changes mediated by hunger in honeybees infected with *Nosema ceranae*. *Apidologie* **40**: 595–599.
- Neumann, P., and Carreck, N.L. (2010) Honey bee colony losses. *J Apic Res* **49**: 1–6.
- Oldroyd, B.P. (2007) What's killing American honey Bees? *PLoS Biol* **5**: 1195–1199.
- Paxton, R.J., Klee, J., Korpela, S., and Fries, I. (2007) *Nosema ceranae* has infected *Apis mellifera* in Europe since at least 1998 and may be more virulent than *Nosema apis*. *Apidologie* **38**: 558–565.
- Pettis, J.S., and Delaplane, K.S. (2010) Coordinated responses to honey bee decline in the USA. *Apidologie* **41**: 256–263.
- Pettis, J.S., van Engelsdorp, D., Johnson, J., and Dively, G. (2012) Pesticide exposure in honey bees results in increased levels of the gut pathogen *Nosema*. *Naturwissenschaften* **99**: 153–158.
- Pirk, C.W.W., Human, H., Crewe, R.M., and van Engelsdorp, D. (2014) A survey of managed honey bee colony losses in the Republic of South Africa – 2009 to 2011. *J Apic Res* **53**: 35–42.
- Pohorecka, K., Skubida, P., Miszczak, A., Semkiw, P., Sikorski, P., Zagibajlo, K., *et al.* (2012) Residues of neonicotinoid insecticides in bee collected plant materials from oilseed rape crops and their effect on bee colonies. *J Apic Sci* **56**: 115–134.

- Potts, S.G., Biesmeijer, J.C., Kremen, C., Neumann, P., Schweiger, O., and Kunin, W.E. (2010) Global pollinator declines: trends, impacts and drivers. *Trends Ecol Evol* **25**: 345–353.
- Retschnig, G., Neumann, P., and Williams, G.R. (2014a) Thiacloprid-*Nosema ceranae* interactions in honey bees: host survivorship but not parasite fecundity is dependent on pesticide dose. *J Invertebr Pathol* **118**: 18–19.
- Retschnig, G., Williams, G.R., Mehmman, M.M., Yañez, O., de Miranda, J.R., and Neumann, P. (2014b) Sex-specific differences in pathogen susceptibility in honey bees (*Apis mellifera*). *PLoS ONE* **9**: e85261.
- Rosenkranz, P., Aumeier, P., and Ziegelmann, B. (2010) Biology and control of *Varroa destructor*. *J Invertebr Pathol* **103**: 96–119.
- Sandrock, C., Tanadini, L.G., Pettis, J.S., Biesmeijer, J.C., Potts, S.G., and Neumann, P. (2014a) Sublethal neonicotinoid insecticide exposure reduces solitary bee reproductive success. *Agric Forest Entomol* **16**: 119–128.
- Sandrock, C., Tanadini, M., Tanadini, L.G., Fauser-Misslin, A., Potts, S.G., and Neumann, P. (2014b) Impact of chronic neonicotinoid exposure on honeybee colony performance and queen supersedure. *PLoS ONE* **9**: e103592.
- Schäfer, M.O., Ritter, W., Pettis, J.S., and Neumann, P. (2010) Winter losses of honeybee colonies (Hymenoptera: Apidae): the role of infestations with *Aethina tumida* (Coleoptera: Nitidulidae) and *Varroa destructor* (Parasitiformes: Varroidae). *J Econ Entomol* **103**: 10–16.
- Scheiner, R., Abramson, C.I., Brodschneider, R., Crailsheim, K., Farina, W.M., Fuchs, S., et al. (2013) Standard methods for behavioural studies of *Apis mellifera*. *J Apic Res* **52**(4): 52.4.04 <http://dx.doi.org/10.3896/IBRA.1.52.4.04>
- Schmuck, R., Stadler, T., and Schmidt, H.W. (2003) Field relevance of a synergistic effect observed in the laboratory between an EBI fungicide and a chloronicotiny insecticide in the honeybee (*Apis mellifera* L., Hymenoptera). *Pest Manag Sci* **59**: 279–286.
- Schneider, C.W., Tautz, J., Grunewald, B., and Fuchs, S. (2012a) RFID tracking of sublethal effects of two neonicotinoid insecticides on the foraging behavior of *Apis mellifera*. *PLoS ONE* **7**: e30023.
- Schneider, S., Eisenhardt, D., and Rademacher, E. (2012b) Sublethal effects of oxalic acid on *Apis mellifera* (Hymenoptera: Apidae): changes in behaviour and longevity. *Apidologie* **43**: 218–225.
- Siede, R., Faust, L., Meixner, M., Maus, C., Grünwald, B., and Büchler, R. (2014) Testing thiacloprid: does a long-term exposure affect bee colonies? Sixth European Conference of Apidologie: 22.
- Tanner, G., and Czerwenka, C. (2011) LC-MS/MS analysis of neonicotinoid insecticides in honey: methodology and residue findings in Austrian honeys. *J Agric Food Chem* **59**: 12271–12277.
- Teeters, B.S., Johnson, R.M., Ellis, M.D., and Siegfried, B.D. (2012) Using video-tracking to assess sublethal effects of pesticides on honey bees (*Apis mellifera* L.). *Environ Toxicol Chem* **31**: 1349–1354.
- Tsigouri, A.D., Menkissoglu-Spiroudi, U., and Thrasivoulou, A. (2001) Study of tau-fluvalinate persistence in honey. *Pest Manag Sci* **57**: 467–471.
- Vidau, C., Diogon, M., Aufauvre, J., Fontbonne, R., Viguès, B., Brunet, J.L., et al. (2011) Exposure to sublethal doses of fipronil and thiacloprid highly increases mortality of honeybees previously infected by *Nosema ceranae*. *PLoS ONE* **6**: e21550.
- Wang, D.I., and Mofler, F.E. (1970) The division of labor and queen attendance behavior of *Nosema*-infected worker honey bees. *J Econ Entomol* **63**: 1540–1541.
- Williams, G.R., Shafer, A.B.A., Rogers, R.E.L., Shutler, D., and Stewart, D.T. (2008) First detection of *Nosema ceranae*, a microsporidian parasite of European honey bees (*Apis mellifera*), in Canada and central USA. *J Invertebr Pathol* **97**: 189–192.
- Williams, G.R., Shutler, D., Little, C.M., Burgher-MacLellan, K.L., and Rogers, R.E.L. (2011) The microsporidian *Nosema ceranae*, the antibiotic Fumagilin-B (R), and western honey bee (*Apis mellifera*) colony strength. *Apidologie* **42**: 15–22.
- Williams, G.R., Alaux, C., Costa, C., Csáki, T., Doublet, V., Eisenhardt, D., et al. (2013) Standard methods for maintaining adult *Apis mellifera* in cages under in vitro laboratory conditions. *J Apic Res* **52**: 52.1.04. <http://dx.doi.org/10.3896/IBRA.1.52.1.04>
- Williams, G.R., Shutler, D., Burgher-MacLellan, K.L., and Rogers, R.E.L. (2014) Infra-population and -community dynamics of the parasites *Nosema apis* and *Nosema ceranae*, and consequences for honey bee (*Apis mellifera*) hosts. *PLoS ONE* **9**: e99465.
- Wu, J.Y., Anelli, C.M., and Sheppard, W.S. (2011) Sub-lethal effects of pesticide residues in brood comb on worker honey bee (*Apis mellifera*) development and longevity. *PLoS ONE* **6**: e14720.
- Yang, E.C., Chang, H.C., Wu, W.Y., and Chen, Y.W. (2012) Impaired olfactory associative behavior of honeybee workers due to contamination of imidacloprid in the larval stage. *PLoS ONE* **7**: e49472.
- Yoshiyama, M., and Kimura, K. (2011) Distribution of *Nosema ceranae* in the European honeybee, *Apis mellifera* in Japan. *J Invertebr Pathol* **106**: 263–267.