Neonicotinoids disrupt circadian rhythms and sleep in honey bees

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Honey bees are critical pollinators in ecosystems and agriculture, but their numbers have significantly declined. Declines in pollinator populations are thought to be due to multiple factors including habitat loss, climate change, increased vulnerability to disease and parasites, and pesticide use. Neonicotinoid pesticides are agonists of insect nicotinic cholinergic receptors, and sub-lethal exposures are linked to reduced honey bee hive survival. Honey bees are highly dependent on circadian clocks to regulate critical behaviors, such as foraging orientation and navigation, time-memory for food sources, sleep, and learning/memory processes. Because circadian clock neurons in insects receive light input through cholinergic signaling we tested for effects of neonicotinoids on honey bee circadian rhythms and sleep. Neonicotinoid ingestion by feeding over several days results in neonicotinoid accumulation in the bee brain, disrupts circadian rhythmicity in many individual bees, shifts the timing of behavioral circadian rhythms in bees that remain rhythmic, and impairs sleep. Neonicotinoids and light input act synergistically to disrupt bee circadian behavior, and neonicotinoids directly stimulate wake-promoting clock neurons in the fruit fly brain. Neonicotinoids disrupt honey bee circadian rhythms and sleep, likely by aberrant stimulation of clock neurons, to potentially impair honey bee navigation, time-memory, and social communication.

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Results

Honey bee (Apis mellifera ligustica) hives were maintained on the Vanderbilt University campus. Vanderbilt does not use neonicotinoid pesticides, and naive forager bees from our hives did not show detectable levels of thiamethoxam or clothianidin, the two neonicotinoids used in this study (Supplemental Fig. 1).

To study effects of neonicotinoid pesticides on individual honey bee circadian rhythms and sleep, forager honey bees were captured at hive entrances and maintained in individual infrared activity monitors in the laboratory for up to 8 days with ad libitum access to bee candy (honey and powdered sugar, with or without the addition of thiamethoxam or clothianidin)24. Neonicotinoids in the food ranged from 0 to 140 ng/g (or parts per billion, ppb), which is within the concentration range reported in flower nectar and pollen encountered by foraging bees20–23. To determine the amounts of neonicotinoids present in dosed bees, we isolated whole brains from control bees that consumed bee candy with no added neonicotinoid, and from bees that consumed bee candy dosed with thiamethoxam at two concentrations (25 ppb and 70 ppb). We collected brains from each group at 24-h intervals over 4 days and then tested for thiamethoxan and its metabolite clothianidin by mass spectrometry (see "Materials and methods"). Thiamethoxam and clothianidin were undetectable in control bee brains (Supplemental Fig. 1). Thiamethoxam was undetectable, or detected at low picogram levels, in bee brains at both doses. The active metabolite of thiamethoxam, clothianidin24,25, was readily detected at higher levels (0–220 pg/brain, Supplemental Fig. 1), consistent with the previously described rapid conversion of thiamethoxam to clothianidin in insects24,25. Forager honey bees exposed to thiamethoxam-dosed food exhibited sustained levels of clothianidin of ca. 50–70 pg/brain through day 4 at the 70 ppb dose, with lower levels detected at the 25 ppb dose (Supplemental Fig. 1).

To obtain sufficient circadian cycles for analysis, only data from control and dosed bees that survived a minimum of 4 days in light-dark (LD) experiments, or a minimum of 5 days in constant dark (DD) and constant light (LL) experiments, were included in datasets (see "Materials and methods"). For those bees meeting these criteria for inclusion, survivorship analysis indicated that there was no additional mortality of neonicotinoid-exposed bees compared to control unexposed bees at any of the neonicotinoid concentrations (25–140 ppb) used in the LD, DD, and LL datasets (N = 944, Supplemental Fig. 2). Survivorship analysis of all bees that began LD, DD and LL experiments (including those that did not survive sufficient days to contribute to the dataset) showed that there was an initial significant loss of dosed bees compared to controls primarily limited to the first day of exposure to both 70 ppb and 140 ppb levels of thiamethoxam or clothianidin in the food, followed by comparable survivorship to controls on subsequent days (Supplemental Fig. 3). Thus, our study results report from bees for which consumption of food with these levels of neonicotinoids for 4–5 days was not lethal and did not reduce survivorship compared to controls, while consumption of food with the 70 ppb and 140 ppb levels of thiamethoxam and clothianidin was lethal to some bees in the first 24 h.

We first measured entrained and free-running circadian rhythms of captured foragers for 4 days of 12:12 LD followed by 4 days of DD in the presence and absence of neonicotinoids in the food. Ingestion of neonicotinoids completely disrupted locomotor circadian rhythms in a significant portion of bees, inducing arrhythmic behavior in an increasing proportion of bees in a dose-dependent manner (Fig. 1A,B). While only a small fraction (12%) of unexposed control bees exhibited arrhythmic locomotor behavior, nearly half of thiamethoxam-dosed (46%) and clothianidin-dosed (42%) bees lost circadian behavioral rhythms following consumption of these neonicotinoids (140 ppb, Fig. 1E,F; Pearson Chi-Square test, \( p = 0.0043 \) Thiamethoxam; \( p = 0.0142 \) Clothianidin).

In addition to disrupting honey bee locomotor circadian rhythms in many individuals, neonicotinoid ingestion also dramatically altered the rhythmic behavior of the dosed bees that remained rhythmic. Individual and average actogram activity profiles show that honey bees that ingested neonicotinoids delayed termination of their active period well into the dark phase following lights off, and increased their activity at night (Fig. 1C,D,G,H). There was a more than 3-h delay in the time of activity offset in bees that consumed either thiamethoxam or clothianidin compared to undosed bees (Fig. 1J; One-Way ANOVA \( p < 0.0001 \)). This marked delay in activity offset persisted into the first day of DD in bees that consumed clothianidin (Fig. 1K, \( p = 0.0007 \)), indicating a shift in the alignment of circadian rhythms relative to the light cycle. While honey bees that consumed neonicotinoids had statistically significant increases in the proportion of locomotor activity at night, and in the overall duration of activity each day, there was no significant increase in the total activity counts, indicating that neonicotinoids induced changes in the overall temporal structure of activity, but not the total amount of daily activity (Supplemental Fig. 4). In addition, ingestion of thiamethoxam, but not clothianidin, lengthened the free-running period of honey bee locomotor rhythms measured in DD following LD (Fig. 1L; One-Way ANOVA, \( p = 0.0002 \)), demonstrating an effect on a core property of the underlying circadian clock.

Surprisingly, when honey bees were exposed to neonicotinoids solely in DD (without first being exposed during LD), the treatment effects were blunted. There was no significant increase in behavioral arrhythmicity (Fig. 2A,C) and only a trend toward lengthening of the period (Supplemental Fig. 5). This result, and the central role of cholinergic signaling in clock light input in Drosophila, suggested to us the possibility of a key interaction between light input and neonicotinoid ingestion. To examine this potential interaction further, we combined continuous background illumination (LL, 230 lx) with exposure to thiamethoxam-dosed bee candy for a period of 5 days (Fig. 2B,D). Constant illumination is an abnormal signal to the circadian clock which renders some bees arrhythmic26. We found that that LL itself disrupted rhythms in 28% of undosed bees compared to just 9% arrhythmicity in DD. When thiamethoxam was combined with LL, a dose-dependent pattern in circadian disruption emerged, such that 34% of bees exposed to 25 ppb and 46% of bees exposed to 70 ppb became arrhythmic (Fig. 2B,D), but no significant effects occurred in response to exposure to these doses of thiamethoxam in DD (Fig. 2A). The differential effects between neonicotinoid exposure in LL vs DD suggests synergy between continuous light input and neonicotinoids in altering honey bee circadian behaviors.
Light input mediated by nicotinic cholinergic neurotransmission targets the large ventral lateral neurons (L-LNαs), a specific subset of PDF-expressing clock neurons that antagonize sleep in fruit flies.27,28 We therefore tested for neonicotinoid effects on bee sleep using previously established methods for sleep analysis in flies and bees that define sleep as episodes of inactivity lasting 5 min or more.29–31 Ingestion of neonicotinoids significantly impaired sleep in honey bees (Fig. 3), with average daily sleep profiles showing that neonicotinoid-dosed bees in LD exhibited reduced sleep across both night and day (Fig. 3A,B). Overall sleep duration was reduced by up to 50% (Fig. 3C,D, One-way ANOVA: p = 0.0004 Thiamethoxam; p = 0.00067 Clothianidin), while the number of sleep bouts were reduced as much as 24% (Fig. 3E,F, One-way ANOVA: p = 0.0314 Thiamethoxam; p = 0.0549 Clothianidin). Neonicotinoids also significantly reduced sleep upon application in DD and in LL (Supplemental Fig. 6). Thus, while neonicotinoids did not change the overall levels of activity (above), they changed the temporal structure of both activity and rest, disrupting sleep.

To examine whether neonicotinoids affect PDF-expressing circadian clock neurons, we next measured the Ca²⁺ responses of Drosophila L-LN, neurons. Intact whole-brain explants expressing the GCaMP5G Ca²⁺ indicator in the PDF⁺ clock neurons were imaged for Ca²⁺-dependent changes in fluorescence intensity in the L-LN, wake-promoting neurons upon exposure to either clothianidin or vehicle. We applied clothianidin because mass spectroscopy indicated an accumulation of clothianidin, as a metabolite of thiamethoxam, in bee brains when
bees consumed thiamethoxam-dosed food (Supplemental Fig. 1). Exposure of isolated brains to clothianidin resulted in a ca. 350% increase in Ca2+ fluorescence in the l-LNv PDF+ clock neurons (ΔF/F, vehicle, −0.01 ± 0.04, Clothianidin, 3.62 ± 0.47, p < 0.0001, Fig. 4A–C). These results are consistent with findings that *Drosophila* l-LNv neurons are excited directly by nicotinic cholinergic stimulation32,33, and likely express nicotinic acetylcholine receptors that are targeted by neonicotinoids. Recent results from Tasman et al. (2020, PREPRINT) demonstrate effects of chronic ingestion of neonicotinoids on *Drosophila* clock neurons as well as circadian rhythms and sleep34.

**Discussion**

This study demonstrates that neonicotinoid ingestion disrupts circadian rhythms in exposed honey bees. In addition, neonicotinoids alter the rhythmic behavior of bees that remain rhythmic following ingestion, delaying activity offset and extending activity into the night. Neonicotinoids also impair honey bee sleep, reducing total sleep and the number of sleep bouts.

The effects of neonicotinoids on honey bee circadian rhythms are enhanced by light input. In the presence of light cycles (LD) or constant light (LL), neonicotinoids induce loss of circadian behavioral rhythms in a dose-dependent manner. However, in the absence of light cycles (DD), neonicotinoids did not significantly disrupt honey bee circadian rhythms at similar exposure levels. In the field, forager honey bees experience the daily light cycle, and thus the effects of neonicotinoids measured in LD cycles are most relevant outside the laboratory.

In *Drosophila*, the PDF+ l-LNv, wake-promoting arousal neurons receive nicotinic cholinergic input from the compound eyes28,35,36. Neonicotinoid ingestion in bees mimics the effects of experimental overstimulation of *Drosophila* l-LNv,s, extending activity into the night, increasing night-time activity, and dramatically reducing sleep38. Our results suggest that neonicotinoids aberrantly excite nicotinic cholinergic light input pathways to overstimulate homologous wake-promoting clock neurons in bees, leading to disruption and alteration of circadian rhythms and sleep. The neonicotinoid stimulation of *Drosophila* l-LNv, neurons we observed (Fig. 4), and the previously observed delay of honey bee circadian rhythms in response to PDF injection into the putative bee clock nuclei39, support this proposed mechanism. It is also notable that chronic neonicotinoid ingestion also
affects honey bee locomotor responses to light (phototaxis), and acute exposure can induce increased locomotor activity.37

Chronic ingestion of neonicotinoids resulted in three principal effects in our studies—disruption of behavioral circadian rhythms, shifts in the alignment of circadian activity with the light cycle, and impairment of sleep. Each exposure was accompanied by significant decreases in total sleep duration and number of sleep episodes. Figure 3. Neonicotinoids disrupt sleep in honey bee foragers. Four day average sleep profiles for control bees and bees exposed to either (A) thiamethoxam (Two-way RM ANOVA, Time \( p < 0.0001, **\); Dose \( p = 0.0004, ***\); Interaction \( p = 0.0015, **\)) or (B) clothianidin (Two-way RM ANOVA, Time \( p < 0.0001, ****\); Dose \( p = 0.0054, **\); Interaction \( p < 0.0001, ****\)). Significant decreases in total sleep duration for exposure to (C) thiamethoxam (One-way ANOVA \( F = 6.436; p = 0.0004, ***\); Dunnett’s multiple comparison 70 ppb \( p = 0.0038, **\); 140 ppb \( p = 0.0026, **\)) or exposure to (D) clothianidin (One-way ANOVA \( F = 4.205; p = 0.0067, **\); Dunnett’s multiple comparison 140 ppb \( p = 0.0115, *\)). In addition, there were significant decreases in the number of sleep episodes for exposure to (E) thiamethoxam (One-way ANOVA \( F = 3.020; p = 0.0314*, \) Dunnett’s multiple comparison 140 ppb \( p < 0.0233, *\)) and for (F) clothianidin (One-way ANOVA \( F = 2.586; p = 0.0549, ns\); Dunnett’s multiple comparison 140 ppb \( p < 0.0396, *\)) dosed bees. Significant values only are shown for Dunnett’s post hoc multiple comparisons (C–F).
of these effects on the honey bee circadian system have the potential for deleterious impacts on critical honey bee foraging behaviors which could ultimately impact hive health and survival. The abrogation of circadian rhythms that we found in significant proportions of bees that ingested neonicotinoids, could be expected to disrupt forager time-compensated sun-compass navigation, as it does in butterflies where the critical clocks are in the antennae\textsuperscript{38}. Compromised circadian rhythms would also be expected to disrupt circadian clock-dependent time-memory for food source availability (Zeitgedächtnis). Each of these effects would significantly decrease the efficiency of honey bee foraging.

Ingestion of neonicotinoids delayed activity offset in LD, and this delay persisted when bees were then transitioned into DD, suggesting that neonicotinoids alter the timing of the honey circadian clock. The timing of activity offset relative to lights off has been shown to be a key marker of the entrainment, of honey bee circadian rhythms with the daily light cycle\textsuperscript{39}, which is a dominant cue defining local environmental time. Thus, shifts in the alignment of circadian locomotor rhythm relative to the daily light cycle, as we observed in honey bees that remained rhythmic following neonicotinoid ingestion, would be expected to induce navigational errors due to changes in alignment of clock-based time-compensation for sun-compass orientation\textsuperscript{16,40}. Neonicotinoid consumption induced ca. 3-h delays in locomotor activity alignment of dosed bees compared with the light cycle. Shifts of this magnitude are predicted to engender large navigational errors due to changes in alignment of clock-based time-compensation for sun-compass orientation\textsuperscript{16,40}. Neonicotinoid consumption induced ca. 3-h delays in locomotor activity alignment of dosed bees compared with the light cycle. Shifts of this magnitude are predicted to engender large navigational errors due to changes in alignment of clock-based time-compensation for sun-compass orientation\textsuperscript{16,40}.

Interestingly, catch-move-release experiments following chronic neonicotinoid exposures showed that many of the neonicotinoid-exposed bees became lost and did not return to their hive due to mis-orientation of initial
Locomotor activity monitoring experiments. Foragers were caught outside the hive entrance upon their return from foraging trips, confirmed by the presence of pollen on their hind legs. Foragers were individually housed in plastic tubes containing either dosed bee candy (ground white cane sugar and honey) containing a specified amount of thiamethoxam, clothianidin, or control bee candy with no pesticide. Tubes were loaded into Locomotor Activity Monitors (LAMs, Trikinetics) and housed in an environmental chamber with a controlled light:dark cycle, constant temperature of 35 °C and humidity of 60%. A filter paper wick supplied water from a reservoir to each tube. Data recording (infrared beam interruptions) began immediately after loading the monitors, and data beginning with the first full day of activity was used for analysis. Experiments of 12:12 light:dark cycle (LD) for 4 days, followed by 4 days of constant darkness (DD), took place in the spring/summer of 2017. Experiments of 250 lx constant light (LL) and constant darkness (DD) were performed during the spring/summer of 2019. In the case of the LL experiments, bees were placed into the incubator during the daytime and lights were not turned off for the duration of the experiment. For DD, foragers were placed in the incubator in the daytime and lights were turned off at 19:00 CDT that day and not turned back on for the duration of the experiment. For the LD experiments, locomotor activity rhythms and sleep data were analyzed from bees that survived in the activity monitors a minimum of 4 days, and up to 8 days of data were used. For the DD and LL experiments only bees that survived 5 full days of the experiment were used for data analysis.

Behavioral metrics (data analysis). Locomotor activity rhythms are displayed as double plotted actograms (Fig. 1A–D). Free-running period calculations were calculated using autocorrelation analysis (Fig. 1L, Supplementary Fig. 5) generated using the previously published MATLAB toolboxes. Individuals were scored as rhythmic or arrhythmic by a combination of eye inspection of each activity record, the absence of a significant peak in chi-square periodogram, and evaluation of the waveform (showing a significant and cyclical pattern) of the correlogram produced by the autocorrelation analysis (Fig. 1E,F, Fig. 2). The individual activity onset and offsets for each day were obtained using ClockLab (Actimetrics) with manual corrections as needed (Fig. 1I–L). Average activity duration was calculated by subtracting the onsets from the offsets of each day (offset time–onset time) and then averaging (Supplemental Fig. 4A,B). The proportion of activity occurring in the dark was calculated by dividing the number of infrared beam breaks recorded throughout the dark phase of a particular day

Materials and methods
Honey bees. Honey bees (Apis mellifera ligustica) were maintained in standard Langstroth 10-frame boxes (Kelly Beekeeping, Clarkson, KY) on the Vanderbilt campus. All bees used for experiments came from healthy colonies with a mated queen. Vanderbilt does not use neonicotinoid pesticides on its property, and forager honey bees (Kelly Beekeeping, Clarkson, KY) on the Vanderbilt campus. All bees used for experiments came from healthy colonies with a mated queen. Vanderbilt does not use neonicotinoid pesticides on its property, and forager honey bees from our hives did not show detectable baseline levels of thiamethoxam or clothianidin, the two neonicotinoids that we used in our study (Supplemental Fig. 1).

Our results show dramatic impacts of neonicotinoids on circadian rhythms and sleep in honey bees. However, two limitations of our studies stem from the laboratory setting in which they were performed—(1) experiments were conducted over a relatively short study period (up to about 1 week), and (2) we measured the circadian rhythms of individuals, not entire hives. The relatively short duration of these experiments may actually underestimate the impact of long-term environmental neonicotinoids on the honey bee circadian system, as in the field exposures may be more prolonged and may include developmental exposures as well. It would be of interest in the future to measure circadian aspects of whole honey bee colony behavior in the field experiments such as studies that have demonstrated reduced hive productivity and whole hive survival due to exposure to neonicotinoids.

Overall, we find that chronic ingestion of neonicotinoids disrupts circadian behavioral rhythms and sleep in forager honey bees, and that the predicted downstream neurobehavioral impacts of disrupted rhythms and sleep are potentially deleterious to hive health. Future studies are needed to identify and characterize the fundamental neural substrates that generate circadian and sleep behaviors in honey bees, and that synchronize circadian rhythms to local time through the daily light cycle. The cholinergic signaling neural circuits by which light and neonicotinoids impact honey bee locomotor rhythms also need to be identified. With the future development of additional experimental tools for honey bee neuroscience, new lines of research may elucidate the mechanism behind the interaction between pesticides and light signaling. The understanding of these pathways would open the way to elucidating the mechanisms by which neonicotinoids impact the circadian rhythms of honey bees, and suggest possible means to limit their disruptive impact.
by the number of beam breaks recorded throughout both the light and dark phases of that day. For each bee, all
days were then averaged and these values grouped by treatment (Supplemental Fig. 4C,D). Total activity counts
for LD portion of the experiment shown as total activity counts over 4 days of light: dark cycle (Supplemental
Fig. 4E,F).

Sleep was analyzed from the activity data collected from the LAMs. Several studies have shown that lack
of movement (beam interruptions) for five consecutive minutes serves as a reliable measure for sleep in honey
bees29,45,46. Consistent with these studies we defined sleep as 5 min or longer bouts of inactivity. Data was ana-
yzed using a custom toolbox for MATLAB that was originally developed to measure sleep in flies31. For each
individual, we calculated sleep in 30 min bins, total sleep duration throughout the day, and the number of average
sleep episodes in 24 h. (Fig. 3, Supplemental Fig. 6).

**LC–MS/MS analysis.** Sample analyses were carried out by the Vanderbilt Mass Spectrometry Core Lab
using a *Vanquish* ultrahigh performance liquid chromatography (UHPLC) system interfaced to a *Q Exactive*
HF quadrupole/orbitrap mass spectrometer (Thermo Fisher Scientific) equipped with an Ion Max API source, a
HESI-II electrospray probe, and a 50 um ID stainless steel capillary. Chromatographic separation was per-
formed with a reverse-phase Acquity HSS C18 column (1.7 μm, 2.1 × 50 mm, Waters, Milford, MA) at a flow rate
of 300 μL/min. Mobile phases were made up of (A) 0.2% HCOOH in H₂O and (B) 0.2% HCOOH in CH₃CN.
Gradient conditions were as follows: 0–1 min, B = 0%; 1–6 min, B = 0–100%; 6–6.5 min, B = 100%; 6.5–7.0 min,
B = 100–0%; 7.0–10 min, B = 0%. The total chromatographic run time was 10 min; the sample injection volume
was 10 μL. A software-controlled divert valve was used to transfer eluent from 0 to 1.5 min and from 5 to 10 min
of each chromatographic run to waste. The instrument was calibrated weekly over a mass range of m/z 195 to
m/z 1822 with a mixture of caffeine, MRFA, and Ultramark-1621 using the manufacturer’s HF-X tuning soft-
ware. The mass spectrometer was operated in positive ion mode. The following optimized source parameters
were used for the detection of analyte and internal standard: N₂ sheath gas 40 psi; N₂ auxiliary gas 10 psi; spray
voltage 5.0 kV; capillary temperature 320 °C; vaporizer temperature 100 °C. Quantitation was based on parallel
reaction monitoring (PRM) using a precursor isolation width of 2.0 m/z and the following MS/MS transitions.

| Analyte          | Precursor mass (m/z) | Product mass (m/z) | HCD collision energy | S-lens RF level |
|------------------|----------------------|-------------------|----------------------|-----------------|
| Clothianidin     | 250.0160             | 169.05450         | 10                   | 10              |
| Clothianidin-d₃  | 253.0348             | 172.0733          | 10                   | 10              |
| Thiamethoxam     | 292.0255             | 211.06510         | 10                   | 10              |
| Thiamethoxam-d₃  | 295.0454             | 214.0839          | 10                   | 10              |

HCD product ion spectra were acquired in the Orbitrap at a resolving power of 30,000, an AGC target of
2e5, and a maximum injection time of 100 ms. Data acquisition and quantitative spectral analysis were done
using Xcalibur version 2.2 sp1 and LCQuan version 2.7.0, respectively. Calibration curves were constructed for
clothianidin and thiamethoxam by plotting peak area ratios (analyte / internal standard) against analyte con-
centrations for a series of seven calibrants, ranging in concentration from 2 pg to 1 ng. A weighting factor of 1/
C² was applied in the linear least-squares regression analysis to maintain homogeneity of variance across the
concentration range (% error ≤ 20% for at least four out of every five standards). Foragers were exposed to either
0 ng/g, 25 ng/g or 70 ng/g of thiamethoxam and kept in individual chambers for five days. Individuals from each
group were taken at 2 h, 26 h, 50 h, 74 h, and 98 h after exposure to the pesticide food and flash-frozen in liquid
nitrogen and kept at −80 °C until processed. Brains were dissected in 100% ETOH chilled with dry ice. Each
brain was homogenized in 5% 1:1 MeOH:H₂O solution, acetonitrile. Internal standards for d₃-thiamethoxam
and d₃-clothianidin were added at a concentration of 250 nM in solution. Homogenate was spun at 3000 rpm for
3 min after which supernatant was decanted and dried. Dried samples were stored at −20 °C until processed for
LC–MS. The dried residue was reconstituted in 100 μL of H₂O/CH₃OH (3:1) vigorously vortexed, and transferred
to 200-μL silanized autosampler vials equipped with Teflon-lined bonded rubber septa.

**Calcium imaging.** Calcium imaging was performed as previously described47. Adult male flies (4–7 days
post-eclosion) from the cross PDF-Gal4 > 20XUAS-IVS-GCamp5G 41 were immobilized on 300 mm petri
dishes in 3 mL physiological saline containing 128 mM NaCl, 2 mM KCl, 4 mM MgCl₂, 35.5 mM sucrose, 5 mM
HEPES, and 1.8 mM Ca²⁺, pH 7.2. Labeled PDF soma were identified under halogen lamp illumination and
allowed to rest for 2–3 min to photo bleach soma prior to baseline recording. 1-LN, neurons we distinguished
by soma size and position. Soma were imaged with a 40 × water-immersion objective with maximal pinhole
aperture on a Zeiss LSM510Meta laser-scanning confocal microscope, using 256 × 256 resolution and a region
of interest (ROI) box to achieve ~186 ms scans. Baseline fluorescence was recorded for 250 frames, followed by
direct application of given drug, with a main recording for 1,000 total frames. To confirm that explants were
viable even when no response to treatment/vehicle was observed, 50 μL of 2.5 M KCl was applied after 1,000
frames to confirm a normal calcium response to high potassium.
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**Author contributions**

M.C.T. and D.G.M. conceived of the project. M.A.G.G. and M.C.T. performed pesticide/honey bee behavioral experiments. M.A.G.G., E.S.P. and M.C.T. performed honey bee LL experiments, DD experiments. M.A.G.G., M.C.T., E.S.P. analyzed the data. C.A.D, M.C.T., and M.A.G.G., performed *Drosophila* Ca²⁺ imaging experiments. M.C.T., M.A.G.G., C.A.D., D.M., and D.G.M. designed experiments. D.G.M., M.C.T. and M.A.G.G. wrote the paper. J.A.R., K.B., and D.M. provided editing, comments, and experimental guidance.

**Competing interests**

The authors declare no competing interests.

**Additional information**

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