Circadian regulation of olfaction and an evolutionarily conserved, nontranscriptional marker in Caenorhabditis elegans

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Edited by J. Woodland Hastings, Harvard University, Cambridge, MA, and approved November 2, 2012 (received for review July 9, 2012)

Circadian clocks provide a temporal structure to processes from gene expression to behavior in organisms from all phyla. Most clocks are synchronized to the environment by alternations of light and dark. However, many organisms experience only muted daily environmental cycles due to their lightless spatial niches (e.g., caves or soil). This has led to speculation that they may dispense with the daily clock. However, recent reports contradict this notion, showing various behavioral and molecular rhythms in Caenorhabditis elegans in and blind cave fish. Based on the ecology of nematodes, we applied low-amplitude temperature cycles to synchronize populations of animals through development. This entrainment regime reveals rhythms on multiple levels: in olfactory cued behavior, in RNA and protein abundance, and in the oxidation state of a broadly conserved peroxiredoxin protein. Our work links the nematode clock with that of other clock model systems; it also emphasizes the importance of daily rhythms in sensory functions that are likely to impact on organism fitness and population structure.

Daily cycles of light and dark are at once highly predictable and extremely stressful to living organisms. Life has thus evolved numerous strategies to use the physical properties of light—for example, to maintain a distinct orientation in space (vision) and time (the circadian clock). Circadian biological clocks regulate processes from gene expression to behavior via a complex network—from molecule to cell to organism to population—that requires exquisite coordination between all levels such that events occur at an optimal time (1–3). At the most basic, molecular level, many components that regulate daily timing are known (so-called clock genes), functioning as a transcriptional negative feedback loop. As more components are identified, forming additional and interlocked regulatory loops, the model of the molecular clock has become one of a complex molecular network (4, 5). In theoretical terms, the molecular network could be viewed as similar to the network that is formed by cells and organs with respect to their daily oscillations (3).

We remain ill-equipped to probe the system as a network—formed by cells and organs with respect to their daily oscillations (3). Circadian regulation of olfaction and an evolutionarily conserved, nontranscriptional marker in Caenorhabditis elegans

The authors declare no conflict of interest.

Author contributions: M.O. and M.M. designed research; M.O., J.S.O., R.S.E., and U.K.V. performed research; A.B.R. contributed new reagents/analytic tools; M.O. and M.M. analyzed data; and M.O. and M.M. wrote the paper.

This article is a PNAS Direct Submission.

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This article contains supporting information online at www.pnas.org/lookup/suppl/doi:10.1073/pnas.1211705109/-/DCSupplemental.

www.pnas.org/cgi/doi/10.1073/pnas.1211705109

PNAS | December 11, 2012 | vol. 109 | no. 50 | 20479–20484
temperature cycle, as if in an entrainment situation, or they were released to constant conditions (as shown in Fig. 1, start a t 120 h) to reveal the circadian property called free-running rhythm, whereby daily oscillations persist in the absence of, for example, light or temperature cycles (24).

We first tested if our protocol generated populations with synchronous, free-running circadian rhythms, by measuring RNA levels from nematodes harvested over 2 d after release into constant temperature (13 °C). There exists a putative clock gene ortholog in C. elegans, namely lin-42 (F47F6.1). To date, lin-42 mRNA has not been found to be expressed according to a circadian rhythm; rather, its expression correlates with developmental stages such as larval transitions (25, 26). In agreement with earlier work, we also found that this RNA species is expressed constitutively in adults, even after our synchronization protocol (Fig. 1 and Fig. S1). We then investigated expression of transcripts that had been reported as rhythmic in protocols that used higher amplitude temperature cycles (11). Although the expression of many of these genes was dynamic (decreasing or increasing on release to constant conditions), only one of nine (B0507.8) (11) was significantly rhythmic (Fig. 1C and Table S1), suggesting that this low-amplitude temperature cycle effectively entrains populations of C. elegans and that they have a free-running, circadian rhythm at the level of RNA transcription.

To extend our repertoire of rhythmic outputs, we asked if the redox state of PRX follows a circadian cycle. Recent reports show this protein, involved in hydrogen peroxide metabolism, oscillating between reduced and oxidized forms in organisms from all phyla, even in nonnucleated cells (27, 28). C. elegans expresses at least two typical 2-Cys PRX genes, prdx-2 (F09E5.15) and prdx-3 (R07E5.2). The PRX active site is highly conserved in these proteins (Fig. S2), and an antiserum raised against the oxidized peptide DFTFVC*PTEI detects predominantly PRDX-2 (Fig. S3). Protein extracts from animals released into constant conditions from temperature cycles

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**Fig. 1.** A protocol for supporting circadian rhythms in C. elegans. (A) Egg preps were inoculated onto a lawn of E. coli (arrow). The plates were placed in incubators that were programmed for temperature cycles of 16 h at 13 °C (shown in blue) and 8 h at 16 °C (shown in pink). On the sixth day, the plates were released to constant temperature (13 °C), unless otherwise indicated, and the experiments were started. Time point 120 h corresponds to the starting point for all of the experiments (time = 0 h). The pictures show the developmental stage of C. elegans at the end of each 24 h cycle. (B) The mRNA of the putative clock gene ortholog, lin-42, is not rhythmic in our cultivation protocol. lin-42 mRNA levels were measured by quantitative RT-PCR, and the results were normalized to the amount of act-4 mRNA and then to the sample with the highest expression level. The samples were collected for 48 h in constant conditions starting at 120 h after inoculation (red box in panel A). We express the timing of the harvesting protocol (x axis) as time from the end of the warm to cold transition at 120 h. The plot shows the average ± SEM of four biological replicates. (C) Quantitative RT-PCR was used to verify that the protocol supports circadian rhythms. The levels of mRNA corresponding to several transcripts, rhythmic under a different protocol (11), are shown. The plots show the mean ± SEM of three biological replicates. A sinusoidal curve was fitted to the data for B0507.8 using Circwave (P < 0.01). Additional statistical analysis with Circwave and JTK-Cycle is shown in Table S1. The results were normalized and x axes are labeled as for panel B.
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mRNA levels are not rhythmic under the same protocol used for protein
sample with the highest expression level. The statistical analysis with Circwave and JTK-Cycle is shown in Table S1.

Representative immunoblot probed for over-/hyper-oxidized 2-Cys PRX (PRX-SO2/3). β-warm to cold. Loading control shows an image of the same blot probed for conditions (13 °C, darkness). Time point 0 h corresponds to the release from darkness), the oxidation of PRDX-2 shows a circadian rhythm in free-running

After entrainment to a temperature cycle (16 h at 13 °C and 8 h at 16 °C, sinusoidal curve was served molecular marker of the circadian clock in

Olfaction, for example, is regulated by the circadian
robust behaviors described in this nematode (30)
C. elegans. Olfaction is typically assessed in
genic food choices (34). Olfaction is measured using a chemotaxis assay, whereby the animals are harvested
during

We applied this method to assay the aversive response to the chemical 1-octanol over the course of a day at constant temperature (Fig. 3A). The animals showed lower responses in the subjective

Fig. 2. Oxidation state of PRX in constant conditions in C. elegans. (A) Representative immunoblot probed for over-/hyper-oxidized 2-Cys PRX (PRX-SO2/3). After entrainment to a temperature cycle (16 h at 13 °C and 8 h at 16 °C, darkness), the oxidation of PRDX-2 shows a circadian rhythm in free-running conditions (13 °C, darkness). Time point 0 h corresponds to the release from warm to cold. Loading control shows an image of the same blot probed for β-actin. For each time series, the data were quantified by densitometry and normalized to the highest value within each blot. The x axes are labeled as described in Fig. 18. The mean ± SEM for three biological replicates is shown. A sinusoidal curve was fitted to the data using Circwave (P < 0.01). Additional statistical analysis with Circwave and JTK-Cycle is shown in Table S1. (B) prdx-2 mRNA levels are not rhythmic under the same protocol used for protein quantification. The expression of prdx-2 was measured by quantitative RT-PCR. The results were normalized to the amount of act-4 mRNA and then to the sample with the highest expression level. The x axes are labeled as described in Fig. 18. The plot shows the mean ± SEM of three biological replicates.

showed statistically significant rhythms in this form of PRDX-2 (Fig. 24 and Table S1). This rhythm represents a phylogenetically conserved molecular marker of the circadian clock in C. elegans. Interestingly, the expression of prdx-2 mRNA is not clock regulated (Fig. 2).

Given that we find daily rhythms in RNA and metabolism, we wanted to know if the clock is regulating quantifiable behaviors. In seemingly all circadian systems, sensory functions are clock-regulated. Olfaction, for example, is regulated by the circadian clock in many animals (19–21, 29). Indeed, it is one of the most robust behaviors described in this nematode (30–33) and is a key component of fitness, as it guides between nutritious and pathogenic food choices (34). Olfaction is typically assessed in C. elegans with a chemotaxis assay, whereby the animals are harvested from their culture plates and washed several times to remove the interfering effects of Escherichia coli. We developed a unique, quantitative, in situ assay for olfaction so that we could measure olfaction without perturbing the animals, thus avoiding possible masking (35) that might occur due to the washing step. Eggs were inoculated on a small drop of food (Fig. S4A), and as described above, the developing animals were submitted to temperature cycles for 5 d (Fig. 1A). The olfaction assay was initiated by applying a drop of 1-octanol (a chemorepellant for C. elegans) to one side of the food source. This odorant was chosen because it was the only compound for which chemotaxis could be demonstrated in the presence of E. coli among many odorants that were tested (Table S2). At the end of the chemotaxis assay, a picture was taken showing the animals as they were distributed over the lawn of bacteria. Each picture was divided into a proximal and a distal half, and nematodes were counted to yield a Chemotaxis Index (CI; Fig. S4A). The in situ assay was optimized for quantitative studies by comparing a dilution series of the odorant over assay end points from 5 to 60 min (Fig. S4B). We determined that an end point of 15 min in combination with a 1/27 dilution of 1-octanol is optimal.

We applied this method to assay the aversive response to the chemical 1-octanol over the course of a day at constant temperature (Fig. 3A). The animals showed lower responses in the subjective

Fig. 3. Circadian modulation of olfaction in response to 1-octanol. (A) Response to 1-octanol over 24 h in constant conditions. Blue panels represent cool temperature (13 °C); pink panels represent warm temperature (16 °C). After 5 d of growth in a temperature cycle (13 °C to 16 °C), the response to 1-octanol was measured for 24 h at constant temperature, 13 °C. The data are plotted as the percentage deviation from the average CI of the experiment. The x axes are labeled as described in Fig. 18. The values for each time point are shown as the mean ± SEM calculated from 5 to 10 plates in three independent experiments. The average CIs for the three experiments are 0.13, 0.31, and 0.49. A sinusoidal curve was fitted to the data (P < 0.001) using Circwave. (B) Response to 1-octanol over 24 h in entraining conditions (a temperature cycle). Blue panels represent cool temperature (13 °C); pink panels represent warm temperature (16 °C). After 5 d of growth in a temperature cycle, chemotaxis was measured over 24 h at the same cycling conditions. The data are plotted as the percentage deviation from the average CI of the experiment. The x axes are labeled as described in Fig. 18. The values for each time point are shown as the mean ± SEM calculated from 5 to 10 plates in three independent experiments. The average CIs for the three experiments are 0.19, 0.40, and 0.59. For a plot of the CI before normalization, see Fig. S5.
night compared with the subjective day, yielding a fitted sine curve with high significance ($P < 0.001$). Interestingly, when the animals were assayed for chemotaxis under cycling temperature conditions, although they showed a similar pattern of increased responsiveness during the day, this increase in olfaction was extended longer and later than in constant conditions (Fig. 3B). This suggests that the increase in temperature itself induces a masking effect or possibly that the chemical is more volatile at the higher temperature. Importantly, the chemotaxis increased before the onset of warm temperature, showing anticipation of the zeitgeber transition, a feature that is expected under appropriately structured circadian entrainment conditions.

Another feature of circadian clocks is how they behave in non-24-h entraining cycles. For instance, in cycles that are slightly shorter than 24 h, a circadian program would be expected to entrain later, whereas a masked or driven rhythm should show similar phase reference points relative to zeitgeber transitions. The imposition of a 23 h temperature cycle on the nematodes perturbs the entrainment such that—on release to constant conditions—a later phase of 1-octanol olfaction is observed on the first day in constant conditions (Fig. S6). The data from entrainment (in two different cycle lengths) and release to a free run suggest the presence of an underlying circadian oscillator regulating olfaction.

Much is known concerning the molecular aspects of olfaction. For instance, odorant receptors are a large family of G protein-coupled receptors (GPCRs) (36) [or highly homologous to these (37)]. GPCRs, in turn, are often regulated by G protein-coupled receptor kinases (GRKs) and arrestins that facilitate desensitization of these molecules in mammals (38). The genome of *C. elegans* contains two predicted GRKs, grk-1 (F19C6.1) and grk-2 (W02B3.2). GRK-2 regulates the response to 1-octanol in *C. elegans*, most likely through the phosphorylation of the correspondent GPCR (39), although the gene encoding the receptor for 1-octanol in *C. elegans* remains unknown. Furthermore, nematodes lacking GRK-2 function show disrupted chemosensation, including responses to 1-octanol (40) due to the activation of inhibitory pathways that dampen signaling in the absence of GRK-2 (41). 1-octanol is octanol (40) due to the activation of inhibitory pathways that

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**Discussion**

Circadian clocks confer a fitness advantage in the cycling light environment (43), presumably via the temporal structure that they create for processes ranging from gene expression to complex behavior such as sleep. The adaptive advantage of the clock in environments with neither light nor high-amplitude environmental cycles (such as in soil) is less obvious. Despite this, daily rhythms in blind cavefish were recently reported (44). The circadian system of the cavefish *Phreatichthys andruzii* is not entrained by light/dark cycles but rather to feeding time (once per 24 h), showing a characteristic increase in activity in anticipation of food. In the context of such a feeding protocol, rhythms in gene expression were also observed. In constant conditions, gene expression of this light-blind oscillator shows a very long free-running rhythm (47 h).

Over the last decade, there have been numerous reports of daily rhythms in the soil-dwelling nematode, *C. elegans* (6–12), although the connection between molecules and behavior has been lacking. Reasons for this may include a lack of known clock gene homologs as starting points, a number of low-amplitude behavioral rhythms, and the use of conditions that are far from ecological. We noted that although *C. elegans* have no eyes, they retain some capability to sense light acutely (45–47). However, because they also have no pigmentation, we adopted the hypothesis that they do not regularly encounter light and hence would not use light to entrain their circadian clock. Here, we have developed a protocol using temperature as a zeitgeber that entrains the clock through development. Evidence of this synchronization is shown here as rhythms in molecules and behavior upon release to constant conditions. In contrast to the cavefish (44), the period of the nematode circadian rhythm is close to 24 h (Figs. 1C, 2A, 3A, and 4A and Table S1).
Quantitative Real-Time PCR (qPCR)  
Nematodes were collected from the plates every 4 h for 2 d in constant conditions (13 °C, darkness) using cold M9 buffer, centrifuged, and washed once to remove bacteria. Protein was isolated using TRIzol (Invitrogen), and cDNA synthesis was performed (30 min at 48 °C) using Random Hexamers and MultiScribe Reverse Transcriptase (Applied Biosystems). Quantitative PCR analyses (SYBR Green PCR Master Mix, Applied Biosystems) were performed using a 7500 Real Time PCR System thermal cycler (Applied Biosystems). The reaction included denaturation (10 min at 95 °C) and 40 PCR cycles (15 s at 95 °C and 1 min at 60 °C). The results for each gene were normalized to the corresponding results obtained with act-4 by the ΔΔct method and then to the sample with the highest RNA level. Primers were designed using Primer Express Software (Applied Biosystems) (Table S3). Circadian rhythmicity was evaluated for each gene, and a sinusaloid curve was fitted to the data for B0507.8 using Circave (by R. Hut, available at www.eucock.org).

In Situ Chemotaxis Assay. For the 1-octanol assay, NGM plates were inoculated with 15 μL of a concentrated E. coli culture (100 g wet weight/L) and allowed to dry overnight at room temperature. To start the experiment, about 100 eggs were placed in the E. coli drop, and the worms were grown for 5 d in temperature cycles (13 °C to 16 °C, as in Fig. 1A) in constant dim light. The chemotaxis assay was performed either during the sixth day in a temperature cycle or on the sixth day following release to constant temperature. Every hour, chemotaxis assays were performed on naïve plates by placing 1 μL of 1-octanol [3.7% (vol/vol) in ethanol] at a distance of 0.8 cm from the center of the E. coli drop. The plates were incubated for 15 min, at which point a picture was taken using a light microscope (10×, Stereo Discovery V8, Zeiss). The CI was calculated for each time point and each plate (Fig. S4). A sinusaloid curve was fitted to the data using Circave (by R. Hut, available at www.eucock.org).

Protein Preparation. Nematodes were collected using cold M9 buffer, centrifuged, and washed one time to remove bacteria. Proteins were extracted in Nonidet P-40 protein lysis buffer (150 mM NaCl, 1% Nonidet P-40, 50 mM Tris pH 8.0) containing protease inhibitors (Complete Mini, Roche) and allowed to dry overnight at room temperature. To start the experiment, about 100 eggs were placed in the E. coli drop, and the worms were grown for 5 d in temperature cycles (13 °C to 16 °C, as in Fig. 1A) in constant dim light. The chemotaxis assay was performed either during the sixth day in a temperature cycle or on the sixth day following release to constant temperature. Every hour, chemotaxis assays were performed on naïve plates by placing 1 μL of 1-octanol [3.7% (vol/vol) in ethanol] at a distance of 0.8 cm from the center of the E. coli drop. The plates were incubated for 15 min, at which point a picture was taken using a light microscope (10×, Stereo Discovery V8, Zeiss). The CI was calculated for each time point and each plate (Fig. S4). A sinusaloid curve was fitted to the data using Circave (by R. Hut, available at www.eucock.org).

Materials and Methods

Strains and Experimental Conditions. The N2 wild-type strain of C. elegans was used for all circadian experiments. The mutant strains VC289 and SO2/3 (Abcam, ab16830) was used as an earlier version of this manuscript; P. Hardin for helpful suggestions; T. Roenneberg (Ludwig-Maximilians-Universität-München), J. Hogeness (University of Pennsylvania), and R. Hut (University of Groningen) for input on data analysis; and the Caenorhabditis Genetics Center supported by the National Institutes of Health for supplying animals. Our work is supported by the European Commission (Euclock, 6th Framework Programme Integrated Project), the Netherlands Organization for Scientific Research (Dutch Science Foundation, Vici, and Open Programme), and the Rosalind Franklin Fellowships of the University of Groningen. A.B.R. is supported by the Wellcome Trust (083643/2/).
07/2), the European Research Council (ERC) (Starting Grant 281348, Meta-


Supporting Information

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Fig. S1. Expression of the heterochronic gene *lin-42* through development in our temperature entrainment protocol. Experiments represented in all other figures in this article were performed within the time window of 120–168 h after inoculation of the eggs, when *lin-42* is expressed at low levels despite an ongoing temperature cycle. Blue panels represent 13 °C, and pink panels represent 16 °C.

Fig. S2. Multiple sequence alignment showing PRX amino acid sequences. The highly conserved active site is underlined. The sequences analyzed correspond to *At* (*Arabidopsis thaliana*; NP_187769.1), *Se* (*Synechococcus elongatus* PCC 7942; YP_401326.1), *Hs* (*Homo sapiens*; NP_005800.3), *Mm* (*Mus musculus*; NP_035693.3), *Ce2* (*C. elegans*; NP_001122604.1; *prdx-2*), *Dm* (*Drosophila melanogaster*; NP_477510.1), *Ce3* (*C. elegans*; NP_497892.1; *prdx-3*), and *Nc* (*Neurospora crassa*; XP_959621.1).

Fig. S3. Antiserum against PRX-SO\(_{2/3}\) recognizes the oxidized form of PRDX-2. Wild-type (N2) and mutant *prdx-2* (VC289) and *prdx-3* (VC1151) worms were treated with 1 mM H\(_2\)O\(_2\) for 30 min (to induce expression of the PRX proteins) and then lysed for immoblotting. The immunoblot was probed for PRX-SO\(_{2/3}\). The antiserum raised against the oxidized peptide DFTFVCPTEI detects both PRDX-2 and PRDX-3 in wild-type *C. elegans* (N2), indicated by a doublet with H\(_2\)O\(_2\) treatment. The lower band detected by the antiserum is absent in the mutant *prdx-2* and present in the *prdx-3* mutant, indicating that PRDX-2 is the dominant *C. elegans* ortholog of PRX that is detected in the time-course assays.
Fig. S4. (A) An in situ assay for chemotaxis: population assay in olfaction using 1-octanol. Approximately 100 nematodes were inoculated onto a drop of *E. coli* as eggs. They were then subjected to the development protocol with temperature cycles (Fig. 1A). On day 6, olfaction assays were started by placing a 1 μL drop of 1-octanol (shown as a black drop) to one side of the *E. coli*. After a given amount of time, a picture was taken and the CI was determined by counting the animals on the proximal (p) and distal (d) sides of the drop of bacteria. The difference (d – p) was divided by the total number of animals. (B) Optimization of the chemotaxis assay. The assay was optimized by comparing different time points after addition of 1-octanol (between 5 and 60 min) and dilutions of the chemorepellant (none to 1/243 in ethanol).

Fig. S5. CI from the three experiments in Fig. 3B are plotted independently. A sinewave was fitted to each time series using Circwave.
Fig. S6. Response to 1-octanol in constant conditions after entrainment to either a 24 h cycle (Upper) or a 23 h cycle (Lower). One complete cycle is in both cases represented as 360°. A sinewave was fitted to each series using Circwave. The acrophase of the sinewave adjusted to the T = 24 series (with a $P < 0.001$) is 225.45°, and the acrophase of the sinewave adjusted to the T = 23 series (with a $P = 0.0019$) is 283.35°.

Table S1. Period estimates and statistical parameters calculated by Circwave and JTK_Cycle

<table>
<thead>
<tr>
<th>Molecular marker</th>
<th>Circwave</th>
<th>JTK_Cycle</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Tau</td>
<td>$F$ statistic</td>
</tr>
<tr>
<td>B0507.8</td>
<td>22.6</td>
<td>7.7037</td>
</tr>
<tr>
<td>PRX-SO2/3</td>
<td>24.5</td>
<td>6.008</td>
</tr>
<tr>
<td>GRK-2</td>
<td>31.3</td>
<td>18.9573</td>
</tr>
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</table>

Circwave is an analysis tool for determining circadian profiles and their significance using harmonic regression in combination with the $F$-test statistic. A fundamental sinusoidal wave is fitted through the data, and its significance is tested against a fitted horizontal line through the overall average (www.euclock.org). JTK_CYCLE is a nonparametric statistical algorithm designed to identify and characterize cycling variables. JTK_CYCLE provides optimal phase, amplitude, and period estimates for each variable, and permutation-based $P$ values (1).

Table S2. List of volatile odorants tested in the in situ chemotaxis assay

<table>
<thead>
<tr>
<th>Chemical*</th>
<th>Type of response (ref. 1)</th>
<th>Type of chemical</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-Butanol</td>
<td>Attraction</td>
<td>Alcohol</td>
</tr>
<tr>
<td>2-Butanol</td>
<td>Attraction</td>
<td>Alcohol</td>
</tr>
<tr>
<td>Isoamyl alcohol</td>
<td>Attraction</td>
<td>Alcohol</td>
</tr>
<tr>
<td>Acetone</td>
<td>Attraction</td>
<td>Ketone</td>
</tr>
<tr>
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<td>Attraction</td>
<td>Ketone</td>
</tr>
<tr>
<td>Ethyl acetate</td>
<td>Attraction</td>
<td>Ester</td>
</tr>
<tr>
<td>Propyl acetate</td>
<td>Attraction</td>
<td>Ester</td>
</tr>
<tr>
<td>n-Butyl acetate</td>
<td>Attraction</td>
<td>Ester</td>
</tr>
<tr>
<td>Isoamyl acetate</td>
<td>Attraction</td>
<td>Ester</td>
</tr>
<tr>
<td>Aniline</td>
<td>Attraction</td>
<td>Aromatic compound</td>
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<td>Weak attraction</td>
<td>Alcohol</td>
</tr>
<tr>
<td>Benzylalcohol</td>
<td>Weak attraction</td>
<td>Alcohol</td>
</tr>
<tr>
<td>1-Propanol</td>
<td>Weak attraction</td>
<td>Alcohol</td>
</tr>
<tr>
<td>1-Octanol</td>
<td>Repulsion</td>
<td>Alcohol</td>
</tr>
</tbody>
</table>

*All of the chemicals were diluted with ethanol at a ratio of 1:1 except for 1-butanol and the three weak attractants, which were used undiluted.


Table S3. List of primers for quantitative RT-PCR

<table>
<thead>
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<th>Gene</th>
<th>Forward primer</th>
<th>Reverse primer</th>
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<tr>
<td>F47F6.1 (lin-42)</td>
<td>5′-CCACTGACCCCGAGAAGCAC-3′</td>
<td>5′-GAGTTGGTGCCACTTGTCGG-3′</td>
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<td>F01D5.5</td>
<td>5′-AACCTGTAACATGGCAGGAA-3′</td>
<td>5′-GCCGTCACCCAGTTAGAC-3′</td>
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<td>Y110A2AL.9</td>
<td>5′-ACCAAGGATGGTTTGACCCGG-3′</td>
<td>5′-TTGGTGACACTGATGCACCGTT-3′</td>
</tr>
<tr>
<td>T16D1.2 (pho-4)</td>
<td>5′-GAAGTTGATGATGGTTCAGG-3′</td>
<td>5′-GAGTTGGTGGCCTTGAC-3′</td>
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<td>M199.4 (clec-190)</td>
<td>5′-ATGATTGGAACCTGAAACG-3′</td>
<td>5′-CCAGAAAATCCGGTCCGT-3′</td>
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<td>F15A4.6</td>
<td>5′-CAATGCAATCGGTCTTCTTG-3′</td>
<td>5′-CCATCGTACACTTGTGCTCA-3′</td>
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<td>C30G12.2</td>
<td>5′-CTGCAGAAGGAGATGAGAAGA-3′</td>
<td>5′-CCATCGAATGGCGGTTCA-3′</td>
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<tr>
<td>F15E6.8 (dtc-7)</td>
<td>5′-TCTCCTGCGCCCTATTGCTG-3′</td>
<td>5′-CCATCGAATGGCGGTTCA-3′</td>
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<td>B0507.8</td>
<td>5′-AAAGAGAAGCAGGATGACG-3′</td>
<td>5′-CCATCGAATGGCGGTTCA-3′</td>
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<tr>
<td>ZC308.1 (gld-2)</td>
<td>5′-TCACTTCTTCTCTCTTGTCG-3′</td>
<td>5′-CCATCGAATGGCGGTTCA-3′</td>
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<td>F09E5.15 (prdx-2)</td>
<td>5′-GGGAAGCGAGAAGATGCGGTTCC-3′</td>
<td>5′-CCATCGAATGGCGGTTCA-3′</td>
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<td>W02B3.2 (grk-2)</td>
<td>5′-AGGATGTAACAGGAGATGCGGGA-3′</td>
<td>5′-CCATCGAATGGCGGTTCA-3′</td>
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<tr>
<td>M03F4.2 (act-4)</td>
<td>5′-GGCATCACACCTTCTACACAGA-3′</td>
<td>5′-TGGATTGAGTGAGACTCAGT-3′</td>
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