Catecholamine receptor polymorphisms affect decision-making in *C. elegans*

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**Abstract**

Innate behaviours are flexible: they change rapidly in response to transient environmental conditions, and are modified slowly by changes in the genome. A classical flexible behaviour is the exploration-exploitation decision, which describes the time at which foraging animals choose to abandon a depleting food supply. Here we use quantitative genetic analysis to examine the decision to leave a food patch in *Caenorhabditis elegans*. We find that patch-leaving is a multigenic trait regulated in part by naturally-occurring noncoding polymorphisms in *tyra-3*, which encodes a G protein-coupled catecholamine receptor related to vertebrate adrenergic receptors. *tyra-3* acts in sensory neurons that detect food-related cues, suggesting that the internal catecholamines detected by *tyra-3* regulate responses to external conditions. These results indicate that genetic variation and environmental cues can converge on common circuits to regulate behaviour, and suggest that catecholamines have an ancient role in regulating behavioural decisions.

Despite abundant evidence for heritability of behavioural traits within and between species, only a few naturally varying traits have been associated with polymorphisms in specific genes. Foraging for food is an ecologically relevant, environmentally regulated behaviour that is suitable for genetic analysis, as it can differ between populations of a species that live in different habitats. An essential foraging decision is the choice between exploiting...
existing resources and exploring other options that may provide new resources. This decision can be described by Charnov’s marginal value theorem, which proposes that the optimal time for an animal to leave a foraging ground occurs when local resource levels fall below the average level in the entire habitat. The marginal value theorem was developed for animals foraging for food in patchy environments, but has analogies with diverse decision-making processes in field biology, cognitive neuroscience, and economics.

Studies of patch-leaving behaviour in the nematode *C. elegans* have revealed innate, environmental, and experience-dependent factors that affect its foraging decisions. *C. elegans* rarely leaves a dense lawn of high-quality bacterial food, but more frequently leaves lawns of pathogenic bacteria or lawns that are spiked with chemical repellents. Males will leave lawns that do not contain potential mates, while hermaphrodites leave lawns when animal density is high. In addition, wild-type strains vary in their propensity to leave bacterial lawns based on a genetic polymorphism that affects the G protein-coupled neuropeptide receptor NPR-1. This polymorphism affects many foraging behaviours; low-activity NPR-1 strains aggregate into social feeding groups, move quickly on food, and have altered responses to oxygen, carbon dioxide, and pheromones compared to the N2 laboratory strain. The high-activity allele of NPR-1 in N2 arose in the laboratory, probably as an adaptation to laboratory conditions, so it is not known whether genetic variation affects *C. elegans* foraging in natural environments.

Natural genetic variation within a species can generate diversity in foraging behaviour, as exemplified by the polymorphic *Drosophila melanogaster foraging (for)* gene, which encodes a cGMP-dependent protein kinase. A low-activity allele of for is present in *Drosophila* sitter larvae, which move slowly on a food patch; a high-activity allele of for is present in rover larvae, which move quickly and disperse rapidly. A for-related cGMP-dependent kinase affects foraging in honeybees, ants, and nematodes, suggesting that diverse animals share molecular mechanisms for behavioural regulation.

To gain further insight into the genetics and neurobiology of lawn-leaving behaviour in *C. elegans*, we here use quantitative genetic analysis to examine its genetic architecture in wild-type strains, and show that genetic variation in multiple loci, including a catecholamine receptor, interacts with environmental conditions to regulate the exploitation-exploration decision.

### Multiple loci affect leaving behaviour

Different wild-type strains of *C. elegans* vary in their tendency to leave or remain on a standardized small lawn of bacterial food (Fig. 1a). For example, adult hermaphrodites from the laboratory strain N2 leave the lawn only once every 100 minutes, whereas animals from the CB4856 (HW) strain isolated from pineapple fields in Hawaii leave the lawn once every 5–6 minutes (Fig. 1b, Supplementary Movies 1 and 2). To determine the genetic architecture of this behavioural difference between N2 and HW, we quantified leaving rates in 91 N2-HW recombinant inbred advanced intercross lines (RIAILs). 58 of the RIAILs had low leaving rates comparable to N2, only 6–10 had high leaving rates comparable to HW, and 23
had intermediate rates (Fig. 1c). The excess of low leaving rates and the continuous
behavioural distribution in RIAILs suggest that leaving is a multigenic quantitative trait.

Quantitative trait locus (QTL) analysis of the RIAILs uncovered two regions with
significant effects on leaving rates, one on the X chromosome and one on chromosome II
(Fig. 1d). The X chromosome QTL overlapped with the location of the polymorphic G
protein-coupled neuropeptide receptor NPR-1, which affects many food-related
behaviours. The npr-1 polymorphism has previously been shown to affect leaving, as
well as locomotion speed on food, a behaviour that partially correlates with leaving rate
(Supplementary Fig. 2). The asymmetric distribution is consistent with a role for npr-1 in leaving behaviour, but indicates that npr-1 has epistatic
interactions with other loci segregating in the RIAILs.

The involvement of npr-1 in leaving behaviour was confirmed by analyzing near-isogenic
lines (NILs) containing the N2 and HW npr-1 alleles in the reciprocal strain background,
and by examining npr-1 null mutants (Supplementary Fig. 3). Specific transgenic expression
of the N2 npr-1 allele in its essential site of action, the RMG motor neurons, sharply
reduced the leaving rate of HW animals (Supplementary Fig. 3). Thus npr-1 is a regulator of
HW leaving rates, but not the only contributing gene.

### tyra-3 affects leaving behaviour

Studies in yeast, flies, mice, and plants have shown that individual QTLs often resolve into
several genes that contribute to phenotypic variance. Similarly, fine-mapping of the ~
1 Mb QTL that contained npr-1 suggested the existence of multiple loci that affected leaving
rates. A NIL with <150 kb of N2 DNA spanning the npr-1 locus introgressed into HW had
N2-like leaving rates (leav-1 QTL, Fig 2a and Supplementary Fig. 3). A second NIL with
700 kb of N2 DNA that did not cover npr-1 introgressed into HW also had a low leaving
rate, with about half the leaving rate of HW (leav-2 QTL, Fig. 2a). These results suggest the
existence of a second X-linked locus that affects leaving rates, which we called leav-2. The
leav-2 region did not affect leaving in the N2 genetic background (Fig. 2a), so all subsequent
experiments were conducted in the HW background.

A 100 kb minimal region for leav-2 was identified by analyzing the breakpoints of
individual RIAILs (Supplementary Fig. 4 and Supplementary Methods). We characterized
the genetic properties of leav-2 by crossing the leav-2 NIL strain with HW. The
heterozygous F1 progeny had leaving rates similar to the leav-2 NIL (Fig. 2a), indicating
that the N2 leav-2 locus was dominant to HW and suggesting that N2 transgenes covering
the relevant gene should reduce the leaving rate of HW animals. Therefore, overlapping N2
genomic DNA fragments from the 100 kb minimal leav-2 region were introduced into HW
animals by microinjection (Fig. 2b and Supplementary Fig. 5). A single gene in this region
reduced leaving rates: tyra-3, which encodes a G protein-coupled receptor for the
invertebrate norephinephrine-like neurotransmitters tyramine and octopamine. Tyramine
and octopamine receptors are related to vertebrate adrenergic receptors, and are thought to carry out analogous functions. tyra-3 genomic fragments from the N2 strain were more active than tyra-3 fragments from the HW strain injected at the same concentration, consistent with the possibility that tyra-3 is a polymorphic gene that differs between N2 and HW (Fig. 2b).

If leav-2 corresponds to tyra-3, a tyra-3 mutation should eliminate its activity. To test this prediction genetically, a null allele of tyra-3 in an N2 background was introgressed into a HW background. The N2 region in the resulting NIL covered from 4.9 to 5.4 MB of the X chromosome, the inferred position of leav-2. The tyra-3(ok325) null NIL had high (HW-like) leaving rates, suggesting that N2 leav-2 activity was not present in the strain (Fig. 2a). Heterozygotes between HW and the near-isogenic tyra-3(ok325) null strain also had high leaving rates (Fig. 2a). These results are as expected if the active locus in leav-2 is tyra-3; however, other genes within the introgressed regions could also contribute to the different leaving rates.

To strengthen the connection between tyra-3 and leav-2, RNAi against tyra-3 was performed in the leav-2 NIL that has low leaving rates due to the presence of the N2 QTL. Knockdown of tyra-3 increased the leaving rate of the leav-2 NIL to levels observed in HW animals, the result predicted if the tyra-3 locus from N2 reduces leaving (Fig. 2c). Comparable experiments in a pure HW strain had minimal effects, as expected if tyra-3 activity in HW is already low.

Further confirmation that the HW allele of tyra-3 has reduced biological activity was provided by examining the one phenotype previously associated with tyra-3, avoidance of dilute octanol. tyra-3 null mutants avoid octanol more strongly than wild-type N2; the NIL strain with the HW tyra-3 allele had a similar enhanced octanol response, suggesting that the HW tyra-3 allele has reduced tyra-3 function (Supplementary Fig. 6).

**Noncoding changes affect tyra-3 activity**

The differential activity of N2 and HW genomic tyra-3 fragments in the leaving assay suggested that N2 and HW alleles are functionally distinct (Fig. 2b). To identify polymorphisms between N2 and HW alleles of tyra-3, we sequenced ~19 kb surrounding the tyra-3 locus in HW. There were 34 differences between HW and the N2 consensus genomic sequence (Fig. 3a): 33 noncoding changes and a single coding difference that changed a glutamate in the tyra-3b isoform to glycine.

Sequences that contribute to the differential activity of N2 and HW tyra-3 alleles were localized further using transgenic assays. We fused N2 and HW tyra-3b cDNAs to 4.9 kb of noncoding N2 or HW sequence upstream of the tyra-3b start site and introduced each of the four resulting clones into the HW strain. tyra-3 transgenes with the N2 noncoding sequence were significantly more potent than comparable transgenes with the HW sequence, regardless of whether they preceded N2 or HW tyra-3 cDNAs (Fig. 3b), excluding the coding polymorphism and localizing a functional difference between N2 and HW tyra-3 genes to a 4.9 kb region that harbours 5 noncoding SNPs, 1 single nucleotide insertion, and a
184 bp deletion in HW. To narrow the relevant change down further, the 184 bp deletion was engineered into the N2 tyra-3 genomic fragment; this clone was significantly less potent in the leaving assay than the full N2 genomic fragment (Supplementary Fig. 7). These results indicate that the 184 bp deletion represents at least part of the functional difference between N2 and HW tyra-3 alleles.

Sequence variation in tyra-3 noncoding regions could affect the level or location of tyra-3 expression. Quantitative RT-PCR of tyra-3 mRNA levels in mixed-stage animals indicated that N2 expressed approximately twice as much tyra-3 mRNA as HW, consistent with increased tyra-3 activity in the N2 strain (Fig. 3c). The leav-2 NIL with N2 tyra-3 introgressed into HW also had high tyra-3 mRNA levels, suggesting that cis-acting changes affect tyra-3 expression (Fig. 3c).

Since both N2 and HW were cultivated in the laboratory for many years before permanent cultures were frozen, we wished to exclude the possibility that the tyra-3 polymorphisms were laboratory-derived. Therefore, 19 kb of the tyra-3 locus was sequenced in all wild strains tested for leaving behaviour in Fig. 1, including three strains that were frozen immediately after their isolation. Each strain represents a different C. elegans haplotype group. Both N2-like and HW-like tyra-3 sequences were represented in the wild-caught strains, confirming the wild ancestry of both alleles (Supplementary Table 1 and Supplementary Methods). Notably, the tyra-3 locus of MY1 was identical to N2 and, correspondingly, the leaving rate of MY1 was similar to that of N2.

**tyra-3 acts in sensory neurons**

The identification of tyra-3 provided an opportunity to characterize the neuronal basis of the decision to leave or remain on a food patch. The biological activity of a transgene with 4.9 kb upstream of the tyra-3b start site fused to a tyra-3 cDNA (Fig. 3b) implied that it was expressed in cells that regulate leaving behaviour. When this 4.9 kb region was fused to GFP, it drove reliable expression in ASK, ADL, AIM, AUA, BAG, CEP, OLQ, and SDQL neurons, in other unidentified neurons in the ventral ganglion and the tail, occasionally in AFD and AWC neurons, and in two non-neuronal cell types, the spermatheca and the distal tip cell (Fig. 4a and data not shown). The same set of cells was observed with reporter genes bearing either N2 or HW tyra-3 upstream regions, and in both N2 and HW genetic backgrounds. Together with the quantitative RT-PCR data (Fig. 3c), these results suggest that different tyra-3 expression levels, not different sites of expression, distinguish N2 and HW alleles.

The neurons whose activity is regulated by tyra-3 were localized further by expressing tyra-3 cDNAs from cell type-specific promoters. tyra-3 expression in ASK or BAG sensory neurons significantly reduced leaving, but expression in the CEP or ADL sensory neurons did not (Fig. 4b). The ASK neurons sense attractive food-derived amino acids and regulate search behaviours after animals are removed from food. The BAG neurons sense CO2 and O2, two cues associated with bacterial metabolism. Lowering O2 to levels that activate BAG reduced leaving rates (Supplementary Fig. 8).
To ask whether the *tyra-3* noncoding polymorphism affects expression in relevant neurons, single-copy N2 or HW *tyra-3b* promoters driving GFP were inserted into a single, defined chromosomal location using the MosSCI technique. GFP levels in ASK neurons were significantly higher for transgenes containing the N2 promoter compared to those containing the HW promoter. These results suggest that the N2 *tyra-3* locus is associated with higher *tyra-3* expression in ASK, as well as higher *tyra-3* mRNA expression at a whole-animal level; expression in BAG was not examined.

The behavioural functions of ASK and BAG, and *tyra-3*’s effect on those functions, were assessed by killing the neurons in different genetic backgrounds. Killing the ASK neurons reduced the leaving rate of HW animals, indicating that ASK can promote leaving. The ablation resembled the effect of the ASK::*tyra-3* transgene, suggesting that *tyra-3* reduces ASK activity. In agreement with this idea, killing the ASK neurons in a strain with the N2 high-activity *tyra-3* allele did not reduce their leaving rates further. The effect of *tyra-3* on ASK was selective for this assay; *tyra-3* did not reduce lysine chemotaxis, a second ASK-dependent behaviour.

Killing the BAG neurons increased leaving rates in the strain with the N2 *tyra-3* allele, demonstrating that BAG neurons prevent leaving. However, killing BAG had no effect in the strain with the HW *tyra-3* allele, suggesting that BAG activity is already low in this strain under the assay conditions. The ablation and genetic results suggest that the N2 *tyra-3* allele decreases ASK activity and increases BAG activity, two changes that act together to prevent leaving.

**Gene-gene-environment interactions**

Like most natural behaviours, the decision to leave a food patch is regulated by multiple genes and the environment; it responds to genetic variation in *tyra-3*, *npr-1*, and additional genes on the autosomes as well as food quality and quantity. Our results suggested that the N2 *npr-1* allele was epistatic to *tyra-3*; animals with the N2 *npr-1* allele had low leaving rates regardless of the *tyra-3* genotype. However, N2 *npr-1* reduced the leaving rate to almost zero, making it difficult to detect any further reduction. To make the assay more powerful, leaving was assayed on bacterial lawns of different densities. Leaving rates of all genotypes increased on thinner lawns and decreased on thicker lawns, but the thickness of the lawn changed the genetic interaction between *tyra-3* and *npr-1*. In the standard leaving assay, *tyra-3* polymorphisms had different effects only in the presence of the HW *npr-1* allele; on a thinner lawn, only in the presence of the N2 *npr-1* allele. Thus the epistatic relationship between *npr-1* and *tyra-3* is defined by the specific environment, not by an intrinsic regulatory relationship between the genes.

**Discussion**

Our results show that natural variation in *tyra-3* affects patch leaving, a behaviour representative of the exploration-exploitation decision. *tyra-3* encodes a G protein-coupled receptor activated by the invertebrate transmitters tyramine and octopamine, which are
structurally related to vertebrate epinephrine and norepinephrine. Catecholamines are known to regulate arousal systems that affect many behaviours and behavioural decisions. In C. elegans, octopamine drives sensory, molecular, and behavioural responses to starvation, and tyramine affects specific aspects of locomotion. In insects, octopamine affects locomotory activity, arousal, and aggression. Mammalian norepinephrine is generally implicated in arousal behaviours, and norepinephrine release from the locus coeruleus is associated with switching between different tasks, a cognitive function with analogies to the exploration-exploitation decision.

Relatively few natural behavioural variations have been mapped to the single-gene level in any animal, and it is interesting that several of these variations affect G protein-coupled receptor signalling systems. We speculate that these receptor pathways may serve as common substrates of behavioural variation. All animal genomes encode many G protein-coupled receptors with different expression patterns. These receptors may provide a reservoir for genetic changes, as alteration in an individual receptor could cause relatively discrete effects without disrupting the entire system.

QTL mapping in rodents and in Drosophila indicates that most behavioural traits are polygenic, with widespread epistatic effects. In agreement with this conclusion, our analysis suggests the existence of epistatic interactions between tyra-3, npr-1, and at least one additional locus. Importantly, the non-additive interactions between tyra-3 and npr-1 are not stable, but vary based on the genetic background and the environment, similar to what has been found with yeast sporulation QTLs.

QTL analysis was performed on the mean leaving rates of N2-HW recombinant inbred advanced intercross lines (RIALs) by nonparametric interval mapping in R/qtl.
Significance levels were estimated from 10,000 permutations of the data. Near-Isogenic Lines were created by backcrossing a chromosomal region or allele into the desired genetic background at least 9 times.

Extrachromosomal transgenes were made by injection of DNA clones into the gonads of young adult hermaphrodites together with a fluorescent coinjection marker. To control for variation between transgenes, between two and five independent lines from each injection were characterized.

Single-copy insertion of transgenes was performed using the direct MosSCI transposition technique, targeting the *ttTi5605* Mos allele on chromosome II.

**Methods**

**Analysis of Behaviour in the Leaving Assay**

6 cm NGM agar plates were seeded with 70 μL (conditioning plate) or with 10 μL (assay plate) of a fresh overnight culture of *E. coli* HB101 diluted in LB to OD$_{600nm}$=2.0. 90 min after seeding the plates, ten young adult hermaphrodites were picked onto the conditioning plate. 30 min after being placed on the conditioning plates, seven of the animals were transferred onto the lawn of the assay plate. The 30 min leaving assay began 1 hr after placing the seven animals on the assay plate. The number of leaving events was recorded manually by examining the video recordings, and further behavioural analysis was conducted with a Matlab code adapted from the Parallel Worm Tracker. A leaving event was defined as an episode in which the whole body of an animal left the bacterial lawn and the animal did not reverse immediately to return to the lawn. The leaving rate was calculated as the number of leaving events per worm minute spent inside the bacterial lawn. Experiments on each strain were repeated at least three times.

**Quantitative Trait Locus Analysis**

The N2-HW recombinant inbred advanced intercross lines (RIALs) used in this study represent the terminal generation of a 20-generation pedigree founded by reciprocal crosses between N2 and HW. The lines were constructed through 10 generations of intercrossing followed by 10 generations of selfing. They have been genotyped at 1454 nuclear and one mitochondrial markers and have a 5.3-fold expansion of the F2 genetic map. QTL analysis was performed on the mean leaving rates of N2-HW recombinant inbred advanced intercross lines (RIALs) by nonparametric interval mapping in R/qtl. Significance levels were estimated from 10,000 permutations of the data.

**Quantitative RT-PCR**

Total RNA from mixed stage worms was isolated with Trizol. 1.5 μg of RNA and oligo-dT were used for reverse transcription using SuperScript III First-Strand Synthesis (Invitrogen) according to the manufacturer’s instructions. Real-time PCR was performed with Fast SYBR Green Master Mix (Applied Biosystems) on a 7900HT Real-Time PCR System (Applied Biosystems). *act-3* was used as the calibrator for relative quantitation. 5' primers
corresponded to upstream exons that distinguished tyra-3 isoforms, and 3′ primers corresponded to shared exon sequence. Primers used were:

- tyra-3a&c.2_F, ccacctggcaatagcagcag
- tyra-3b_F, ggcctattggtgcggtttg
- tyra-3a & tyra-3b_R, tccttctggctgcaaatac
- act-3_F, ttcagatcgagaccattcaaa
- act-3_R, gcaaatgtaggggtcctctatg

**tyra-3 Expression Pattern**

The N2 and HW 4.9 kb tyra-3b promoters were amplified using primers: tcaacctggaacctacaaggg and cGataagcaagatgctaggt, which overlaps the coding region by 4 bp. The ATG start codon is mutated to ATC (mutation is uppercase in primer). These promoters were individually fused by PCR to a fragment containing GFP followed by the unc-54 3′-UTR, as described. These PCR products were injected individually into both HW and N2 animals at 20 ng/μL. Cells expressing GFP were identified by Nomarski microscopy in both L1 and adult hermaphrodites. The identification of some cells was aided by injecting Ptyra-3b::GFP-expressing animals with promoter-mCherry fusions with established expression patterns. In this manner, the AIM neurons were identified as Ptyra-3b::GFP-expressing cells based on their position and the absence of co-localization with Ptx-3::mCherry. The BAG neurons co-expressed Ptyra-3b::GFP and Pflp-17::mCherry. The CEP neurons co-expressed Ptyra-3b::GFP and Pdsrta-9::mCherry. The ADL neurons co-expressed Ptyra-3b::GFP and Psri-51::mCherry.

**Extrachromosomal transgenes**

Transgenes were made by injection of DNA clones into the gonads of young adult hermaphrodites together with a fluorescent coinjection marker. To control for variation between transgenes, between two and five independent lines from each injection were characterized.

**Generation of MosSCI Lines and Quantitation of GFP Fluorescence in ASK**

Single-copy insertion of transgenes was performed using the direct MosSCI technique targeting the tfTi5605 Mos allele on chromosome II, as described. A schematic of the mechanism underlying MosSCI is shown in Fig. 4c.

The pCFJ151 targeting vector was modified by the introduction of an FseI restriction site into the multiple cloning site by site-directed mutagenesis using the primers gtaatacgacttaaagctgagccgccgccctgctagaggtcagctaggacctgccgtaggagttac and ggtgagctctgtgcctagctacccggcgggctgacggtctacgtattac to make pAB1. An FseI-SpeI fragment from a pSM vector containing N2-Ptyra-3b::N2-tyra-3b::SL2 GFP::unc-54 3′-UTR or HW-Ptyra-3b::N2-tyra-3b::SL2 GFP::unc-54 3′-UTR was cloned into pAB1.
For each tyra-3-containing test plasmid, about fifty EG4322 animals were injected with a mixture of tyra-3 plasmid, pGH8, pCFJ90, pCFJ104, and pJL43.1. After positive and negative selection and full sequencing of the insert, two inserted transgenes each of N2-Ptyra-3b and HW-Ptyra-3b were backcrossed to HW males seven times, selecting GFP-fluorescent hermaphrodites each generation. The transgene-containing chromosome was then homozygosed.

The strains containing the single-copy transgene in a HW background were injected with Psra-9::mCherry to identify ASK. Young adult hermaphrodites were examined on a Zeiss Imager Z.1 with a 60X objective focused on ASK using mCherry to prevent bleaching of GFP signal. Fluorescence signals were acquired with fixed acquisition times (30–50 msec for mCherry, 100 msec for GFP). Background mean fluorescence intensity adjacent to ASK was subtracted from the ASK signal.

RNAi

RNA interference was performed essentially as described. A fragment common to all tyra-3 isoforms was amplified. The following primers were used, which include the T7 sequence (underlined):

\[
\text{taatacgactcactatagggagagaaaatggcagcaggacttt}
\]
\[
\text{taatacgactcactatagggagaatcctcgcagtctgtggagt}
\]

\textit{in vitro} transcription was performed with RiboMAX kit (Promega). dsRNA was injected at 1.2 μg/μL into the gonads of adult hermaphrodites. Eggs laid 24 and 48 hours after injection were used for the behavioural assays.

Octanol Avoidance Assay

Avoidance assays were conducted essentially as described. In brief, ~20 three-day old animals were picked off of their growth plates food into a transfer plate without bacteria where they were allowed to crawl and rid themselves of bacteria. Animals were then transferred onto an NGM plate without food. After 40 minutes, a microcapillary with 30% octanol (v/v diluted fresh every day in ethanol) was presented in front of the animal’s nose. The time to reverse was recorded. If animals did not reverse within 20 seconds, the assay was stopped. Animals were presented with odor 1–3 times per experiment, with at least 3 minutes of rest interval. We replicated published results demonstrating that tyra-3 null mutants had more rapid responses than N2 in the presence of exogenous serotonin and tyramine but also observed more rapid responses in the absence of exogenous neuromodulators, as shown in Supplementary Fig. 6.

Cell Ablations

For leaving behaviour assays ASK was ablated with a laser microbeam as described. BAG was killed using split human caspase 3 fragments expressed from \textit{flp-17} and \textit{glb-5} promoters that overlapped only in BAG. For lysine chemotaxis assays, ASK was killed using a mouse caspase 1 gene expressed from the \textit{sra-9} promoter. The ASK strain was a generous gift from Ryuzo Shingai.
Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1. Lawn-leaving behaviour varies between wild-type *C. elegans* strains

**a)** Lawn-leaving assays. Top: Six adult HW hermaphrodites on a bacterial lawn. One animal has left the lawn and one is leaving. Bottom: Track of a HW animal during 5 min of an assay; colour shows passage of time. The border of the lawn is outlined. Scale bar, 6 mm.

**b)** Leaving rates of six wild-type strains.

**c)** Leaving rates of 91 N2-HW recombinant inbred advanced intercross lines (RIAILs) and parental strains.

**d)** QTL analysis of RIAILs shown in c. The horizontal line denotes the P<0.01 genome-wide significance threshold. Error bars indicate s.e.m.
Figure 2. N2 and HW *tyra-3* alleles differentially affect leaving rates

**a)** Dissection of the QTL on X into two loci: *leav-1* (4.70–4.78 Mb) and *leav-2* (4.78–5.75 Mb). ‘Genotype’ shows chromosomes; thick line is X chromosome. Blue denotes HW DNA, red denotes N2 DNA, and yellow denotes the *tyra-3(ok325)* null mutant. In heterozygous strains, both X chromosomes are diagrammed.

**b)** *tyra-3* genomic fragments (Fig. 3a) reduce HW leaving rates. Blue, HW transgenes; red, N2 transgenes. Two-way ANOVA showed significant effects of both transgene concentration and DNA strain of origin.

**c)** Effect of *tyra-3* RNAi. Error bars indicate s.e.m. * P<0.05, ** P<0.01, or *** P<0.001 by t-test or ANOVA with Dunnett test.
Figure 3. Noncoding changes in *tyra-3* affect its activity and expression level

a) HW polymorphisms in the *tyra-3* locus relative to N2. *tyra-3* encodes three predicted G protein-coupled receptors. The genomic region examined in Fig. 2b and the 4.9 kb promoter used in Figs. 3b and 4a are indicated.

b) Leaving rates of transgenic HW animals with *tyra-3b* promoters fused to *tyra-3b* cDNAs. Error bars indicate s.e.m. ** P<0.01 by two-way ANOVA; no statistical interaction between the promoter and the cDNA.

c) Relative amounts of *tyra-3* isoform mRNAs in HW, N2, and *leav-2* strains (Fig. 2a). Error bars indicate s.d. ** P<0.01 compared to HW, ANOVA with Dunnett test.
Figure 4. *tyra-3* acts in ASK and BAG sensory neurons

**a)** Expression of 4.9 kb N2 *tyra-3b* promoter::GFP fusion (Fig. 3a) in HW animal; HW *tyra-3b* promoter::GFP is expressed in the same cells. Posterior signal is gut autofluorescence. Scale bar = 20μm. **b)** Leaving rates of HW strains expressing *tyra-3b* in specific cells. **c)** Left: GFP fluorescence intensity in ASK of HW animals with a MosSCI insertion of N2 or HW 4.9 kb *tyra-3b* promoter::GFP. Right: Schematic of MosSCI technique. **d)** Leaving rates after killing ASK or BAG in HW and *leav-2* strains (Fig. 2a). Error bars indicate s.e.m. * P<0.05, ** P<0.01, or *** P<0.001 by t-test or ANOVA with Dunnett test.