CREB mediates the *C. elegans* dauer polyphenism through direct and cell-autonomous regulation of TGF-β expression

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Abstract

Animals can adapt to dynamic environmental conditions by modulating their developmental programs. Understanding the genetic architecture and molecular mechanisms underlying developmental plasticity in response to changing environments is an important and emerging area of research. Here, we show a novel role of cyclic AMP response element binding protein (CREB)-encoding *chr-1* gene in developmental polyphenism of *C. elegans*. Under conditions that promote normal development in wild-type animals, *chr-1* mutants inappropriately form transient pre-dauer (L2d) larva and express the L2d marker gene. L2d formation in *chr-1* mutants is specifically induced by the ascaroside pheromone ascr#5 (asc-ωC3; C3), and *chr-1* functions autonomously in the ascr#5-sensing ASI neurons to inhibit L2d formation. Moreover, we find that CRH-1 directly binds upstream of the *daf-7* TGF-β locus and promotes its expression in the ASI neurons. Taken together, these results provide new insight into how animals alter their developmental programs in response to environmental changes.

Author summary

* C. elegans* exhibits polyphenic development that is regulated by environmental conditions. Newly hatched larvae of *C. elegans* sense and integrate food availability and population density to determine whether to undergo normal reproductive development or arrest as non-aging, stress-resistant dauer. Dauer entry in *C. elegans* is proceeded by a commitment to either L2d, an obligate dauer precursor, or reproductive growth. Here, we find a *flp-8* neuropeptide gene as the bona fide L2d marker and show that the *chr-1* gene controls the L2d formation by direct regulation of *daf-7* TGF-β expression. Identification of this gene, as well as the novel L2d marker, now provides a unique opportunity to study the under-explored L2d formation of *C. elegans*. 
Introduction

Animals exhibit phenotypic traits whose expression can be influenced by environmental signals, including temperature, food availability, and crowding. These signals are sensed, processed, and integrated by the nervous system, and govern appropriate developmental and/or behavioral phenotypes through effecting local and systemic changes in gene expression and/or endocrine signals. For example, population density and food supply determine wing development (winged vs. wingless) of the pea aphid (*A. pisum*) via ecdysone signaling [1]. This extreme form of phenotypic plasticity is referred to as polyphenism—the development of alternate and distinct phenotypes in animals of the same genotype—and is found across taxa [2,3]. Although the ability of animals to modulate their developmental programs in response to changing environmental conditions is crucial for their survival, the molecular mechanisms underlying such developmental plasticity are not yet fully understood.

The nematode *Caenorhabditis elegans* exhibits polyphenic development that is regulated by environmental conditions. Newly hatched L1/L2 larvae of *C. elegans* sense and integrate environmental signals, including food availability, temperature, and the abundance of pheromones (a population density indicator) to determine whether to undergo normal reproductive development into the L3 larval stage, or arrest at an alternate L3 stage, known as dauer. Dauer larvae are developmentally arrested, non-aging, stress-resistant, and can re-enter normal reproductive development if conditions improve [4,5] (Fig. 1A). Under less favorable conditions, L1 larvae may transiently develop as L2d larvae, an obligate precursor to dauer commitment, but which still retain the capacity to resume reproductive development [4] (Fig. 1A). Thus, worms assess and weigh multiple factors at multiple time points along the dauer trajectory in making the dauer decision. However, it is not clear how worms evaluate external and internal signals at specific developmental time windows and execute dauer formation. Specifically, the signals and molecules that promote the L1-to-L2d developmental transition are also not well understood.

*C. elegans* secretes pheromones, which are a complex cocktail of small molecules called ascarosides [6–9]. Moreover, worms produce and secrete distinct concentrations and combinations of ascarosides depending upon developmental stages and environmental conditions [7,10,11]. For example, ascarosides ascr#2 (asc-C6MK; C6), ascr#3 (asc-ΔC9; C9), and ascr#5 (asc-ωC3; C3), act as a dauer-inducing cue and worms produce specific concentrations of each ascaroside depending on cultivation conditions [6,7]. Although the potency and property of pheromone components in inducing dauer formation have been described, little is known about the contribution of each component to specific time windows along the dauer trajectory.

The molecular and genetic mechanisms underlying the decision to enter the dauer stage have been extensively studied, and have established two neuroendocrine *daf-7*/TGF-β and *daf-2*/insulin signaling pathways as critical mediators of this binary fate decision [12–14]. Expression of *daf-7* in the ASI sensory neuron type is regulated by environmental signals, including temperature, food availability, pheromone, and infection [12,13,15,16]. However, the mechanisms that regulate *daf-7* expression as a function of environmental conditions have not been fully described.

*crh-1*-encoded CREB (cAMP response element binding protein) plays critical roles in *C. elegans* metabolism, sensory responses, development, lifespan, and learning and memory [17–29] Similar to mammalian CREB, CRH-1 directly regulates target gene expression via the evolutionarily conserved cAMP-responsive element (CRE) and represents a nutritive or metabolic indicator [30]. However, a role for CREB in developmental plasticity, such as the decision to enter the dauer stage in *C. elegans*, has not been explored.
Here, we show that CRH-1 plays a role in the decision to enter the L2d stage. In rich food conditions, pheromone is generally insufficient to induce L2d/dauer formation in wild-type animals. However, \textit{crh-1} mutants exhibit transient L2d formation and express the L2d marker.

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**Fig 1.** The \textit{flp-8} expression is turned on in a subset of touch receptor neurons of L2d larvae. (A) \textit{C. elegans} life cycle at 25°C. In favorable conditions, newly hatched L1/L2 larvae undergo normal reproductive development into L3 larvae. Under unfavorable conditions, L1 larvae arrest as dauer larvae via the L2d pre-dauer stage. Under less favorable conditions, L1 larvae transiently become L2d larvae, which can resume reproductive growth. Numbers show the approximate duration in hours of development of each stage or hours after hatching (hAH). (B) Representative images of wild-type animals expressing of \textit{flp-8::gfp} in a subset of touch receptor neurons in the presence of heat-killed OP50 food without (left column) or with (right three columns) ascr#5 pheromone. The color-matched boxed regions of the second column are shown at higher magnification. hAH, hour after hatching. Scale bars: 50 μm. (C) Percent of \textit{flp-8::gfp} expression in the AVM, ALM, and PLM neurons by wild-type animals when grown in the presence of heat-killed OP50 food and the presence or absence of ascr#5 pheromone. n ≥ 30 for each. Error bars indicate SEM. * and ** indicate different from wild-type at p < 0.05 and p < 0.001, respectively (student t-test).

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gene \textit{flp-8} under such conditions. L2d formation of \textit{crh-1} mutants is induced by ascr\#5, but not by ascr\#2 or ascr\#3, and expression of CRH-1 in the ascr\#5-sensing ASI neurons is sufficient to rescue inappropriate L2d formation in \textit{crh-1} mutants. We find that CRH-1 acts cell-autonomously in the ASI sensory neurons to regulate \textit{daf-7} TGF-\(\beta\) expression and that ASI-specific overexpression of \textit{daf-7} is sufficient to block L2d formation in \textit{crh-1} mutants. We also identify a functional CRE in \textit{daf-7} regulatory sequences and show that it is specifically bound by CRH-1, confirming that CRH-1 regulates \textit{daf-7} expression in ASI. Together, these results reveal that CREB modulates developmental plasticity by repressing L2d entry in the presence of rich food sources and abundant ascr\#5, via direct regulation of \textit{daf-7} TGF-\(\beta\) expression in the ascr\#5-sensing ASI neurons.

**Results**

A \textit{flp-8} FMRFamide-like neuropeptide gene is expressed in a subset of touch receptor neurons in pre-dauer larvae

To better understand the mechanisms underlying L2d larvae formation, we first screened for a marker that specifically identifies the L2d developmental stage. The \textit{flp-8} FMRFamide-like gene was previously shown to be differentially expressed in a subset of touch receptor neurons (AVM, ALM, PLM; referred to as AAP-TRN henceforth) in dauer larvae in addition to expression in the AUA, URX and PVM neurons throughout larval development [31]. We monitored the onset and maintenance of \textit{flp-8p::gfp} reporter gene expression in the AAP-TRN of animals grown under non-dauer-inducing conditions and dauer-inducing conditions with limited heat-killed OP50 food and with and without the ascr\#5 pheromone, respectively [32]. Consistent with the previous study [31], we detected the expression of \textit{flp-8p::gfp} in AVM only in the dauer-inducing conditions (Fig 1B and 1C). Compared to under non-dauer-inducing conditions, expression in AVM was first detected 40 h after hatching (hAH) with more prominent expression evident at 60 hAH; 60 hAH is approximately 20 h prior to dauer entry/commitment under these conditions (Fig 1B and 1C). Animals expressing \textit{flp-8p::gfp} in the AAP-TRN of larvae at 60 hAH were morphologically similar to dauer larvae but did not survive under 1% sodium dodecyl sulfate (SDS) [5] treatment (S1 Fig), implying that \textit{flp-8} expression in the AAP-TRN begins at the L2d stage, before the completion of dauer cuticular remodeling that confers resistance to this detergent. We noted the onset of \textit{flp-8} expression in the ALM and PLM neurons at 55 hAH, five hour prior to dauer entry (Fig 1B and 1C). All dauer larvae expressed \textit{flp-8p::gfp} in the AAP-TRN (Fig 1B and 1C). Moreover, \textit{flp-8p::gfp} was expressed in the AAP-TRN of \textit{daf-7} mutant larvae (S2 Fig). Taken together, these results suggest that the onset of \textit{flp-8p::gfp} expression in the AAP-TRN and possibly the ALM and PLM neurons can be used as a marker for the L2d stage.

\textit{crh-1} CREB mutants express L2d marker genes in non-dauer-inducing conditions

In a specific non-dauer-inducing condition with abundant live OP50 food and high concentrations of ascr\#5, we found that about 25% \textit{crh-1} mutant larvae inappropriately expressed the \textit{flp-8} reporter gene in the AAP-TRN, which is significantly different from wild-type larvae (Fig 2A and 2B). However, in dauer-inducing conditions with limited heat-killed OP50 food and high concentrations of ascr\#5 pheromone, the expression of \textit{flp-8} in \textit{crh-1} mutant larvae in AAP-TRN was similar to that of wild-type, although a slightly smaller percentage of \textit{crh-1} mutants expressed GFP in the AAP-TRN as compared to wild-type (S3 Fig). All assays were conducted in this non-dauer inducing condition with abundant live OP50 food unless noted.
other, the onset of expression in these neurons in *crh-1* mutants under non-dauer-inducing conditions was approximately 35 hAH, five hours earlier than in the dauer-inducing condition, likely due to their rapid growth from having live food available [33]. *flp-8* expression was not observed in the AAP-TRN of wild-type or *crh-1* mutant larvae in the absence of pheromone (S4 Fig), indicating that the exposure to pheromone is necessary for induction of *flp-8* expression in the AAP-TRN. Additional genetic marker, *nhr-246* transcription factor, for the L2d stage was also expressed in the intestine of *crh-1* mutant larvae [34] (S5 Fig). These results show that *crh-1* mutants inappropriately express L2d marker genes in conditions that do not promote dauer formation in wild-type animals.

**crh-1** mutants inappropriately form transient pre-dauer L2d larvae upon exposure to ascr#5

To better understand the *crh-1* mutant phenotype, we next examined the ability of *crh-1* mutants to enter the dauer stage in response to different concentrations of ascaroside pheromones, under standardized dauer-inducing conditions. We found that *crh-1* mutants exhibited normal or weakly decreased pheromone-mediated dauer formation (S3 Fig). However, in a non-dauer inducing condition with abundant live OP50 food and high concentrations of crude pheromone, *crh-1* mutants formed thin dauer-like larvae 40–60 hAH (Figs 2C and S4) consistent with inappropriate induction of the L2d stage. To assess whether these larvae were indeed dauers, we treated them with 1% SDS. None of the *crh-1* dauer-like larvae survived under 1% SDS treatment (S1 Fig), indicating that these larvae have not fully remodeled their cuticle as expected for dauer larvae. We next characterized their morphology using scanning electron microscopy. The mouths of *crh-1* dauer-like larvae remained open as in L3 larvae (S6 Fig), suggesting, in combination with their other phenotypic attributes, that they are indeed L2d larvae. Moreover, by 65 hAH, these larvae had resumed reproductive development, consistent with transient passage through the L2d state and incongruent with dauer commitment and exit.

To determine whether the transient L2d formation of *crh-1* mutants is induced by specific pheromone components, we tested several ascaroside pheromones, including ascr#2, ascr#3, and ascr#5. All assays were conducted beginning 40 hAH unless noted otherwise. We found that *crh-1* mutants specifically formed transient L2d in the presence of micromolar concentrations of ascr#5, but not ascr#2 or ascr#3 (Figs 2D and S7). Moreover, *flp-8* was not expressed in the AAP-TRN of *crh-1* mutant larvae in response to ascr#2 and ascr#3 (S8 Fig).

ascr#5-mediated dauer formation has been shown to be primarily mediated by a pair of GTP-binding protein (G protein)-coupled receptors (GPCRs), SRG-36 and SRG-37 expressed specifically in ASI, and partly by another pair of GPCRs, SRBC-64 and SRBC-66 expressed in the ASK sensory neurons [15,35]. We found that the transient L2d formation of *crh-1* mutants was suppressed completely by mutations in both *srg-36* and *srg-37* and partially by mutations in both *srbc-64* and *srbc-66* (Fig 2E), supporting the specificity of ascr#5 in the transient L2d formation of *crh-1* mutants. Taken together, these results imply that *crh-1* mutants...
inappropriately and transiently enter into the L2d stage in response to ascr#5 prior to resuming reproductive development.

crh-1 acts in the ASI neurons to regulate ascr#5-mediated L2d formation

To identify the molecular mechanisms by which CRH-1 regulates ascr#5-mediated L2d formation, we identified the site of action of CRH-1. Previously, crh-1 had been shown to be expressed in a set of head neurons of adult C. elegans [17,24]. To further verify the expression patterns of crh-1, we generated transgenic animals expressing GFP under the control of sequences 2601 bp, 2411 bp, 1094 bp, or 737 bp (crh-1p1::gfp to crh-1p4::gfp) upstream of the translation start sequence of crh-1 gene. We found that sequences 2601, 2411, and 1094 bp regulatory sequences drove limited-expression possibly due to lack of certain cis-regulatory sequences [21,23] and showed expression specifically in three pairs of sensory neurons in the head amphid organs, including in the AWC, ASE, and AFD neurons, consistent with previous observations [24]. However, we also routinely detected weak GFP expression in the ASI sensory neurons (Fig 3A and 3B). Strong GFP expression of the crh-1p4::gfp transgene was detected exclusively in the ASI neurons throughout larval development (Fig 3A and 3B). Expression of wild-type crh-1 cDNA sequences driven by crh-1p1 or crh-1p4 promoters were both partially sufficient to suppress L2d formation of crh-1 mutants (Fig 3C), suggesting that crh-1 acts in the ASI sensory neurons. The morphology and dye-filling properties of the ASI neurons were not altered by the crh-1 mutation (n > 50 each) (S9 Fig), suggesting that CRH-1 does not play a general role in ASI neuron development.

To confirm these findings, we next rescued crh-1 mutant phenotypes by expressing crh-1 cDNA under the control of cell-specific promoters, including srg-47 (ASI), ceh-36Δ (AWC), and ttx-1 (AFD) [17]. Whereas expression of crh-1 in single AWC or AFD neurons did not rescue the L2d formation defects of crh-1 mutants, ASI-specific expression rescued the phenotype to the same extent as expression driven under the endogenous crh-1 promoter (Fig 3D). ASI-specific expression of crh-1 also fully rescued the flp-8 expression defects of crh-1 mutants (Fig 3E). Taken together, we conclude that the crh-1 gene is expressed in and functions in the ASI neurons to regulate L2d formation.

daf-7 TGF-β expression is decreased in the ASI neurons of crh-1 mutants

Identification of the ASI neurons as the site of action for CRH-1 prompted us to investigate whether CRH-1 also plays a role in regulating the expression of the daf-7 transforming growth factor-β (TGF-β), which is expressed mainly in ASI, and which plays a major role in dauer formation [12,13]. A feasible hypothesis for the inappropriate entry of crh-1 mutants into the L2d stage is that the expression of daf-7 is downregulated in crh-1 mutants under non-dauer-inducing conditions. Previously, the daf-7 expression has been monitored using either an integrated daf-7p:gfp reporter gene, or by single-molecule fluorescence in situ hybridization (smFISH) [16,36]. We found that the daf-7p:gfp reporter exhibited decreased expression in the ASI neurons of crh-1 mutants (Fig 4A and 4B). The ASI-specific expression of crh-1 restored daf-7p:gfp expression in crh-1 mutants (Fig 4B). We next used smFISH to accurately examine the endogenous transcriptional activity of daf-7 and observed that the fluorescence intensity of daf-7 FISH probes in the ASI neurons was also decreased in crh-1 (Fig 4A and 4C). Moreover, ascr#5 further decreased daf-7 transcriptional activity (Fig 4C). These results demonstrate that CRH-1 is necessary to maintain a robust expression of daf-7 in the ASI neurons.

Similar to daf-7 expression, the expression of a candidate str-3 chemoreceptor gene in the ASI neurons is regulated by exposure to pheromones [15,36]. Expression of an str-3p:gfp reporter gene was strongly decreased in crh-1 mutants. This gene expression phenotype was
Fig 3. crh-1 is expressed and acts in the ASI neurons to regulate ascr#5-mediated L2d formation. (A) The percentage of adult transgenic animals expressing crh-1p1: gfp, crh-1p2: gfp, crh-1p3: gfp or crh-1p4: gfp reporter construct in the indicated neurons is shown. The strength of GFP expression is indicated by the number of + symbols. At least two independent extrachromosomal lines for each construct were examined. n ≥ 60 for each. (B) Representative images of adult or L1 larval transgenic animals expressing gfp expression under the control of crh-1 promoter1 (crh-1p1) and crh-1 promoter4 (crh-1p4). Adult animals were dye-filled with DiI. Scale bar: 10 μm. (C and D) Percent of L2d formed by animals of the indicated genotypes when grown in the presence of live OP50 food and ascr#5 pheromone. N ≥ 6 for each. Error bars indicate SEM. *, **, and *** indicate different from crh-1 mutants at
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again rescued by ASI-specific expression of crh-1 (Fig 4D and 4E). Taken together, these results are consistent with the hypothesis that CRH-1 mediates transcriptional regulation of daf-7 TGF-β and STR-3 GPCR gene expression in the ASI neurons.

daf-7 TGF-β signaling is required for the ascr#5-mediated L2d formation of crh-1 mutants
daf-7 null mutants form dauers constitutively (Daf-c) regardless of environmental conditions, although other conditions and mutants in which daf-7 expression is perturbed result in less penetrant dauer-related phenotypes [12]. We, therefore, investigated whether reduced daf-7 expression plays a role in the inappropriate induction of L2d larvae in crh-1 mutants. As expected, 100% of daf-7 mutants formed L2d with abundant live OP50 food and high concentrations of ascr#5 (Fig 5A). Compared to the transient L2d formation of crh-1 mutants, daf-7 L2d larvae committed to dauer entry. In addition, flp-8 expression in the AVM neurons of daf-7 mutants initiated at 40 hAH and lasted throughout the L2d and dauer stages (Fig 5B and 5C). To confirm that DAF-7 acts downstream to CRH-1 in regulating dauer formation, we overexpressed DAF-7 specifically in the ASI neurons of crh-1 mutants and found that it partially rescued the L2d formation phenotype of daf-7 mutants (Fig 5E), further supporting a role of the ASI neurons in the L2d formation of crh-1 mutants.

It was previously shown that the Daf-c phenotype of daf-7 mutants was suppressed by loss-of-function mutations of the daf-3 SMAD or daf-5 SNO/SKI transcription factor genes [37,38], whereas mutation of daf-16 represses Daf-c mutations in the insulin signaling pathway [14]. We found that the loss of daf-5 function fully suppressed the L2d phenotype of crh-1 mutants, whereas the loss of daf-3 or daf-16 function did not affect the L2d formation of crh-1 mutants (Figs 5F and S10). These results suggest that daf-5 acts downstream of crh-1 in ascr#5-mediated L2d formation and that the daf-7 TGF-β signaling pathway mediates the L2d formation of crh-1 mutants.

CRH-1 directly regulates daf-7 expression via a conserved CRE motif
Like mammalian CREB, CRH-1 can act as a sequence-specific transcription factor to regulate gene expression via cAMP-response elements (CRE; TGACGTCA) [39] (Fig 6A). To determine whether CRH-1 directly regulates the expression of daf-7 in the ASI neurons, we first analyzed the promoter regions of the daf-7 gene. Although we could not identify any DNA sequences which showed exact matches to the CRE site within 3.1 kb upstream of the daf-7 initiator codon, we identified ten DNA sequences that contain the consensus ACCT CRE core sequence within this sequence (Fig 6B). We mutated these sequences in the context of the daf-7p:gfp reporter gene, and found that mutations in one specific sequence, “GTACGTAC” located ~2380 bp upstream of the translational start site caused decreased daf-7p:gfp expression in the ASI neurons (Fig 6B and 6C). This mutation did not affect the faint reporter expression we observed in ASJ neurons (S11 Fig), indicating the specificity of this motif in the regulation of ASI expression of daf-7 [16]. Moreover, the decreased ASI daf-7 expression by mutations of the CRE site was not altered in crh-1 mutant adults (Fig 6D). These results support our hypothesis that CRH-1 binds this putative CRE in the daf-7 promoter to regulate its expression in ASI neurons.
Fig 4. *crh-1* mutants exhibit a decrease of *daf-7* expression in the ASI neurons. (A) Representative images of L1 larval wild-type or *crh-1* mutant animals expressing *daf-7::gfp* transgene in the presence (+) or absence (-) of *ascr*#5. Shown is GFP expression or *daf-7* smFISH signal in the ASI neurons. All images for wild-type versus *crh-1* mutant animals were taken with identical camera settings, including the same exposure time. Anterior is to the left. Scale bar: 50 μm. (B) The relative level of *daf-7::gfp* fluorescence in the ASI neurons of L1 larval animals of the indicated genotypes in the absence (-) of *ascr*#5. (C) Scatter plot of fluorescence intensity of *daf-7* smFISH signal in the ASI neurons of wild-type or *crh-1* mutant animals in the presence (+) or absence (-) of *ascr*#5. The median is indicated by a horizontal line. Each dot is the fluorescence intensity of a single neuron; n ≥ 30 neurons total each, at least two independent experiments. (D) Representative images of L1 larval wild-type or *crh-1* mutant animals expressing *str-3::gfp* in the ASI neurons. Scale bar: 10 μm. (E) The relative level of *str-3::gfp* fluorescence in the ASI neurons of L1 larval animals of the indicated genotypes in the absence (-) of *ascr*#5. n ≥ 30 for each. Error bars indicate SEM. ***, †††, or ### indicates different from wild-type, absence of *ascr*#5, or *crh-1* mutants at p < 0.001, respectively (student t-test). Arrowheads indicate the ASI neurons (A and D).

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Fig 5. daf-7 mediates the ascr5-mediated L2d formation of crh-1 mutants. (A) Percent of L2d formed by animals of the indicated genotypes when grown in the presence of live OP50 food and ascr5 pheromone. N ≥ 4 for each. (B) Representative images of wild-type (left column), crh-1 mutant (middle column), or daf-7 mutant animals expressing of flp-8p-gfp in a subset of touch receptor neurons in the presence of live OP50 food and ascr5 pheromone. The color-matched boxed regions of the second column each are shown at higher magnification. hAH, hour after hatching. Scale bars: 50 μm. Images of wild-type and crh-1 mutants are originated from Fig 2B. (C) Percent of flp-8p-gfp expression in the AVM neurons by animals of the indicated genotypes when grown in the presence of live OP50 food and ascr5 pheromone. n ≥ 30 for each. *** indicates different from wild-type at p < 0.001 (student’s t-test). (C-E) Percent of L2d formed by animals of the indicated genotypes when grown in the presence of live OP50 food and ascr5 pheromone. N ≥ 8 (C), 6 (D), 4 (E) for each. Error bars represent the SEM. *** or ††† indicates different from crh-1 or daf-7 mutants at p < 0.001, respectively (student’s t-test).

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Fig 6. CRH-1 directly binds a CRE motif in the *daf-7* promoter. (A) Shown are the conserved mammalian CRE motif and the newly identified CRE motif in the *daf-7* promoter. (B) The percentage of transgenic animals expressing *daf-7p::gfp* reporter construct in the ASI neurons is shown. GFP fluorescence was observed either in wild-type or *crh-1* mutant animals. The strength of GFP expression is indicated by the number of + symbols. Wild-type nucleotides are indicated in red, mutated nucleotides in blue. An arrowhead indicates *daf-7p*-mutated CRE promoter. At least two independent extrachromosomal lines for each construct were examined. N \( \geq 30 \) for each. (C) Representative images of L1 larval wild-type or *crh-1* mutant animals expressing *daf-7p::gfp* or *daf-7p*-mutated CRE::gfp in the ASI neurons. Scale bar: 50 \( \mu \)m. (D) Scatter plot of fluorescence intensity of *daf-7p::gfp* fluorescence in the ASI neurons of adult wild-type or *crh-1* mutant animals with (+) or without (-) of CRE site mutation. The median is indicated by a horizontal line. Each dot is the fluorescence intensity of a
To confirm the direct binding of CRH-1 to this CRE motif in vitro, we performed electrophoretic mobility shift (EMSA) assays on cultured HEK293T cells. Specific CRE binding activity was evident in GFP-CRH-1 overexpressing nuclei, and absent in control nuclear extract (Fig 6E). Furthermore, this CRE binding activity was significantly diminished by co-incubation with wild-type non-biotinylated probes, whereas mutant unlabeled probes failed to effectively compete (Fig 6E). While these results need to be further confirmed in vivo, these results indicate that CRH-1 can directly bind to the CRE motif identified in the daf-7 promoter sequence, which, in turn, may enhance daf-7 expression in the ASI neurons.

Thus, we have identified CRH-1 as a critical mediator of ASI state in developing C. elegans. CRH-1 promotes the expression of daf-7 in the presence of pheromone and abundant food to suppress inappropriate passage through L2d, which ultimately delays reproductive maturation of the animal and may be disadvantageous to fitness.

Discussion

Dauer entry in C. elegans is gated by sequential developmental decisions, first, at L1 to transition to either L2 or L2d, and second, a commitment to either diapause or reproductive growth (Fig 1A). Precociously or inadvertently entering dauer, has the potential to carry a large fitness cost for an animal, given the rapid development and fecundity of C. elegans. Here we have shown that the CREB homolog CRH-1 is a key regulator of the first decision, whether or not to enter L2d. crh-1 mutants inappropriately express the L2d marker gene flp-8 in ALM, AVM, and PLM neurons and transiently form L2d larvae in non-dauer-inducing conditions. However, crh-1 mutants do not commit to dauer entry in the absence of additional environmental cues and thus ultimately make the correct developmental decision. Our data support a model in which CRH-1 positively regulates dauer-inhibiting daf-7 TGF-β expression in the ascr#5-sensing ASI neurons (Fig 7). The loss of crh-1 function leads to decreased daf-7 TGF-β expression, which provides a sensitized genetic background for further ascr#5-mediated daf-7 TGF-β repression via SRG-36/37 GPCRs (Fig 7). When the daf-7 expression is eliminated, animals constitutively enter the dauer state, and thus daf-7 expression serves as a critical rheostat for the determination of C. elegans developmental trajectory [12,13,15]. Our results demonstrate that worms exploit distinct strategy and molecules at different dauer decision points and CRH-1 acts in ASI and gates the initial L1-to-L2d decision in response to pro-growth environmental signals (food), prior to dauer commitment, and thus crh-1 mutant provides a unique opportunity to study the under-explored L1 to L2/L2d transition. Previously, Lakhina and coworkers showed that crh-1 mutants develop more slowly than wild-type animals at 20˚C in the presence of food due to a change of basal CRH-1 activity [23]. Altogether, these results suggest that CRH-1 regulates distinct aspects of C. elegans development.

We have shown that an abundance of live OP50 food and high (2 μM) concentrations of ascr#5 (the most abundant ascaroside secreted when animals have grown at 20˚C) [7], which may reflect a natural condition of crowding at 20˚C but high food, do not induce dauer formation (L2d to dauer transition) but do induce L2d formation (L1 to L2d transition) in crh-1 mutants. These results suggest CRH-1 is necessary to give the appropriate weight to pro-growth food cues in the environment such that the presence of pheromone in the environment is sufficient to promote L2d formation but not dauer commitment [4]. Moreover, ascr#5 but
not ascr#2 or ascr#3 causes L2d formation in crh-1 mutants, indicating that specific, but not all, pheromone component(s) may affect the decision for L1 to transition to L2 or L2d and worms weigh pheromone signals differently at different dauer decision points. ascr#5 is the most potent component of the dauer inducing ascaroside pheromones [7] and is produced in higher amounts in well-fed worms than in starved worms, indicating that ascr#5 may be a bona fide indicator of healthy populations or favorable environments [11]. Despite this hypothesis, very high (6 μM) concentrations of ascr#5 can elicit L2d formation in wild-type animals, indicating that crh-1 mutants are precocious in L2d entry. Given that ascr#5 further reduces daf-7 expression in the ASI neurons of crh-1 mutants, we infer that CRH-1 acts as a food sensor to set the basal level of daf-7 expression. Furthermore, the two-step process to enter dauer likely allows animals to be more flexible and adjust to rapidly changing environments [40].

Previously, a large number of genes have been shown to be differentially expressed in dauer larvae [41,42]. Interestingly, expression of most, but not all, flp genes, including flp-8 are up-regulated during dauer development [42], and several flp genes have shown to be involved in dauer development and behaviors [42,43]. Since the flp-8 expression in the AAP-TRN begins at the L2d stage and is maintained throughout the dauer stage, the function of FLP-8 neuropeptide in the AAP-TRN of L2d and dauer larvae is intriguing. Although the exact function of FLP-8 has not been determined, a family of FLP neuropeptides appears to inhibit overall circuit activity [44]. Previously, Chen and Chalfie [45] showed that dauer larvae exhibited reduced touch sensitivity. Thus, it is possible that FLP-8 expression in the AAP-TRN could inhibit circuit activity underlying touch sensation, which leads to decreased mechanosensation in dauer and likely L2d larvae. It would be the next question to be explored in the future.

Fig 7. A model of the role of CRH-1 in ascr#5-mediated L2d formation. CRH-1 positively regulates dauer-inhibiting daf-7 TGF-β expression in the ascr#5-sensing ASI neurons. In crh-1 mutants, the decreased daf-7 TGF-β expression provides a sensitized genetic background for further ascr#5-mediated daf-7 TGF-β repression via srg-36/37 GPCRs which leads to transient L2d formation and flp-8 expression of AAP-TRN.

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Transforming growth factor β (TGF-β) signaling mediates diverse physiological roles, including cell differentiation, morphogenesis, tissue homeostasis, regeneration, and immune response, while the misregulation of TGF-β signaling promotes distinct disease states such as cancer progression (see review by Massague [46]). While cellular response pathways to secretion and diffusion/trafficking of the TGF-β ligand are well-characterized, relatively little is known about the regulation of its expression [47]. In C. elegans, TGF-β signaling regulates body size, innate immunity, aging, cell fate specification, differentiation, and dauer formation (see review by Savage and Padgett [48]). For example, TGF-β signaling regulates CREB activity in the hypodermis to mediate reproductive aging, and crh-1 acts downstream of sma-3 R-SMAD and regulates wrt-10 Hedgehog-related signaling factor[49]. Although the downstream components of TGF-β signaling are evolutionarily conserved and well-characterized, upstream transcription factors and cis-regulatory elements by which expression of TGF-β genes is controlled are not as well characterized. Here, we have shown that the CREB homolog CRH-1 directly binds a novel CRE in the daf-7 TGF-β promoter, which must integrate multiple external and internal signals and positively regulates its expression in the ASI neuroendocrine cells. Future studies are needed to identify potential CRH-1 co-factors, which may help to refine or modulate its actions at the daf-7 promoter. Similar to mammalian CREB, C. elegans CRH-1 functions in many developmental and behavioral processes, including longevity, learning and memory, and now, developmental plasticity. Thus, CREB acts as a determinant of a context-dependent role of TGF-β signaling and provides a useful model to study TGF-β gene expression.

Materials and methods

Strains and constructs

The N2 Bristol strain was used as wild-type. All strains were maintained and grown on NGM agar plates seeded with E. coli OP50 at 20˚C [50]. All used strains in this study are listed in S1 Table. Promoters and cDNA used for rescue experiments of crh-1 and daf-7 were described in Park et al. [51]. The daf-7 promoter (3127bp) for expression and CRE site analysis was produced by polymerase chain reactions using forward and reverse primers (see S2 Table) and subcloned into a pPD95.77 vector. The nhr-246 promoter (3067bp) for expression was produced by polymerase chain reactions using forward primer 5’-aaaaaacctgcaggCACCGATGCG TTGTTATAGG-3’ and reverse primer 5’-ttttttcccgggTGTTGAAA TTGAAAATTATTTTGAA -3’ [34] and subcloned into a pPD95.77 vector. All plasmids were injected at 0.1 ng (flp-8 expression analysis), 10 ng (rescue experiments), or 50 ng (expression analysis) with unc-122p::dsRed as an injection marker.

Pheromone preparation and assay plates

Crude pheromone was prepared following the protocol described by Golden and Riddle (1984). Before assay, each crude pheromone batch was tested for its efficiency to induce dauer formation. The assay plates containing crude pheromone were prepared by spreading pheromone and drying for 3–4 h. The ascaroside pheromone components were chemically synthesized following Butcher et al. [6,7]. Before use, pheromone was diluted with dH2O from a 3 mM stock solution of pheromone in 100% ethanol. For the assay plates containing synthetic ascarosides, the pheromone solution was mixed with agar solution before pouring.

L2d formation assay

Synchronized adult animals were placed onto 3.5 cm assay plates seeded with live OP50 and containing ethanol or pheromone at 25˚C and allowed to lay over 50 eggs for 1–2 h. Then,
adult animals were removed, and assay plates were placed at 25°C for 40 h before observing the morphology of worms or marker gene expression and counting as L2d larvae. Dauer larvae and L2d larvae were selected based on the survival with 1% SDS treatment.

Dauer formation assay

A dauer formation assay was performed as previously described by Golden and Riddle [4] and Neal et al. [52]. Synchronized adult animals were allowed to lay eggs at 25°C for 4–5 h on 3.5 cm assay plates. After the plates contained approximately 65–85 eggs/plate, adult animals were removed, and the plates were placed at 25°C for 68–72 h. Dauer and non-dauer animals were identified by visual inspection. Assay plates were made with Noble agar (BD Biosciences) lacking peptone. Heat-killed OP50 bacteria were seeded on each plate. For each experiment, all strains were assayed in parallel in at least four independent experiments.

daf-7 smFISH

daf-7 smFISH was performed as previously described by Meisel et al. [16] by using the Stellaris smFISH fluorescent probe and buffer sets (Biosearch). Worms were washed in 1XPBS and fixed in 3.7% formaldehyde in PBS for 45 min at 4°C and rinsed twice with 1XPBS. Then, worms were immersed in 70% ethanol for 24 h at 4°C and washed by buffer A (Biosearch) for 5 minutes at room temperature. Then, a hybridization buffer containing the daf-7 probe (125 nM, Biosearch) was added and incubated overnight at 30°C. Worms were washed twice with buffer A. Then, worms were mounted onto glass slides by mounting medium (Sigma). All smFISH images were acquired by a Zeiss Axio Imager using 63x objectives and a CCD camera (Hamamatsu). Quantification was performed using ImageJ software to extract mean intensity and regions of interest. The relative mean intensity of fluorescence was normalized by area.

Quantification of GFP expression

For GFP quantification, the worms were anesthetized in sodium azide on an agar pad, and GFP fluorescence was observed with a Zeiss Axio Imager using 40x (for the adult stage) and 63x (for L1) objectives and a CCD camera (Hamamatsu). The relative expression level of GFP was measured at each developmental stage. The relative GFP levels were rated from 1 (dim) to 5 (bright) by visual inspection, and these values were confirmed using Image J software.

SEM (Scanning Electron Microscope) imaging

Samples were prepared by critical point dryer (Leica EM CPD300) after sequential ethanol dehydration. Dried samples were coated with gold using a sputter coater and then were imaged with a HR FE-SEM (Hitachi, SU8020, CCRF DGIST).

Electrophoretic mobility shift assay (EMSA)

Nuclear extract preparation from HEK293T cells and in vitro EMSA reactions were carried out as previously described Kim et al. [53], and biotinylated EMSA probes and unlabeled competitor probes were synthesized by using a LightShift Chemiluminescent EMSA Kit (Thermo Scientific). The daf-7p-WT-CRE and daf-7p-mut-CRE probes with the following sequences were employed:

daf-7p-WT-CRE: 5’ attaggtagctagctaatattaggtagctagctaatattaggtagctagctaa 3’ (60 mer)
daf-7p-mut-CRE: 5’ attaggtagttttaaatattaggtagttttaaatattaggtagttttaaatat 3’ (60 mer)
Statistical analysis

Statistical analysis were performed using GraphPad Prism 8.0 and Microsoft Excel 365. Data used for statistical analysis are in the S1 Data file.

Supporting information

S1 Fig. *crh-1* mutants form transient predauer (L2d) larvae. Shown are the survival rate of wild-type, *crh-1* mutant and *daf-7* mutants animals under 1% sodium dodecyl sulfate (SDS) treatment. Two independent assays with n > 50.

(SPDF)

S2 Fig. *daf-7* mutants exhibit strong expression of *flp-8* in the AVM in non dauer-inducing conditions. Shown are images of *flp-8p::gfp* expression in the AVM in *daf-7* mutants at 45 and 60 hAH. The boxed regions of the first column are shown on the right columns at higher magnification. hAH, hour after hatching. Scale bars: 50 μm.

(SPDF)

S3 Fig. *crh-1* mutants exhibit expression of *flp-8* in the AAP-TRN in dauer-inducing conditions. Percent of *flp-8p::gfp* expression in the AVM, ALM and PLM neurons by wild-type or *crh-1* mutant animals when grown in the presence of heat-killed OP50 food and ascr#5 pheromone. n ≥ 30 for each. hAH, hour after hatching. Error bars indicate SEM. * and ** indicate different from wild-type at p < 0.05 and p < 0.01, respectively (student t-test).

(SPDF)

S4 Fig. ascr#5 is required for *flp-8p::gfp* expression in the AAP-TRN in wild-type or *crh-1* mutant animals. Percent of *flp-8p::gfp* expression in the AVM, ALM and PLM neurons by wild-type or *crh-1* mutant animals when grown in the presence of heat-killed OP50 food (left) or live OP50 (right). n ≥ 30 for each. hAH, hour after hatching. Error bars indicate SEM. * indicates different from wild-type at p < 0.05 (student t-test).

(SPDF)

S5 Fig. *crh-1* mutants exhibit expression of *nhr-246* in the intestine in non dauer-inducing conditions. Shown are left images of *nhr-246p::gfp* expression in the intestine of *crh-1* mutants at 45, 50, and 55 hAH. The boxed regions of the second column are shown on the right columns at higher magnification. Scale bars: 50 μm. Shown is the right panel of the percent of *nhr-246p::gfp* expression in the intestine by wild-type or *crh-1* mutant animals when grown in the presence of live OP50 food and ascr#5 pheromone at 45, 50, and 55 hAH. hAH, hour after hatching. n ≥ 20 for each. Error bars indicate SEM. ** indicates different from wild-type at p < 0.001 (student t-test).

(SPDF)

S6 Fig. *crh-1* mutants exhibit normal or weakly decreased pheromone-mediated dauer formation. Percent of dauer formed by wild-type or *crh-1* mutant animals when grown in the presence of heat-killed OP50 food and ascr#5, ascr#2, or ascr#3 pheromone.

(SPDF)

S7 Fig. *crh-1* mutants form L2d larvae in a non-dauer inducing condition. Percent of L2d formed by wild-type or *crh-1* mutant animals when grown in the presence of live OP50 food and crude pheromone. N ≥ 5 for each. Error bars indicate SEM. *, **, and *** indicate different from wild-type at p < 0.05, p < 0.01, and p < 0.001, respectively (student t-test).

(SPDF)
S8 Fig. Morphology of mouths of wild-type L3 larva, \textit{crh-1} mutant L2d larva, and wild-type dauer larva. Shown are images of scanning electron microscopy. Scale bar: 5\,$\mu$m (top) and 100\,$\mu$m (bottom). (PDF)

S9 Fig. \textit{crh-1} mutants do not form L2d larvae in the presence of ascr#2. Percent of L2d formed by wild-type or \textit{crh-1} mutant animals when grown in the presence of live OP50 food and three different concentrations of ascr#2 pheromone. N \geq 5 for each. (PDF)

S10 Fig. \textit{crh-1} mutants do not exhibit \textit{flp-8} expression in the AAP-TRN in response to ascr#2 and ascr#3. Percent of \textit{flp-8p::gfp} expression in the AVM, ALM and PLM neurons by wild-type or \textit{crh-1} mutant animals when grown in the presence of live OP50 and ascr#2 or ascr#3. n \geq 30 for each. hAH, hour after hatching. Error bars indicate SEM. (PDF)

S11 Fig. The ASI morphology and dye-filling properties are not altered in the \textit{crh-1} mutants. Shown are images of wild-type or \textit{crh-1} mutant animals expressing \textit{gpa-4p::gfp} transgene or stained with DiI. Scale bar: 10\,$\mu$m. (PDF)

S12 Fig. \textit{daf-16} mutation does not affect the L2d formation of \textit{crh-1} mutants. Percent of L2d formed by animals of the indicated genotypes when grown in the presence of live OP50 food and ascr#5 pheromone. N \geq 4 for each. Error bars indicate SEM. N.S. not significantly different (one-way ANOVA with Bonferroni’s post hoc tests). (PDF)

S13 Fig. Mutation in CRE motif of \textit{daf-7} promoter does not affect \textit{daf-7} expression in the ASJ neurons. The percentage of transgenic animals expressing \textit{daf-7p::gfp} reporter construct in the ASJ neurons is shown. GFP fluorescence was observed either in wild-type or \textit{crh-1} mutant animals. The Strength of GFP expression is indicated by the number of + symbols. Wild-type nucleotides are indicated in red, mutated nucleotides in blue. An arrowhead indicates \textit{daf-7p}-mutated CRE promoter. At least two independent extrachromosomal lines for each construct were examined. n \geq 30 for each. (PDF)

S1 Table. List of strains used in this work. (DOCX)

S2 Table. List of primers and probes used in \textit{daf-7} related work. (DOCX)

S1 Data. Excel of raw data file of all relevant figures. (XLSX)

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