Calcineurin and Protein kinase G regulate \textit{C. elegans} behavioral quiescence during locomotion in liquid

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**Abstract**

**Background:** Most rhythmic motor behaviors in nature are episodic i.e. they alternate between different behavioral states, including quiescence. Electrophysiological studies in invertebrate behavioral switching, maintenance and quiescence have elucidated several neuronal mechanisms that generate a temporal pattern in behavior. However, the genetic bases of these processes are less well studied. We have previously uncovered a novel episodic behavior exhibited by \textit{C. elegans} in liquid media where they alternate between distinct phases of rhythmic swimming and quiescence. Here, we have investigated the effect of several genes and their site of action on the behavioral quiescence exhibited in liquid by the nematode \textit{C. elegans}.

**Results:** We have previously reported that high cholinergic signaling promotes quiescence and command interneurons are critical for timing the quiescence bout durations. We have found that in addition to command interneurons, sensory neurons are also critical for quiescence. We show that the protein phosphatase calcineurin homolog \textit{tax-6} promotes swimming whereas the protein kinase G homolog \textit{egl-4} promotes quiescence. \textit{tax-6} expression in the sensory neurons is sufficient to account for its effect. \textit{egl-4} also acts in multiple sensory neurons to mediate its effect on quiescence. In addition our data is consistent with regulation of quiescence by \textit{egl-4} acting functionally downstream of release of acetylcholine (ACh) by motor neurons.

**Conclusions:** Our study provides genetic evidence for mechanisms underlying the maintenance of a behavioral state operating at multiple neuronal levels through the activities of a kinase and a phosphatase. These results in a genetically tractable organism establish a framework for further dissection of the mechanism of quiescence during episodic behaviors.
In addition to the quiescent state exhibited by worms in liquid [9], two other behavioral quiescent states have been studied in C. elegans [11-13]. Movement of C. elegans stops when they undergo lethargus, a developmental state that occurs before each of the four molts during its life cycle [11,12]. This quiescent state exhibits characteristics that resemble sleep in other organisms [11]. On agar plates with high-quality food, movement and feeding of an adult worm is interjected by stops, a behavioral characteristic that has parallels to satiety in mammals [13]. Molecularly, two different signaling systems have been identified to influence these quiescent states. Insulin/TGF-β and EGF signaling induce quiescence on plates with high quality food and during lethargus respectively [11-13]. In both quiescence paradigms, protein kinase G (PKG) activity in sensory neurons is required to promote quiescence. Eventually the activity of PKG at the sensory neuronal level must modulate activities of the motor and/or interneurons to bring about quiescence. How PKG activity in sensory neurons drives quiescence at the motor/interneuron levels is not known. Also it remains to be seen how much overlap exists in the genetics of the three behavioral quiescence states that have been described. The unique features of long term swimming behavior, ease of genetic analysis and the knowledge of the pattern of synaptic connectivity of the entire nervous system [14] that allow mapping of molecular to neuronal correlates underlying behavior makes C. elegans a suitable model to dissect the molecular mechanisms underlying this long-term behavioral pattern.

To further investigate the underlying mechanism(s) of quiescence in liquid, we examined the cellular site of action of several genes that had important effects on swim/quiescence cycling. Several of these do not act at the motor neuron-muscle circuitry but instead act in sensory neurons. We found that maintenance of swimming requires calcineurin activity in the sensory neurons. The PKG homolog egl-4 promotes quiescence by acting at two levels: in the sensory neurons and also downstream of command interneurons. Our analysis suggests that there may be two mechanisms by which swimming motions may be stopped, one by blocking output of the motor system at the level of motor neurons and muscles, a mechanism that appears to come into play when worms swim in liquid, and one by inhibition of initiation of swimming motions upstream of this block, which is strongly influenced by sensory input.

Results
PKG is required for maintenance of the quiescent state during swimming in liquid
The PKG homolog of C. elegans, egl-4, regulates long-term olfactory adaptation [15], different locomotory states (roaming and dwelling) [16] and promotes longer duration of quiescence bouts under different paradigms [11-13]. Though it is widely expressed in the neuromuscular system and has been implicated in modulation of acetylcholine release at neuromuscular junctions [17], the pleiotropic effects of egl-4 have been mostly attributed to its activity in sensory neurons [15-18]. We have previously shown that loss-of-function alleles of egl-4 affected maintenance of the naturally occurring quiescent bouts that occur with regular periodicity when a worm is swimming in liquid [9]. Indeed, when compared to the wild type animals, the median fraction of egl-4(ks60) animals that remained in quiescence during the second phase of swimming (phase one is defined as the first ninety minutes from the start of the assay and the subsequent 90 minutes is defined as the second phase of the assay) was significantly lower (wild type: 0.21; egl-4(ks60): 0.14, p < 0.0001, Mann Whitney test, trends are shown in Figure 1B). Here we show that the loss-of-function mutant, egl-4(ks60), suppresses increased quiescence resulting from including 0.01 mM aldicarb in the assay medium (Figure 1A, B). The median quiescent bout durations of egl-4(ks60) in 0.01 mM aldicarb did not differ from egl-4(ks60) in buffer alone (Figure 1A, Mann Whitney U test, p = 0.5683). This suppression was absent at higher doses (0.1 or 1 mM) aldicarb (data not shown). However egl-4 mutants still went into quiescence albeit for a shorter duration in 0.01 mM aldicarb. Taken together these data suggest that increased quiescent bout durations in liquid due to increased cholinergic activity require egl-4 function.

The increased cholinergic activity during quiescence appeared by several criteria to be at the neuromuscular junction [9]. During quiescence in liquid body-wall muscles are uniformly contracted. Quiescence induced by aldicarb was enhanced by mutation in dgk-1, which increases ACh release from motor-neurons [9,19]. Mutations resulting in reduced synthesis/release of GABA by GABAergic motor neurons, which opposes the action of ACh on muscles, also increased quiescence, whereas the GABA agonist muscimol promoted swimming. Since egl-4 is necessary for ACh-mediated quiescence, we conclude that egl-4 has an activity that promotes quiescence functionally downstream of ACh release by motor neurons.

Further evidence for PKG activity downstream of motor neurons in regulating quiescence
In order to further investigate the quiescence mechanism that appeared to require egl-4 function downstream of ACh release by motor neurons, we examined the relationship of egl-4 function and additional components of the motor circuitry. We had previously found that mutations resulting in defects in GABAergic signaling
result in increased fraction of worms in quiescence [9]. To determine whether quiescence induced by decreased GABA required \textit{egl-4}, we examined double mutants with the GABA-deficient mutant \textit{unc-25(sa94ts)} [20]. Unlike its effect on quiescence induced by Ach, \textit{egl-4 (ks60)} had little effect on increased quiescence due to loss of GABA activity. Although the median quiescent bout duration of \textit{unc-25(sa94)} was (6, mean = 9.6 ± 0.7 minutes) that was slightly longer than the median quiescent bout duration of \textit{unc-25(sa94); egl-4(ks60)} (4.0, mean = 7.5 ± 0.7 minutes) (\textit{p} < 0.0001, Mann-Whitney U test), the distribution of the quiescent bout durations for \textit{unc-25(sa94)} and \textit{unc-25(sa94); egl-4(ks60)} exhibited considerable overlap (Additional File 1). Worms of both these genotypes exhibited long quiescent bouts from which they did not initiate swimming reflected by their overlapping trends of fraction of worms in quiescence and distribution of quiescent bouts (Figure 1C and Additional File 1). We conclude that loss of function of \textit{egl-4} was not sufficient to suppress the increased quiescence due to \textit{unc-25(sa94)} mutation (Table 1) indicating that \textit{egl-4} primarily acts upstream of GABA signaling in regulating duration of quiescence.

Figure 1 PKG promotes quiescence. A) Scatterplot of quiescent bouts for wild type, wild type in presence of 0.01 mM aldicarb (n = 23), wild type in 0.6 OD \textit{E.coli} Op50 (n = 30), \textit{egl-4(ks60)} (n = 24), \textit{egl-4(ks60)} in 0.01 mM aldicarb (n = 23), \textit{egl-4(ks60)} in 0.6 OD Op50 (n = 24), \textit{nmr-1(ak4)} (n = 33), \textit{egl-4(ks60); nmr-1(ak4)} (n = 24), \textit{egl-4(ks62)} (n = 24), \textit{egla(ks66)} (n = 46) and \textit{egla(m77)} (n = 22). The medians are shown by a black or a red horizontal line (for \textit{akIs9}). *** significantly different from wild type, ** significantly different from \textit{nmr-1(ak4)}, ### significantly different from \textit{unc-25(sa94ts)}. Mann-Whitney U test for each pair. NS nonsignificant difference at p < 0.001. B) Fraction of worms in quiescence versus time for wild type (n = 23), wild type in presence of 0.01 mM aldicarb (n = 23), \textit{egl-4(ks60)} (n = 24), \textit{egl-4(ks60)} in 0.01 mM aldicarb (n = 23). C) Fraction of worms in quiescence versus time for wild type (same as Figure 1B), \textit{unc-25(sa94)} (n = 24), \textit{egl-4(ks60)} (same as Figure 1B), \textit{egl-4(ks60);unc-25(sa94)} (n = 24), \textit{akIs9} (n = 21) and \textit{akIs9;egl-4(ks60)} (n = 27).
were partially suppressed by egl-4(ks60) (Figure 1A, C). Taken together, these observations suggest a role of egl-4 genetically downstream of command interneurons in sustaining quiescence. We also observed that suppression of the increased quiescence due to akIs9 by egl-4(ks60) was more potent during the first ~90 minutes of the assay (Figure 1C). Thus egl-4 independent mechanisms are required to shape the behavioral pattern beyond that time period. Overall, these results place a site of action for activity of egl-4 required for quiescence during swimming downstream of command interneurons and release of ACh by motor neurons, but upstream of GABA signaling.

Consistent with a role of command interneurons in timing quiescent bout duration, we found that a deletion in nmr-1, encoding NMDA receptor subunit expressed in the ring motor neuron RIM and all command interneurons except AVB [22], resulted in dramatically increased durations of quiescent bout (Figure 1A). The median quiescent bout durations of nmr-1(ak4); egl-4(ks60) were similar to egl-4(ks60) indicating egl-4 acts downstream of or within command interneurons in maintenance of quiescence (Figure 1A).

PKG acts in sensory neurons to modulate the quiescent state

To investigate whether egl-4 has an activity in the sensory neurons in regulating quiescence, we expressed an egl-4 transgene under different promoters in subsets of sensory neurons in egl-4(n479) loss-of-function mutant background. Like other egl-4 loss-of-function mutants (Figure 1A), these worms had significantly reduced duration of quiescence bouts compared to the wild type worms (Figure 2A, Wilcoxon signed rank test,

| STRAIN                        | Mean (min) | Median (min) | SD  | SE  | Number of bouts |
|-------------------------------|------------|--------------|-----|-----|-----------------|
| Wild type                     | 4.8        | 4.0          | 2.2 | 0.1 | 489             |
| cnb-1(jh103)                  | 20.5       | 1.0          | 49.0| 3.5 | 202             |
| tax-6(db60)                   | 5.5        | 1.0          | 13.3| 0.7 | 214             |
| tax-6(db60)(tax-6cDNA, muscle+) | 10.1      | 1.0          | 25.8| 1.6 | 261             |
| tax-6(db60)(tax-6 sensory+inter+) | 4.7      | 4.0          | 4.4 | 0.3 | 218             |
| tax-6(db60)(tax-6 sensory+inter-) | 3.9      | 3.0          | 3.3 | 0.2 | 192             |
| egl-4(ks60)                   | 2.7        | 2.0          | 2.5 | 0.2 | 173             |
| egl-4(ks60) in 0.6 OD E.coli(OP50) | 4.6      | 2.0          | 5.8 | 0.3 | 461             |
| Wild type in 0.6 OD E.coli(OP50) | 9.3        | 5.0          | 15.7| 1.1 | 189             |
| akIs9                         | 25.9       | 14.5         | 29.8| 2.4 | 158             |
| akIs9;egl-4(ks60)             | 8.9        | 5.0          | 11.5| 0.8 | 206             |
| nmr-1(ak4)                    | 9.0        | 7.0          | 6.1 | 0.5 | 170             |
| unc-25(sa94)                  | 9.6        | 6.0          | 11.7| 0.7 | 309             |
| unc-25(sa94);egl-4(ks60)      | 7.5        | 4.0          | 13.5| 0.7 | 352             |
| nmr-1(ak4);egl-4(ks60)        | 2.8        | 2.0          | 1.6 | 0.1 | 222             |
| osm-6(p811)                   | 7.0        | 5.0          | 7.4 | 0.3 | 479             |
| osm-3(p802)                   | 7.4        | 6.0          | 6.2 | 0.5 | 165             |
| osm-6(p811);egl-4(ks60)       | 3.8        | 3.0          | 2.9 | 0.1 | 452             |
| osm-3(p802);egl-4(ks60)       | 2.9        | 3.0          | 1.8 | 0.1 | 691             |
| nmr-1(ak4);osm-6(p811)        | 14.8       | 9.0          | 16.3| 1.1 | 221             |
| tax-2(p691);tax-4(ky89)       | 4.2        | 4.0          | 1.9 | 0.1 | 281             |
| egl-4(n479)                   | 2.1        | 1.0          | 1.5 | 0.3 | 30              |
| egl-4(n479) [odr-3p::egl-4]   | 3.0        | 3.0          | 2.3 | 0.5 | 19              |
| egl-4(n479) [odr-3p::egl-4]   | 4.4        | 3.0          | 4.5 | 0.7 | 48              |
| egl-4(n479) [odr-3p::egl-4]   | 2.9        | 2.0          | 2.1 | 0.7 | 9               |
| egl-4(n479) [odr-1p::egl-4]   | 3.7        | 1.0          | 4.2 | 0.6 | 47              |
| egl-4(n479) [odr-1p::egl-4]   | 3.6        | 2.0          | 3.6 | 0.5 | 55              |
| egl-4(n479) [tax-4p::egl-4]   | 6.6        | 2.0          | 15.1| 1.6 | 91              |
| egl-4(n479) [tax-4p::egl-4]   | 7.4        | 6.0          | 6.1 | 1.2 | 24              |
| egl-4(n479) [tax-4p::egl-4]   | 4.5        | 2.0          | 6.5 | 0.6 | 114             |

Summary statistics of quiescent bout durations of strains used in the study. The mean, median, standard deviation and the standard error of the mean for the quiescent bout durations in the first 180 minutes of the assay are shown. ‘min’ refers to minute.
When an egl-4 transgene was expressed under a tax-4 promoter there was a partial rescue of this mutant phenotype, the median quiescence bout did not differ from the wild type strain N2 (2 out of 3 transgenic lines) (Figure 2A). However, in all three lines there were long quiescent bouts (>8 minutes, NQ20: 17.6%, NQ21: 41.7%, NQ22: 9.7%) that were not observed in the wild type (3.4% of all bouts), or in egl-4(lf) mutants (0%). In contrast, no rescue was obtained by expression from odr-1 and odr-3 promoters (Figure 2A). Further, long quiescent bouts did not occur or were only somewhat increased under the odr-3 (NQ2: 5.2%, NQ5:14.5%, NQ7: 0%) and odr-1 (NQ11:10.6%, NQ13: 9.1%) promoters. All three promoters drive the expression of the egl-4 transgene in two sensory neurons AWB and AWC, while the tax-4 and odr-3 promoters drive expression in 8 and 3 additional pairs of sensory neurons respectively [11,15,16,23]. Thus tax-4 expressing neurons define a site of action of egl-4 in sustaining quiescent bouts.

Consistent with a role of sensory neurons in quiescence, we found that quiescence duration could be increased by defects in sensory physiology due to mutations in osm-6 or osm-3 (critical for proper cilium structure in sensory neurons). Both of these mutations result in abnormal sensory neuron structure [24] (Figure 2B, Table 1). Increased quiescence in osm-3 and osm-6 mutants was suppressed by egl-4(ks60) mutation (Figure 2B) indicating that egl-4 acts downstream or in sensory neurons. In addition we previously reported that inclusion of food (0.6 O.D. E.coli(OP50)) increased the quiescent bout durations [9]. The increased quiescence bout duration of wild type worms induced by inclusion of E. coli in the assay media was also significantly reduced in egl-4(ks60) mutants (Figure 1A, Table 1).

Since both sensory neurons and command interneurons contribute towards sustaining quiescence and thus there appeared to be two sites of action of egl-4, we asked whether these two pathways were independent. We assayed double mutants of nmr-1(ak4);osm-6(p811). The quiescence bouts in the nmr-1(ak4);osm-6(p811) mutants were significantly longer than either of the single mutants alone (Figure 2B, Table 1). This additive role of nmr-1 and osm-6 is consistent with two independent pathways of quiescence maintenance.

Calcineurin activity in sensory neurons is essential for swimming in liquid

tax-6 and cnb-1, the C. elegans homologues of subunits of calcineurin phosphatase subunits A and B respectively, are expressed in many tissues, including sensory neurons, interneurons and muscles [25,26]. C. elegans calcineurin has been implicated in diverse behaviors including locomotion, thermosensation and chemosensation [25,26]. Calcineurin negatively affects several...
sensory behaviors. For example, loss of tax-6 results in hyper-activation of sensory neurons in an olfactory adaptation paradigm [25].

We have previously observed that in a long-term swimming assay [9], the average fraction of wild type animals in quiescence during the first phase (first 90 minutes) of swimming was low 0.02 +/- 0.004. In the subsequent 90 minutes the average fraction of worms in quiescence increased to 0.23 +/- 0.01. Loss-of-function mutants of either of calcineurin subunit cnb-1(jh300) or tax-6(db60) [25,26] exhibited a striking temporally changing phenotype in which a spike in fraction of animals in quiescence during the first 90 minutes (0.75 ± 0.2 for cnb-1(jh103)) were followed by a decrease in fraction of animals in quiescence (0.62 ± 0.05 for cnb-1(jh103), p < 0.0001, Mann-Whitney U test) during the subsequent period of the assay. In other words a higher fraction of cnb-1(jh103) and tax-6(db60) animals went into quiescence in the first 90 minutes of the assay and remained quiescent during the entire assay duration compared to wild type animals. Thus calcineurin activity is required for preventing animals from going into quiescence. Moreover, the increased quiescence duration in calcineurin loss-of-function mutants is indicative of a role of calcineurin in proper termination of quiescence. Interestingly, the decrease in the fraction of worms in quiescence exhibited by cnb-1(jh103) and tax-6(db60) animals after approximately 90 minutes of the assay (Additional File 2) suggested distinct mechanisms underlying quiescence in the first and second phase of swimming.

In order to determine whether calcineurin activity was required in the nervous system or in other tissues to maintain swimming, we expressed a wild type tax-6 transgene under tissue specific promoters in tax-6(db60) mutants. When expressed under the myo-3 promoter thereby restricting the expression of tax-6 to the body-wall muscles, the increased quiescence phenotype of tax-6(db60) mutants was not rescued (Figure 3A). However, when tax-6 cDNA was expressed in sensory neurons alone from a 1.1 kb fragment of the tax-6 promoter or both sensory and interneurons from an unc-14 promoter [27], the increased quiescence phenotype of tax-6(db60) was rescued, suggesting tax-6 activity in the sensory neurons is critical for maintenance of swimming and proper quiescence duration. Sensory neuronal activity of tax-6 was sufficient for generation of wild type quiescence bouts (Figure 3A, data not shown). It is possible that, in a tax-6 mutant, unopposed phosphorylation by a kinase promotes quiescence. Taken together, these data suggest that activity of calcineurin in the sensory neurons is necessary and sufficient for maintaining a wild type swimming-quiescence pattern in liquid. The altered behavioral patterns in the first and second phases of swimming in calcineurin mutants are likely due to defects in the sensory neurons.

**PKG and calcineurin genetically interact to regulate quiescence**

To investigate whether the quiescence induced due to loss of calcineurin activity was dependent on egl-4, we assayed cnb-1(jh103); egl-4(ks60) double mutants. A significantly lower fraction of worms (0.05 ± 0.04) went into and remained in quiescence compared to the cnb-1 mutant worms (0.62 ± 0.05) throughout the assay suggesting that egl-4 acts genetically downstream of or parallel to cnb-1 in promoting the quiescent state (Figure 3B). Since calcineurin loss-of-function resulted in increased quiescent bout duration, which was dependent on PKG function, our data is consistent with a mechanism where events mediated by unopposed phosphorylation by egl-4 in the cnb-1 loss-of-function mutant increases quiescent bout durations.

**Discussion**

We investigated the genetic basis of a striking behavioral pattern displayed by swimming nematodes in liquid in which they undergo alternate bouts of swimming and quiescence. We found that like other calcineurin-modulated behaviors in C. elegans, activity of calcineurin in the sensory neurons is critical to maintain swimming in C. elegans in liquid. In analogy with the neuronal properties that change in a calcineurin mutant, namely defect in gain control [25], it is likely that increased quiescence in the tax-6/cnb-1 loss-of-function mutants is due to hyperactivation of one or more sensory neurons. In addition, calcineurin mutants displayed a temporally changing phenotype in which there was a significant drop in the fraction of worms in quiescence after 90 minutes. This recovery after 90 minutes from start of assay was more pronounced in case of tax-6(db60) mutants than cnb-1(jh300) mutants, suggesting other phosphatases or autoregulatory mechanisms may play a role in shaping the temporal pattern of swimming-quiescence pattern in liquid. It is possible that in tax-6 loss-of-function mutants unopposed phosphorylation of a yet unidentified substrate induces prolonged quiescence. The phosphorylation possibly involves egl-4 activity as the increased quiescence in calcineurin loss of function mutants was suppressed by loss of function mutations in egl-4. During the first 90 minutes of the assay, PKG and calcineurin play opposing roles in maintenance of swimming. In the second phase (the subsequent 90 minutes) of the assay, however, the fraction of worms in quiescence in egl-4; cnb-1 double mutants was lower than the egl-4 loss of function mutants, suggesting that additional yet unidentified mechanisms may play a role in shaping the temporal pattern of quiescence during swimming in liquid.
We found that the protein kinase G homolog egl-4 acts in the sensory neurons to enhance the quiescence duration liquid. Similar to other paradigms of behavioral quiescence, we found that PKG activity in sensory neurons expressing the cGMP-gated calcium channel tax-4 is critical for regulating quiescent bout duration. However, beyond the involvement of egl-4 in tax-4 expressing neurons, the pathways for induction of different types of quiescence seem to be different. For instance, in quiescence induced by high quality food, loss-of-function mutants of tax-2, encoding the alpha subunit of the cGMP gated ion channels, exhibited very little quiescence [13]. However quiescence in liquid without food doesn’t seem to involve participation of these channels as quiescence bouts of double mutants of alpha and beta subunits, tax-2(p691);tax-4(ky89), were not different from wild type (Table 1), ruling out these channels as phosphorylation targets of egl-4 in modulating quiescence duration in liquid without food. This difference may be attributed to a different target of egl-4 in the sensory neurons or action of egl-4 in additional sites beyond the sensory neurons.

Our data also suggests a second site of action of egl-4 other than the sensory neurons. It seems likely that egl-4 may be acting functionally downstream of acetylcholine release at an as yet unidentified site in regulating quiescence bouts. Our hypothesis is supported by the lack of complete rescue of egl-4 loss of function by expression in the sensory neurons, the suppression in egl-4 mutants of increased quiescence resulting from manipulations of the command interneurons as well as lack of suppression of the GABAergic loss of function mutants. These observations support a site of action of EGL-4 other than within sensory neurons. One possibility is that during quiescence, egl-4 inhibits GABA-ergic signaling and therefore signaling through the ACh pathway is increased causing quiescence. According to this model, in an egl-4 loss-of-function mutant GABA-ergic signaling is enhanced, which terminates quiescence prematurely (Figure 4). It is also possible that egl-4 acts solely in the sensory neurons by activating a neurosecretory pathway that inhibits GABAergic activity (Figure 4). This is consistent with two pathways in maintenance of quiescence. However this scenario requires that sensory neurons with global defects in cilia result in excessive activation of the neurosecretory pathway. Further studies are required to distinguish between these two models. The data is also consistent with a possibility that partial loss of function mutations in egl-4 counters “gain-of-function” like situation that may be mimicked by aldicarb or constitutive depolarization of command interneurons. However this is less likely as increased quiescence due to loss of function mutations in nmr-1, osm-3 and osm-6 was suppressed by egl-4 loss of function mutants.

**Conclusions**

In summary, we have identified parts of a genetic network involving calcineurin and PKG that generate a temporal pattern during swimming in *C.elegans*. Though egl-4 has been implicated in all paradigms of behavioral quiescence in *C. elegans*, our results indicate that the target of EGL-4 in regulating quiescence in liquid is distinct from the
other forms of quiescence. Based on our results we propose dual function of egl-4: one in the sensory neurons that is inhibited by calcineurin and another in inhibiting GABAergic neurotransmission (Figure 4). Our study also implicates opposite roles of a kinase and a phosphatase in generating temporal pattern in behavior. It is likely that the activity of kinase(s) increases the probability of the motor circuit to switch to a quiescent state. Such a process is opposed by calcineurin phosphatase activity in the sensory neurons. Studies of temporal patterns generated by intermittent motor activities have explored statistical properties of the behavioral state transitions or neuro-physiological properties of neural circuits underlying them [1,5,28]. However, very little is known about the genetics of the behavioral intermittencies over time. Our results in a genetically tractable organism establish a framework for further dissection of episodic behaviors.

Methods

Strains
Nematodes were reared and maintained at 20°C on E. coli OP50 on NGM agar plates according to published procedures [29]. Strains that were used are as follows: tax-6(db60), tax-6(db60) [myo3::tax-6], tax-6(db60) [tax-6 sensory+inter-::tax-6], tax-6(db60) [tax-6 sensory +inter+::tax-6] were a kind gift from Ikue Mori (see Figure legend 3 for description of the promoters). The egl-4 transgenic rescue strains (egl-4(n479) [odr-3::egl-4], egl-4(n479) [odr-1::egl-4] and egl-4(n479) [tax-4::egl-4]) were generously provided by David Raizen. akIs9 and tax-2(p691);tax-4(ky89) were kindly provided by A.V. Maricq and Cornelia Bargmann respectively. All other strains were either made during this work or obtained from CGC. Double mutants were generated using standard genetic procedures. osm-6(p811) and osm-3(p802) mutants were monitored by the inability of their sensory neurons to uptake the fluorescent dye DiO [24]. cnb-1 (jh103), egl-4 (ks60) and nmr-(ak4) deletions were confirmed by the following primer pairs:

Left oligo cnb-1: TCTTCTTGTGCACTTCGGTG
Right oligo cnb-1: CAACACAGCCGATCAAATG
EGL-4(KS60) FD: GAAACCTCCAATTCTGCCGAAGG
EGL-4(KS60) RV: GAATTTCCAGTCAACCAAATTC- ATAC
Nmr-1 (ak4) left: GGAAGAGTTTGAAAAACGGCG
Nmr-1 (ak4) right: CGTGTTCTTAGCTCACAGTGTCG

Swimming assays
Swimming assays were performed essentially as described previously [9]. Briefly, one day old adults were transferred to a fresh unseeded plate at 25°C and allowed to forage for 2 min, after which they were picked singly into wells of a microtiter plate containing 200 μl M9 buffer at 25°C. The assays were carried out in a room where the temperature was maintained at 25 ± 1°C. Quiescence was defined as before [9]. Briefly less than two body bends per 5 seconds of observation were considered as quiescent irrespective of movement of the nose/head.

Aldicarb and dye-filing assays
Aldicarb assays in liquid were performed as described before [9]. Dye filing assays were done essentially as published elsewhere with 1:2000 diluted solution of DiO (10 mg/ml stock) [24].

Data analysis
Statistical analysis on the data was done using the Graphpad Prism version 5.0a software for Macintosh. For most cases the data was not normally distributed as determined by the by the D’Agostino and Pearson method. Unless otherwise mentioned we used the non parametric Mann Whitney U test for the quiescence bout durations for two genotypes or Kruskal-Wallis test.
followed by Dunn’s multiple comparison for more than two genotypes. We also used the non-parametric Wilcoxon signed rank test to compare the medians of any given genotype against a hypothetical median of 4.50, which was not significantly different from the median for wild type worms.

Additional file 1: Distribution of frequency of quiescent bout durations for unc-25(sa94ts) and unc-25(sa94ts);egl-4(ks60) worms with bin size = 3 minutes.
Click here for file
[http://www.biomedcentral.com/content/supplementary/1471-2156-11-7-S1.PDF]

Additional file 2: Fraction of worms in quiescence versus time for wild type (n = 25), cnd-1(h105) (n = 24), tax-6(db60) (n = 24) worms.
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[http://www.biomedcentral.com/content/supplementary/1471-2156-11-7-S2.PDF]

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RG and SWE conceived the study, designed the experiments and wrote the manuscript. RG performed the experiments. Both authors read and approved the final manuscript.

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