Microtubule-based localization of a synaptic calcium-signaling complex is required for left-right neuronal asymmetry in *C. elegans*

Chieh Chang¹, Yi-Wen Hsieh¹, Bluma J. Lesch², Cornelia I. Bargmann² and Chiou-Fen Chuang¹,*

### SUMMARY

The axons of *C. elegans* left and right AWC olfactory neurons communicate at synapses through a calcium-signaling complex to regulate stochastic asymmetric cell identities called AWC\textsuperscript{ON} and AWC\textsuperscript{OFF}. However, it is not known how the calcium-signaling complex, which consists of UNC-43/CaMKII, TIR-1/SARM adaptor protein and NSY-1/ASK1 MAPKKK, is localized to postsynaptic sites in the AWC axons for this lateral interaction. Here, we show that microtubule-based localization of the TIR-1 signaling complex to the synapses regulates AWC asymmetry. Similar to unc-43, tir-1 and nsy-1 loss-of-function mutants, specific disruption of microtubules in AWC by nocodazole generates two AWC\textsuperscript{ON} neurons. Reduced localization of UNC-43, TIR-1 and NSY-1 proteins in the AWC axons strongly correlates with the 2AWCON phenotype in nocodazole-treated animals. We identified kinesin motor unc-104(k114) mutants for enhancement of the 2AWCON phenotype of a hypomorphic tir-1 mutant. Mutations in unc-104, like microtubule depolymerization, lead to a reduced level of UNC-43, TIR-1 and NSY-1 proteins in the AWC axons. In addition, dynamic transport of TIR-1 in the AWC axons is dependent on unc-104, the primary motor required for the transport of presynaptic vesicles. Furthermore, unc-104 acts non-cell autonomously in the AWC\textsuperscript{ON} neuron to regulate the AWC\textsuperscript{OFF} identity. Together, these results suggest a model in which UNC-104 may transport some unknown presynaptic factor(s) in the future UNC\textsuperscript{ON} cell that non-cell autonomously control the trafficking of the TIR-1 signaling complex to postsynaptic regions of the AWC axons to regulate the AWC\textsuperscript{OFF} identity.

### KEY WORDS: *C. elegans*, Microtubules, Olfactory development, Left-right asymmetry, Calcium signaling

### INTRODUCTION

The left and right sides of the central nervous system display anatomical, molecular and functional asymmetries throughout the animal kingdom. Functional lateralization in the brain is theorized to increase cognitive performance and social behaviors, as reduced and reverse anatomical brain asymmetry has been linked to a variety of neurodevelopmental disorders (Taylor et al., 2010). However, the mechanisms underlying lateralization of the developing nervous system remain poorly understood.

Like the vertebrate central nervous system, the *C. elegans* nervous system displays molecular and functional asymmetries (Hubert, 2006; Hubert et al., 2002; Sagasti et al., 2001; Taylor et al., 2010). In both ASE taste neurons and AWC olfactory neurons, the left and right neurons of a pair have different patterns of gene expression and respond to different sets of chemicals (Pierce-Shimomura et al., 2001; Troemel et al., 1999; Wes and Bargmann, 2001; Yu et al., 1997). However, the mechanisms that specify ASE and AWC asymmetries are completely distinct. The ASE neurons develop a stereotyped asymmetry early in embryogenesis, whereas the AWC neurons develop stochastic asymmetry late in embryogenesis (Chuang and Bargmann, 2005; Poole and Hubert, 2006). None of the identified ASE asymmetry genes affects AWC asymmetry and, conversely, AWC asymmetry genes do not affect ASE asymmetry (Chang et al., 2003; Koga and Ohshima, 2004; Lanjuin et al., 2003; Lesch et al., 2009).

In wild-type animals, the reporter gene *str-2p::GFP* is expressed in only one of the two AWC neurons, and never in both (Fig. 1A). The two AWC neurons are described as AWC\textsuperscript{ON}, which expresses *str-2p::GFP*, and AWC\textsuperscript{OFF}, which does not. The expression of *str-2p::GFP* in AWC is random: 50% of the animals in a population express *str-2* in the left AWC neuron, whereas the other 50% express *str-2* in the right AWC neuron. Genetic studies and cell killing experiments suggest that AWC\textsuperscript{OFF} is the default state and asymmetric expression of *str-2* requires an interaction between the two AWC neurons (Troemel et al., 1999).

Genetic screens identified several genes that regulate AWC asymmetry. Loss-of-function or reduction-of-function mutations in a calcium-regulated signaling pathway, including a voltage-gated calcium channel (UNC-2, EGL-19, UNC-36), the calcium/calmodulin-dependent protein kinase II (CaMKII) UNC-43, the Toll-interleukin 1 repeat protein TIR-1/SARM, the mitogen-activated protein kinase kinase kinase (MAPKKK) NSY-1/ASK1 and the MAPKK SEK-1 kinase cascade, lead to animals with two AWC\textsuperscript{ON} (Fig. 1C; 2AWCON phenotype) and no AWC\textsuperscript{OFF} cells. These results suggest that these genes normally promote the default AWC\textsuperscript{OFF} identity (Bauer Huang et al., 2007; Chuang and Bargmann, 2005; Sagasti et al., 2001; Tanaka-Hino et al., 2002; Troemel et al., 1999). An innexin gap junction protein, encoded by *str-5*, and a claudin-like protein, encoded by *nsy-4*, function in parallel to inhibit the downstream calcium-regulated signaling pathway in the neuron that acquires the AWC\textsuperscript{ON} identity (Chuang et al., 2007; VanHoven et al., 2006). When the activity of *nsy-4* or *nsy-5* is reduced or lost, mutant animals fail to induce the AWC\textsuperscript{ON} cell, leading to the formation of...
two AWCoff cells (2AWCoff phenotype). NSY-5 forms transient gap junctions between the cell bodies of the AWCs and their neighboring neurons in embryos. Genetic analysis suggests that communication between the two AWCs and other neurons in the NSY-5 gap junction network is required for the induction of AWC asymmetry (Chuang et al., 2007). Once AWC asymmetry is specified in late embryogenesis, both the AWCoff and AWCon identities are maintained throughout the life of the worm by cGMP signaling, dauer pheromone signaling and transcriptional repressors. Animals with mutations in maintenance pathways have wild-type AWC asymmetry in the first larval stage, but display mutant AWC phenotypes in the adult stage (Lesch and Bargmann, 2010; Lesch et al., 2009; Troemel et al., 1999).

TIR-1/SARM functions as an adaptor protein to couple upstream UNC-43/CaMKII and downstream NSY-1/ASK1 MAPKKK signaling at postsynaptic regions in the AWC axons for the regulation of AWCoff identity (Chuang and Bargmann, 2005). Proper localization of these calcium signaling proteins at AWC synapses is important for the regulation of AWC asymmetry, but it is not understood how the AWC asymmetry signaling complex is targeted to the synapses. Here, we show that microtubules and microtubule-dependent kinesin motor UNC-104/KIF1A control the synaptic localization of the TIR-1 signaling complex to regulate left-right AWC asymmetry.

MATERIALS AND METHODS

Strains

Wild-type strains were C. elegans variety Bristol, strain N2. Strains were maintained by standard methods (Brenner, 1974). Mutations and integrated transgenes used in this study included: kysl40 [str-2p::GFP; lin-15(+j) I] (Troemel et al., 1999), vysl2 [odr-3p::tir-1::GFP, ofp-1::DsRed] I, nys-5(ky634) I (Chuang et al., 2007), unc-104(rh142) II, unc-104(rh43) II, unc-104(e1265) II, kysl323 [str-2p::GFP; ofm-1::GFP] II, ben-1(e1880) III, tir-1(tm3036) III, tir-1(ky388ts) III (Chuang and Bargmann, 2005), tir-1(ky649g) III, unc-43(n949g) IV, vysl44 [odr-1::DsRed] V, vysl51 [str-2p::2xnsTagRFP, ofm-1::DsRed] V, and vysl48 [odr-1::DsRed; odr-3p::unc-43; GFP; str-2p::nlsCFP, ofm-1::DsRed].

Transgenes maintained as extrachromosomal arrays included: vyEx1123, 1124 [odr-3p::ben-1::SL2::TagRFP, ofm-1::DsRed], vyEx668, 677, 678 [odr-3p::tir-1::GFP; str-2p::nlsCFP, ofm-1::DsRed], vynl669, 670, 673 [odr-3p::tir-1::GFP; str-2p::nlsCFP, ofm-1::DsRed], kyEx588 [odr-3p::nsy-1::GFP], myo-3prom-unc-104 (Chuang and Bargmann, 2005), vyEx1103, 1104 [odr-3p::nsy-1::GFP, ofm-1::DsRed], vyEx1020, 1029, 1037 [odr-3p::unc-104; odr-1::DsRed; ofm-1::DsRed], vyEx944, 945, 946 [unc-104::unc-104; odr-1::DsRed; ofm-1::DsRed], vyEx925 [unc-104::GFP; odr-1::DsRed], vyEx779, 780, 787 [odr-3p::tir-1::DsRed; ofm-1::unc-43; GFP; str-2p::2xnlsTagRFP, ofm-1::DsRed], vyEx661, 615 [odr-3p::GFP; elp-1::GFP; and ofm-1::GFP; str-2p::3xnsCFP, ofm-1::DsRed].

Plasmid construction and germline transformation

A 4.2 kb PCR fragment of unc-104 promoter was subcloned into pPD95.77 to make unc-104::GFP. ben-1 cDNA (1332 bp) and unc-104 cDNA (4752 bp) were obtained with RT-PCR of total mRNA from mixed stage worms and subcloned to make odr-3p::ben-1::SL2::TagRFP, odr-3p::unc-104, unc-104::unc-104 and odr-3p::unc-104::GFP, odr-3p::tir-1::GFP. For extension of AWC asymmetry, a convenient tool with which to analyze the function of microtubules in biological processes. To test for a potential role of microtubules in the establishment of AWC asymmetry, wild-type animals containing the str-2p::GFP transgene were treated with nocodazole for 2 hours at different embryonic stages and in the first larval (L1) stage, and adults were scored for str-2p::GFP expression in AWC. The AWC neurons are born at about 1 hour after egg laying and their axons extend at about 3.5 hours after egg laying. tir-1 function must be present by late embryogenesis (~8 hours after egg laying) to generate AWC asymmetry, suggesting that the crucial period for the establishment of AWC asymmetry is before the late embryo stage (Chuang and Bargmann, 2005). Nocodazole treatment of embryos before egg laying or at 0.6 hours after egg laying led to the expression of str-2p::GFP in both AWC neurons, a 2AWCOn phenotype (Fig. 1B; see Fig. S1 in the supplementary material). Nocodazole-treated animals with two AWCOn neurons also
displayed slow growth, dumpy morphology, and/or uncoordinated locomotion phenotypes. However, nocodazole treatment of embryos at 6-8 hours after egg laying or during the first larval stage (~13 hours after egg laying) did not cause a 2AWC\textsuperscript{ON} phenotype (see Fig. S1 in the supplementary material). These results suggest that nocodazole needs to be present before and/or during the critical period for \textit{tir}-1 function to affect AWC asymmetry. In subsequent studies, mixed stages of embryos before and after egg laying were used for nocodazole or benomyl (another inhibitor of microtubule polymerization) treatment and only slower-growing adults were scored for AWC phenotypes. We also observed a 2AWC\textsuperscript{ON} phenotype at a lower frequency than nocodazole treatment when embryos were treated with benomyl (Table 1, rows b and c). Benomyl-treated animals sometimes displayed a weak 2AWC\textsuperscript{OFF} phenotype. Some of the 2AWC\textsuperscript{ON} animals resulting from benomyl or nocodazole treatment had truncated and disoriented AWC axons (Fig. 1B). Mutants such as \textit{unc}-76 loss-of-function alleles disrupt axon outgrowth of AWC neurons, but generate a 2AWC\textsuperscript{OFF} phenotype (Troemel et al., 1999), the opposite phenotype from the 2AWC\textsuperscript{ON} phenotype of nocodazole- or benomyl-treated animals. These results suggest that nocodazole and benomyl must affect a different aspect of AWC development in addition to any possible effects on axon outgrowth.

Loss-of-function mutations in \textit{ben-1}, which encodes a β-tubulin, confer resistance to benimidazoles such as nocodazole and benomyl (Driscoll et al., 1989). To determine whether the 2AWC\textsuperscript{ON} phenotype is due to disruption of microtubules, we examined the effect of nocodazole on \textit{ben-1} mutants. \textit{ben-1(e1880)} mutants showed wild-type expression of \textit{str-2}, and treatment of \textit{ben-1(e1880)} mutant embryos with nocodazole did not cause a 2AWC\textsuperscript{ON} phenotype (Table 1, rows d and e). However, expression of wild-type \textit{ben-1} gene primarily in AWC using an \textit{odr-3} promoter (Roayaie et al., 1998) restored the susceptibility of \textit{ben-1(e1880)} mutants to nocodazole in the generation of 2AWC\textsuperscript{ON} phenotype (Table 1, rows f and g). These results suggest that the 2AWC\textsuperscript{ON} phenotype in nocodazole-treated animals is caused by specific disruption of microtubules in AWC.

**Nocodazole treatment may affect the function of the TIR-1 signaling complex in AWC asymmetry**

The molecular signaling pathway that regulates AWC asymmetry involves a calcium-regulated kinase pathway including \textit{unc-43} (CaMKII), \textit{tir-1} (SARM/adaptor protein) and \textit{nsy-1} (ASK1/MAPKKK) (Chuang and Bargmann, 2005; Sagasti et al., 2001; Troemel et al., 1999) (Fig. 1D). Loss-of-function mutants of \textit{unc-43}, \textit{tir-1} or \textit{nsy-1} showed a 2AWC\textsuperscript{ON} phenotype resembling that caused by nocodazole treatment (Fig. 1B,C). This similarity suggests that the nocadazole-induced 2AWC\textsuperscript{ON} phenotype could result from defects in the calcium-regulated CaMKII-TIR-1-NSY-1 signaling pathway.

To determine the potential role of nocodazole in the AWC signaling pathway, we treated different mutants affecting AWC asymmetry with nocodazole. As nocodazole treatment generated two AWC\textsuperscript{OFF} cells, it was administered to the mutants with the opposite phenotype (two AWC\textsuperscript{OFF} cells) to determine the relationship of the nocadazole target with other genes in the AWC pathway. The interaction between nocodazole and mutations was analyzed using the same principles as genetic analysis of double mutants. Loss-of-function (lf) mutations in the \textit{nsy-5}\textsuperscript{m} genes or gain-of-function (gf) mutations in \textit{unc-43}, \textit{tir-1} and \textit{nsy-1}, which cause two AWC\textsuperscript{OFF} cells, (Chuang et al., 2007; Chuang and Bargmann, 2005; Sagasti et al., 2001; Troemel et al., 1999) (Fig. 1D) were treated with nocodazole. Nocodazole treatment of \textit{nsy-5}(lf) or \textit{unc-43}(gf) mutants strongly suppressed their 2AWC\textsuperscript{OFF} phenotype and generated a 2AWC\textsuperscript{ON} phenotype (Table 1, rows h, i, k, l). These results are consistent with the effect of nocodazole on signaling predominantly downstream or parallel to \textit{nsy-5} and \textit{unc-43}. By contrast, the 2AWC\textsuperscript{ON} phenotype resulting from nocodazole treatment was greatly suppressed by \textit{nsy-1}(gf) mutants (Table 1, rows t, u). This result suggests that nocodazole acts on a target mainly upstream of or parallel to \textit{nsy-1}.

These results are consistent with a model in which nocodazole affects the AWC signaling step downstream of/parallel to \textit{unc-43} CaMKII and upstream of/parallel to \textit{nsy-1} MAPKKK. Previous genetic studies suggest that \textit{tir-1} also acts downstream of \textit{unc-43}
CaMKII and upstream of nys-1 MAPKKK to regulate AWC asymmetry (Fig. 1D). Thus, we examined the interaction between tir-1 and nocodazole. The tir-1(ky648) allele, in which a C to T mutation results in an alanine to valine substitution of the 538th amino acid of the TIR-1a isoform, was isolated from a genetic screen for mutants with a 2AWCOFF phenotype (VanHoven et al., 2006). An amino acid of the TIR-1a isoform, was isolated from a genetic screen for mutants with a 2AWCOFF phenotype (VanHoven et al.).

The strong 2AWCOFF phenotype of tir-1(ky648gf) was the opposite of the 2AWCON phenotype (Chuang and Bargmann, 2005). As the TIR-1 signaling complex is regulated by microtubules, we investigated whether the subcellular localization of TIR-1, along with its signaling complex components UNC-43 and NSY-1, is regulated by microtubules.

Microtubules are required for the localization of TIR-1, UNC-43 and NSY-1 in the AWC axons to regulate the AWCOFF identity

The axons of the two AWC neurons form chemical synapses on each other in the nerve ring, where axons from the left and right sides meet (White et al., 1986). Our previous results showed that TIR-1/SARM adaptor, UNC-43/CaMKII and NSY-1 ASK1/ MAPKKK are enriched in postsynaptic regions of the AWC axons (Chuang and Bargmann, 2005). As the TIR-1 signaling complex is a possible target of nocodazole in the AWC signaling pathway, we investigated whether the subcellular localization of TIR-1, along with its signaling complex components UNC-43 and NSY-1, is regulated by microtubules.

unc-43, tir-1 and nys-1 act to execute the AWCOFF decision in late embryogenesis (Chuang and Bargmann, 2005; Sagasti et al., 2001). However, in tir-1(ky648gf) mutants, about 37% of animals had one AWCON neuron in the L1 stage (Table 1, row m), whereas the fraction of animals with one AWCON neuron significantly decreased to 9% in adults (Table 1, row n). These results suggest that tir-1(ky648gf) mutants are also defective in the maintenance of the AWCOFF identity. Similar to tir-1(ky648gf), unc-43(n498gf) and nys-1(gf) also displayed a defect in the maintenance of the AWCOFF state, with more animals exhibiting one AWC neuron in the L1 stage than as adults (Table 1, rows j,k,s,t). These results reveal a role for tir-1, unc-43 and nys-1 in the suppression of the AWCOFF identity and/or promotion of the AWCOFF identity in the L1 stage. Because late embryos constantly move and are poorly permeable to anesthetics such as sodium azide and levamisole, we chose to use the L1 stage for live imaging of fluorescently tagged proteins.

To test directly the effect of nocodazole on the localization of TIR-1, UNC-43, NSY-1 in the AWC axons and the function of these signaling proteins in AWC asymmetry, odr-3p::tir-1::GFP was co-expressed with str-2p::nls-TagRFP, a red nucleus-localized AWCOFF marker. Similarly, odr-3p::unc-43::GFP or odr-3p::nys-1::GFP was co-expressed with str-2p::nls-TagRFP, a red nucleus-localized AWCOFF marker. These strains allowed simultaneous visualization of protein subcellular localization and AWC asymmetry in the same animal. The fluorescence intensity of TIR-1::DsRed, UNC-43::GFP and NSY-1::GFP in the AWC axons was examined and quantified in nocodazole-treated animals that exhibited 1AWCOFF/1AWCON (hereafter, wild type) and 2AWCON phenotypes.

Although both wild-type and 2AWCON animals had TIR-1::DsRed puncta in their AWC axons, the numbers of puncta were significantly different (P=0.0001). Wild-type animals had an average of 5.4 TIR-1::DsRed puncta per animal (n=32), whereas 2AWCON animals had 1.6 puncta per animal (n=21) (data not shown). The average fluorescence intensity of TIR-1::DsRed in the AWC axons was also significantly reduced in 2AWCON animals.

Table 1. Effect of nocodazole and benomyl treatment on str-2 expression in AWC

<table>
<thead>
<tr>
<th>Genetic background</th>
<th>Row</th>
<th>Treatment</th>
<th>2AWCON (%)</th>
<th>1AWCON/1AWCOFF (%)</th>
<th>2AWCON (%)</th>
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<tr>
<td>Wild type</td>
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<tr>
<td></td>
<td>b</td>
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<td>10</td>
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<td></td>
<td>c</td>
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<td>36</td>
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<tr>
<td>ben-1(e1880)</td>
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<td>–</td>
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</tr>
<tr>
<td></td>
<td>e</td>
<td>Nocodazole</td>
<td>3</td>
<td>97</td>
<td>0</td>
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<tr>
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<td>–</td>
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<td>g</td>
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<td>7</td>
<td>62</td>
<td>31****</td>
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<tr>
<td>nys-5(ky634lf)</td>
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<td>33</td>
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<tr>
<td>unc-43(n498gf), L1</td>
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<tr>
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<td>l</td>
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<tr>
<td>tir-1(ky648gf), L1</td>
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<td>odr-3p::nys-1(gf)</td>
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<td>Nocodazole</td>
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Animals were scored as adults, unless otherwise indicated.

+ n, no treatment; **P<0.003; ****P<0.0001; ns, not significant. Z-test for two proportions.
In addition, expression of TIR-1::DsRed in the AWC cell body was stronger in 2AWCON animals than in wild type (Fig. 2A,B). These results reveal a tight link between microtubule-dependent TIR-1 localization in the AWC axons and AWCOFF identity determination.

UNC-43::GFP and NSY-1::GFP were not as tightly localized in the AWC axons as TIR-1, thus we directly quantified their fluorescence intensity instead of measuring the puncta number. In wild type, both UNC-43::GFP and NSY-1::GFP were localized in a punctate pattern in the AWC axons with some diffuse expression in the AWC cell body (Fig. 2D,G). In 2AWCON animals, expression of both UNC-43::GFP and NSY-1::GFP was significantly reduced in the AWC axons (Fig. 2E,F,H,I). The expression level of UNC-43::GFP in the AWC cell body is similar in wild-type and 2AWCON animals. However, expression of NSY-1::GFP in the AWC cell body is faint in 2AWCON animals, suggesting that microtubules may be important for the stability of NSY-1 protein. These results suggest that the presence of wild-type levels of UNC-43 and NSY-1 in the AWC axons is dependent on microtubule function and is required for AWC asymmetry.

As tir-1 promotes the synaptic localization of NSY-1 protein in the AWC axons (Chuang and Bargmann, 2005), reduced amounts of NSY-1 in the AWC axons of nocodazole-treated animals may be due to decreased TIR-1 localization in the axons. To determine whether the effect of nocodazole on the localization of UNC-43 in the AWC axons is dependent on reduced localization of TIR-1, the odr-3p:unc-43::GFP transgene was crossed to tir-1(tm3036lf) mutants. tir-1(tm3036lf), probably a null allele, contains a deletion removing the C-terminal TIR activation domain and causing 100% of animals to become 2AWCON (Table 1, row p). The localization pattern and level of UNC-43::GFP in the AWC axons is similar in wild type and tir-1(tm3036lf) mutants (see Fig. S2 in the supplementary material), suggesting that UNC-43 localization in the AWC axons is regulated by microtubules but is independent of TIR-1 localization.

Together, these results are consistent with a model that microtubules are required for proper localization of TIR-1, UNC-43 and NSY-1 in the AWC axons to regulate the AWCOFF cell identity.

The kinesin motor unc-104/kif1a genetically interacts with tir-1 in the regulation of the AWCOFF identity

Many proteins are transported by dynein/dynactin and kinesin motor proteins along microtubules to different locations within the cells (Muresan, 2000; Susalka et al., 2000). There are at least 25 dynein/dynactin-related and 23 kinesin-related genes predicted in the C. elegans genome (C. elegans Sequencing Consortium, 1998). To overcome potential functional redundancy of microtubule motor proteins in AWC asymmetry, we performed a sensitized RNA interference (RNAi) genetic screen using a tir-1(ky388) temperature-sensitive (ts) mutant, in which the 2AWCON phenotype was incompletely penetrant (34% 2AWCON) at 15°C (Chuang and Bargmann, 2005) (Table 2). We found that RNAi of the kinesin unc-104 significantly enhanced the 2AWCON frequency from 34% to 58% in tir-1(ky388ts) at 15°C (Table 2; P<0.001). To confirm the effect of unc-104(RNAi) on AWC asymmetry, we made tir-1(ky388ts); unc-104(e1265) double mutants and found a strong enhancement of the 2AWCON frequency from 34% to 91% (P<0.0001). These results suggest that unc-104 is required for the regulation of the AWCOFF identity. unc-104(RNAi) or unc-104(e1265) alone did not show a 2AWCON phenotype. Single
unc-104(kif1a) acts non-cell autonomously in the AWC neuron to regulate the AWC\textsuperscript{OFF} identity

Expression of \( \text{unc-104} \) in AWC from the \( \text{odr-3} \) promoter significantly rescued the enhancement of AWC\textsuperscript{ON} frequency from 91% to 39% in \( \text{tir-1(ky388ts)} \); \( \text{unc-104(e1265)} \) double mutants (Table 2; \( P<0.0001 \)). This result suggests that \( \text{unc-104} \) probably acts in AWC neurons to promote the AWC\textsuperscript{OFF} identity. To further refine the site of \( \text{unc-104} \) action in AWC asymmetry, mosaic animals in which \( \text{unc-104} \) activity is different between the two AWC neurons were used to determine whether \( \text{unc-104} \) acts in the AWC\textsuperscript{ON} cell, the AWC\textsuperscript{OFF} cell, or both. An extrachromosomal transgene containing \( \text{unc-104p}::\text{unc-104} \) and \( \text{o Dr-1p}::\text{DsRed} \) (expressed in AWC and AWB) was introduced into \( \text{unc-104(e1265)} \) mutants. The \( \text{unc-104p}::\text{unc-104} \) transgene rescued the uncoordinated locomotion phenotype of \( \text{unc-104(e1265)} \) mutants (data not shown). Transgenes expressing the \( \text{o Dr-1p}::\text{DsRed} \) marker and \( \text{unc-104p}::\text{unc-104} \) in both AWC neurons caused a mixed weak phenotype of AWC\textsuperscript{ON} and AWC\textsuperscript{OFF} (Fig. 4A). Spontaneous loss of the extrachromosomal array in one of the two AWC neurons resulted in mosaic animals in which one of the two AWC neurons expressed the transgene and therefore had \( \text{unc-104} \) activity; this cell could be identified by expression of the DsRed marker. In the majority of these mosaic animals, the \( \text{unc-104(+)} \) AWC neuron became AWC\textsuperscript{ON} and the \( \text{unc-104(-)} \) AWC neuron became AWC\textsuperscript{OFF} (Fig. 4B). These results are consistent with a significant cell-autonomous requirement for \( \text{unc-104} \) within the AWC\textsuperscript{ON} cell to regulate the AWC\textsuperscript{OFF} identity in a non-cell autonomous manner, which is different from the cell autonomous function of UNC-43, TIR-1 and NSY-1 in the regulation of AWC\textsuperscript{OFF} cell.

unc-104(kif1a) is required for the localization of TIR-1, UNC-43, and NSY-1 in the AWC axons

\( \text{C. elegans} \) UNC-104 and the mouse ortholog KIF1A are neuron-specific kinesin motors that transport presynaptic vesicles along microtubules from the cell body to the nerve terminal (Hall and Hedgecock, 1991; Okada et al., 1995). In \( \text{unc-104(e1265)} \) mutants, most synaptic vesicles remain trapped in the cell body and are nearly absent from the axon (Hall and Hedgecock, 1991; Ou et al., 2005).
2010). UNC-49 GABA receptors form postsynaptic clusters at neuromuscular junctions in a unc-104-dependent manner, suggesting that a factor required for postsynaptic receptor clustering is transported from the cell body to presynaptic sites by UNC-104 (Gally and Bessereau, 2003).

To determine whether the postsynaptic localization of TIR-1, like postsynaptic GABA receptors, is dependent on unc-104, we quantitatively analyzed the localization pattern of TIR-1::GFP in wild-type animals and unc-104 mutants. The intensities of TIR-1::GFP puncta were significantly different between wild-type and unc-104(e1265) mutant animals. These results support the hypothesis that UNC-104 may transport some presynaptic factors that regulate the activity, and that anterograde and retrograde movement of TIR-1 is regulated differently. We also measured the velocities of anterograde and retrograde movements of TIR-1::GFP. In wild-type animals, the average speed of anterograde movements is 1.8 μm/second, similar to the speed of unc-104-dependent movements of synaptic vesicles precursors and dense core vesicles measured from in vivo imaging of C. elegans neurons (Ou et al., 2010; Zahn et al., 2004). The average speed of retrograde...
movements is 1.7 μm/seconds, similar to the speed of the retrograde motor dynein in living Dictyostelium cells (Ma and Chisholm, 2002). The average speed of TIR-1 movement in either direction was not significantly altered in unc-104(e1265) mutants (data not shown). These results suggested that altered localization of TIR-1 is dependent on changes in the average direction, rather than the average speed, of transport.

As the localization of UNC-43 and NSY-1, like TIR-1, in the AWC axons was significantly reduced in nocodazole-induced 2AWC\textsuperscript{ON} animals, we also examined the possibility that the localization of these proteins could be regulated by unc-104. In unc-104(e1265) mutants, UNC-43::GFP and NSY-1::GFP showed significantly reduced levels of localization in the AWC axons (Fig. 5D-I). The like effect of nocodazole on their localization in 2AWC\textsuperscript{ON} animals (Fig. 2D-I), the expression level of UNC-43::GFP in the AWC cell body was similar in wild type and unc-104(e1265) mutants, whereas NSY-1::GFP expression was reduced in the AWC cell body in unc-104(e1265) mutants. These results suggest that unc-104 is required for the localization and/or stabilization of UNC-43 and NSY-1 in the AWC axons.

**DISCUSSION**

One challenge for genetic approaches aimed at studying biological processes is the difficulty of identifying specific functions for broadly acting cell biological pathways. Here, we complement classical genetics with a chemical approach to disrupt polymerized microtubules at a specific time in development. Complete microtubule loss leads to lethality and defects in mitosis, but our experiments took advantage of the fact that late embryogenesis, the time at which asymmetric AWC identities are determined, is a time when little cell division occurs. Using a combination of genetics and drug treatment, we have linked microtubule-dependent localization of calcium-regulated signaling proteins, including UNC-43/CaMKII, TIR-1/SARM adaptor and NSY-1/ASK1 MAPKKK to genetic control of neuronal asymmetry in C. elegans. We also identified the requirement of microtubule-dependent kinesin motor gene unc-104\textsuperscript{ON}/kif1\textsuperscript{a} in this process.

Our results suggest that microtubule-dependent and unc-104\textsuperscript{ON} dependent localization of the TIR-1 signaling complex in the AWC axons may be required for maintenance of AWC asymmetry in early larval stages. In addition, the requirement of nocodazole treatment in embryos for generating AWC phenotypes and the potential role of unc-104 in a negative feedback signal from presynaptic vesicles along the AWC axons of unc-104\textsuperscript{ON}/AWC\textsuperscript{OFF} suggest that microtubules and unc-104 may also function in the early AWC\textsuperscript{ON}/AWC\textsuperscript{OFF} decision step during embryogenesis.

Nocodazole needs to be present before and/or during the crucial period in late embryos to affect AWC asymmetry. Two hours of nocodazole treatment would be sufficient to allow the drug to penetrate the embryos. Although the exposure of embryos to nocodazole is transient, the retention of nocodazole in treated early embryos could continue to inhibit microtubule function and microtubule-dependent assembly of the UNC-43/TIR-1/NSY-1 signaling complex in the AWC axons until the crucial stage of AWC asymmetry in late embryos and the maintenance of asymmetry in early larvae. Alternatively, nocodazole may only affect the process in the embryos, and the effect of the drug is then stabilized through the maintenance pathway.

TIR-1 is dynamically transported in both anterograde and retrograde directions in the AWC axons. Analogous to retrograde neurotrophin signaling (Ginty and Segal, 2002), lateral signaling between AWC neurons may be transmitted in a retrograde direction from the synapses to the cell body to regulate distinct patterns of gene expression. It is possible that TIR-1 may serve as an adaptor to link some yet to be identified motor proteins and the UNC-43/TIR-1/NSY-1 calcium-signaling complex for both anterograde and retrograde transport of signals that regulate asymmetric AWC gene expression. Microtubule depolymerization by nocodazole results in defective AWC asymmetry, which could be due to disruption of both anterograde and retrograde transport of the TIR-1 signaling complex. The identification of microtubule-dependent kinesin motor gene unc-104 in this process will lay the groundwork for future studies of the genetic networks that link microtubules and the regulation of calcium signaling in stochastic left-right neuronal asymmetry.

The localization of UNC-43, TIR-1 and NSY-1 to the AWC axons was significantly reduced in unc-104(e1265) mutants, although the reduced level of these calcium signaling proteins was probably sufficient to regulate AWC asymmetry as unc-104(e1265) single mutants did not have AWC\textsuperscript{OFF} phenotype. unc-104 may function redundantly with other unknown motor proteins or microtubule components to regulate the AWC\textsuperscript{OFF} identity. Functional redundancy of different motor proteins has been shown for assembly of the mitotic spindle and transport of mitochondria (Nag et al., 2008; Nangaku et al., 1994; Roof et al., 1992; Tanaka et al., 1998). In tir-1(ky388ts); unc-104(e1265) double mutants, a strong enhancement of the 2AWC\textsuperscript{ON} phenotype could be due to a synergistic effect of reduced tir-1 activity in tir-1(ky388ts) mutants and decreased localization of the UNC-43/TIR-1/NSY-1 signaling complex in the AWC axons of unc-104(e1265) mutants.

UNC-43, TIR-1 and NSY-1 are enriched at postsynaptic sites of the AWC axons (Chuang and Bargmann, 2005), and their localization in the AWC axons is dependent on unc-104, which has been previously shown to transport presynaptic vesicles along microtubules from the cell body to the nerve terminal (Hall and Hedgecock, 1991; Okada et al., 1995). UNC-104/KIF1\textsuperscript{a} may transport some presynaptic factor(s) in the future AWC\textsuperscript{ON} cell that non-cell autonomously regulate the trafficking of UNC-43/TIR-
1/NSY-1 signaling complexes to postsynaptic regions of the AWC axons to regulate the AWC<sup>OFF</sup> identity. <i>cvc-1</i> (cycclin), <i>pct-1</i> (cycclin-dependent Pctaire kinase) and <i>ckt-5</i> (cycclin-dependent kinase) are part of the two cycclin-dependent pathways that act in parallel and partially redundantly to direct polarized trafficking of presynaptic components. <i>unc-104</i> might be involved in the same biological processes as both cycclin-dependent pathways (Ou et al., 2010). Our results showed that <i>cvc-1(RNAi)</i> and <i>pct-1(RNAi)</i>, but not <i>ckt-5(RNAi)</i>, significantly enhanced the 2AWC<sup>COn</sup> phenotype of <i>psc-1(ky388ts)</i> mutants. However, <i>cvc-1(RNAi); unc-104(e1265f)</i> or <i>pct-1(RNAi); unc-104(e1265f)</i> did not show AWC phenotypes (data not shown). These results suggest that <i>cvc-1</i>, <i>pct-1</i> and <i>unc-104</i> may act in the same pathway to transport presynaptic factors that regulate the postsynaptic localization of the TIR-1 signaling complex in the AWC axons. A loss-of-function mutation in <i>unc-13</i>, which encodes a novel conserved protein that regulates neurotransmitter release at the synapse (Maruyama and Brenner, 1991), did not enhance the 2AWC<sup>COn</sup> phenotype of <i>psc-1(ky388ts)</i> mutants (data not shown). It is possible that <i>unc-13(e450f)</i> mutants may still have residual neurotransmission activity. Alternatively, the presynaptic factors regulating the trafficking of UWC-43/TIR-1/NSY-1 in the AWC axons could be non-vesicular proteins transported by the kinesin UNC-104.

NSY-5 gap junctions and NSY-4 claudins are two parallel signaling systems that have opposite intrinsic side biases to induce the AWC<sup>COn</sup> state. It has been proposed that stochastic AWC asymmetry may be driven by relative strengths of the <i>nsy-5</i> signal and the <i>nsy-4</i> signal when the two AWC neurons communicate through the NSY-5 gap junction neuronal network. The AWC cell with a stronger signal then generates a negative-feedback signal to suppress the AWC<sup>COn</sup> state in the contralateral AWC (Chuang et al., 2007). However, the molecular mechanisms of this negative-feedback regulation are not understood. Our genetic results suggest that <i>unc-104</i> may function cell-autonomously in the pre-AWC<sup>COn</sup> cell through an unknown mechanism to regulate the AWC<sup>COn</sup> identity. In addition, <i>unc-104</i> may non-cell autonomously control the localization of UNC-43, TIR-1 and NSY-1 in the pre-AWC<sup>OFF</sup> cell to regulate the AWC<sup>OFF</sup> identity, distinct from the cell-autonomous activity of the TIR-1 signaling complex required in the AWC<sup>OFF</sup> cell. We propose that <i>unc-104</i> may be involved in a negative-feedback signal, sent from pre-AWC<sup>COn</sup> to pre-AWC<sup>OFF</sup>, to ensure enactment of a precise AWC<sup>COn</sup>/AWC<sup>OFF</sup> decision. Identification and characterization of the presynaptic factors transported by UNC-104 would shed light on such negative-feedback mechanisms.

Microtubules have been implicated in different biological processes of the developing nervous system, including neurogenesis, neuronal migration, axon guidance and synapse formation (Singh and Tsai, 2010). Mutations in human tubulin genes or in genes that regulate microtubule function give rise to brain disorders (Demer et al., 2005; des Portes et al., 1998; Gleeson et al., 1998; Jaglin et al., 2009; Keays et al., 2007; Poirier et al., 2007; Tischfield et al., 2010). In addition, dysfunction of microtubule-motor-dependent axonal transport has been linked to several neurodegenerative diseases (De Vos et al., 2008). This study reveals specific roles of microtubules and microtubule-dependent kinesin motor <i>unc-104(k187a)</i> in regulating calcium signaling for asymmetric neuronal differentiation.

TIR-1 is an ortholog of the mammalian TIR domain adaptor protein SARM (Mink et al., 2001), which is also called Mdr88-5 (Kim et al., 2007). SARM is expressed primarily in neurons, and associates with microtubules and mitochondria, as well as JNK3 and MAPK10 isoform 3. Hippocampal neurons from SARM-deficient mice are protected from death during deprivation of oxygen and glucose, suggesting an important role for SARM in regulating neuronal survival (Kim et al., 2007). TIR-1 and NSY-1 are implicated in regulation of anoxic death (Hayakawa et al., 2011), suggesting that this pathway may also have a conserved function in neuronal survival in response to stress. In addition, SARM regulates microtubule stability and neuronal morphology (Chen et al., 2011). Our study of microtubules and motor proteins in trafficking of the TIR-1 calcium-signaling complex in the context of neuronal differentiation and function may provide insights into the regulation of SARM function and calcium signaling in other aspects of neural development.

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