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Troponin I controls ovulatory contraction of non-striated actomyosin networks in the C. elegans somatic gonad

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Summary
The myoepithelial sheath of the Caenorhabditis elegans somatic gonad has non-striated actomyosin networks that provide contractile forces during ovulation, a process in which a mature oocyte is expelled from the ovary. Troponin T and troponin C are known regulators of contraction of the myoepithelial sheath. These are two of the three components of the troponin complex that is generally considered as a striated-muscle-specific regulator of actomyosin contraction. Here, we report identification of troponin I as the third component of the troponin complex that regulates ovulatory contraction of the myoepithelial sheath. C. elegans has four genes encoding troponin-I isoforms. We found that tni-1 and unc-27 (also known as tni-2) encode two major troponin-I isoforms in the myoepithelial sheath. Combination of RNA interference and mutation of tni-1 and unc-27 resulted in loss of the troponin-I protein in the gonad and caused sterility due to defective contraction of the myoepithelial sheath. Troponin-I-depleted gonads were hypercontracted, which is consistent with the function of troponin I as an inhibitor of actomyosin contraction. Troponin I was associated with non-striated actin networks in a tropomyosin-dependent manner. Our results demonstrate that troponin I regulates contraction of non-striated actomyosin networks and is an essential cytoskeletal component of the C. elegans reproductive system.

Key words: Troponin, Contraction, Ovulation, Myoepithelial cells, Actin, Myosin

Introduction
A wide variety of muscle systems occur in different tissues and function as force-generating organs. Muscle is classified into striated and non-striated (or smooth) muscles on the basis of the organization of the actin and myosin filaments. Striated muscle has sarcomeric organization of actin and myosin filaments, and repeats rapid cycles of contraction and relaxation. Non-striated muscle has a lattice-like network of actin and myosin filaments, and generally contracts and relaxes slowly. The difference in their contractile properties can partly be explained by different regulatory systems for actin-myosin interaction. In striated muscle, the actin-myosin interaction is inhibited at low Ca²⁺ concentrations by troponin that is linked to actin filaments, but the inhibition is released by binding of Ca²⁺ to troponin (Squire, 1997). However, in smooth muscle, contraction is activated by phosphorylation of myosin regulatory light chain (Gerthoffer, 2005; Somlyo and Somlyo, 1994). Although troponin is predominantly expressed in striated muscle, ascidian smooth muscle expresses troponin (Endo and Obinata, 1981) and a recent study has demonstrated that troponin is also expressed in mammalian vascular smooth muscle (Moran et al., 2008). Non-striated muscle and other non-striated-muscle-like cells, such as myoepithelial cells, myofibroblasts and pericytes, are quite diverse in morphology and function. Therefore, the regulatory mechanism of contractility is also likely to be diverse but has been studied only in a few representative systems.

The nematode Caenorhabditis elegans is a powerful model organism to study the structure and function of striated muscle, because worms have well-developed body-wall muscle with an organized sarcomeric structure that contains conserved myofibrillar proteins (Moerman and Fire, 1997; Waterston, 1988). C. elegans also has several non-striated muscles. In particular, the myoepithelial sheath of the somatic gonad is an interesting system, in which actin and myosin are organized in non-striated networks (Ardizzi and Epstein, 1987; Hall et al., 1999; Strome, 1986). The myoepithelial sheath surrounds oocytes, and its contraction is tightly coupled with oocyte maturation and is essential for ovulation (McCart et al., 1997; McCarter et al., 1999). However, the regulatory mechanism of ovulatory contraction of the myoepithelial sheath is poorly understood. Despite the fact that the myoepithelial sheath has non-striated actomyosin networks, many of the structural components are expressed from the same genes that are also expressed in the striated body-wall muscle (Ardizzi and Epstein, 1987; Ono and Ono, 2004; Ono et al., 2006; Ono et al., 2007; Rose et al., 1997). The unique feature of the myoepithelial sheath is that components of the troponin complex play major roles in regulating contractility. Troponin is a complex of troponin T (TNT; tropomyosin-binding), troponin I (TNI; inhibitory) and troponin C (TNC; Ca²⁺-binding), and regulates actomyosin contraction in an actin-linked and Ca²⁺-dependent manner generally in striated muscle (Ebashii, 1984; Gomes et al., 2002). Mutation of mup-2 (which encodes TNT) (Myers et al., 1996) or RNA interference of pat-10 (which encodes TNC) (Ono and Ono, 2004) causes sterility due to defective contraction of the myoepithelial sheath during ovulation. These observations suggest that a conventional troponin complex is an actin-linked regulator of contraction of non-striated actomyosin networks in the myoepithelial sheath. However, a TNI
component has not been identified in the *C. elegans* myoepithelial sheath.

TNI is the central component of the troponin complex (Perry, 1999). At low Ca\(^{2+}\) concentrations, TNI binds to actin filament and inhibits its interaction with myosin. At high Ca\(^{2+}\) concentrations, binding of Ca\(^{2+}\) to TNC induces dissociation of TNI from actin and allows actomyosin contraction. *C. elegans* has four TNI-encoding genes: *tni-1, unc-27* (also known as *tni-2*), *tni-3* and *tni-4*. Mutation in *unc-27* causes hypercontraction of the body-wall muscle (Burkeen et al., 2004; Ruksana et al., 2005), indicating that UNC-27 is a major TNI isoform in the body-wall muscle. Biochemical analysis of troponin from the giant nematode *Ascaris* showed that nematode TNI strongly inhibits ATPase activity of actomyosin (Kimura et al., 1987), suggesting that *C. elegans* TNI behaves in a similar manner. Therefore, when TNI is defective, actin and myosin are allowed to interact without regulation, which will lead to continuous muscle contraction. However, *unc-27*-null mutants are fertile and do not show a defect in the reproductive system. In this study, we aimed to identify TNI in the myoepithelial sheath and found that two TNI isoforms, TNI-1 and UNC-27, have partially redundant and essential functions in the myoepithelial sheath. This is the first in vivo demonstration that TNI is essential for reproduction and for regulation of contractility of non-striated muscle.

**Results**

Tropinin I is expressed in the *C. elegans* somatic gonad and colocalizes with non-striated actin networks

As a tool to detect *C. elegans* TNI, we re-evaluated the rabbit polyclonal antibody that was raised against *Ascaris* TNI (Kimura et al., 1987). A previous study has shown that this antibody reacts with *C. elegans* TNI (Nakae and Obinata, 1993), but this was before *C. elegans* TNI was molecularly characterized. *C. elegans* has four TNI genes, *tni-1, unc-27, tni-3* and *tni-4* (Burkeen et al., 2004; Ruksana et al., 2005). By western blot, we confirmed that this antibody reacted with a major band of TNI in wild-type worms (Fig. 1A). In addition, this antibody also recognized TNI isoforms in an *unc-27(e155)* (*unc-27-null*) mutant that lacks the major TNI isoform UNC-27 (Burkeen et al., 2004) (Fig. 1A). By comparing the bands of TNI in wild-type and *unc-27*-mutant worms, we detected at least four bands (Fig. 1A, bands 1-4). In wild-type worms, band 3 was predominant. Bands 2 and 4 were occasionally resolved from band 3 but these bands were often too minor to be separated clearly from band 3. In *unc-27-null* mutants, band 3 was absent, indicating that it corresponded to UNC-27 (Fig. 1A). Band 2 was the most intense band in *unc-27-null* mutants. Because band 2 migrated slower than band 3, band 4 was clearly separated from band 2 in *unc-27-null* mutants. In addition, band 1 was specifically detected in *unc-27-null* mutants (Fig. 1A). Therefore, UNC-27 is the major TNI isoform in wild-type worms. However, bands 1 and 2 in *unc-27-null* mutants are apparently increased as compared with those in wild type, suggesting that there was compensatory upregulation or stabilization of the second and third most abundant TNI isoforms in the absence of UNC-27. Our attempts to resolve TNI proteins by two-dimensional electrophoresis were not successful owing to inconsistent appearance of multiple spots that reacted with anti-TNI antibody, suggesting that TNI proteins are post-translationally modified.

By immunofluorescent staining of the *C. elegans* gonad using the anti-*Ascaris*-TNI antibody, we found that TNI is expressed in the myoepithelial sheath (Fig. 1B-G) and that it localized to non-striated actin networks (Fig. 1H-M). In dissected gonads from wild-type worms, TNI was specifically expressed in the myoepithelial sheath that surrounds oocytes in the proximal gonad (Fig. 1B) but not in the spermatheca, oocytes or distal arm containing non-myoepithelial sheath, germline cells and distal-tip cell (Fig. 1B-D). TNI colocalized to non-striated networks of actin filaments in the myoepithelial sheath (Fig. 1H-J). In an *unc-27*-mutant gonad, TNI was also specifically expressed in the myoepithelial sheath (Fig. 1E-J) and colocalized with actin filaments (Fig. 1K-M). These results indicate that TNI is a component of non-striated actin networks in the myoepithelial sheath, and a TNI isoform(s) other than UNC-27/UNC-2 is expressed in the gonad.

To determine which TNI isoforms are expressed in the myoepithelial sheath, we knocked down each TNI isoform in wild-type worms or *unc-27* mutants and examined TNI expression by western blot and immunofluorescence microscopy. Previous studies
have demonstrated that tni-4 is specifically expressed in the pharyngeal muscle (Burkeen et al., 2004; Ruksana et al., 2005). Therefore, we focused on tni-1, unc-27 and tni-3 in this study. In wild-type worms, neither tni-1(RNAi) nor tni-3(RNAi) had a detectable effect on the total TNI level as compared with control RNAi (Fig. 2A). By immunofluorescent staining of dissected wild-type gonads, it was observed that neither tni-1(RNAi) nor tni-3(RNAi) caused a detectable change in intensity and localization patterns of TNI (Fig. 2E,H) as compared with control-RNAi-treated gonads (Fig. 2B), suggesting that UNC-27 is a major TNI isoform in the myoepithelial sheath. In unc-27-null mutant worms, tni-1(RNAi) greatly reduced the TNI protein level, whereas tni-3(RNAi) caused no detectable change in the TNI level (Fig. 2A). In tni-1(RNAi)-treated unc-27 mutants, the top band of TNI was not detectable, and the most abundant middle band was significantly reduced (Fig. 2A), suggesting that the TNI-1 protein is resolved in these two bands. By immunofluorescent staining of dissected unc-27 gonads, tni-1(RNAi) resulted in a marked reduction of TNI in the myoepithelial sheath (Fig. 2N), whereas tni-3(RNAi) did not affect the staining patterns of TNI (Fig. 2Q) as compared with control-RNAi-treated gonads (Fig. 2K). Therefore, in the absence of UNC-27, TNI-1 is the major isoform in the myoepithelial sheath. Taken together, these results indicate that TNI-1 and UNC-27 are the two major TNI isoforms in the myoepithelial sheath.

Two TNI isoforms are functionally redundant and essential for ovulatory contraction of the somatic gonad

We found that simultaneous depletion of TNI-1 and UNC-27 caused sterility, indicating that these two TNI isoforms are functionally redundant and essential for reproduction in C. elegans. Wild-type worms that were treated with control RNAi (Fig. 3A), tni-1(RNAi) (Fig. 3B) or tni-3(RNAi) (Fig. 3C) produced ~250-300 progeny (brood size) (Fig. 3G). tni-1(RNAi) and tni-3(RNAi) only slightly reduced brood size (Fig. 3G). The unc-27-null mutant with control RNAi was fertile (Fig. 3D) but produced less progeny than wild type (Fig. 3G, compare control RNAi in wild type and unc-27; \( P<0.001 \)). However, tni-1(RNAi)-treated unc-27 mutants (Fig. 3E) were 33% (5/15) sterile with an average brood size of 3.1±2.6 per worm (\( n=10 \)) (Fig. 3G). tni-3(RNAi) did not affect brood size in the unc-27 mutant (Fig. 3F,G). These results strongly suggest that TNI-1 and UNC-27 play redundant and major roles in C. elegans reproduction.

To better understand the role of TNI in reproduction, we analyzed TNI-depleted phenotypes in the hermaphroditic gonad. The myoepithelial sheath, in which TNI is expressed (Fig. 1), provides contractile forces during ovulation to transport a mature oocyte into the spermatheca (McCarter et al., 1997; McCarter et al., 1999). Defective contraction of the myoepithelial sheath often causes accumulation of endomitotic oocytes in the proximal ovary because DNA replication repeatedly occurs in the absence of fertilization, and this phenotype is called Emo phenotype (endomitotic oocytes in the ovary) (Iwasaki et al., 1996). Indeed, Emo phenotype was observed when tropomyosin or TNC was depleted (Ono and Ono, 2004). However, Emo phenotype was not detected after depletion of tni-1 or tni-3 in wild-type or unc-27 worms (Fig. 4). In wild-type worms with control RNAi, oocytes with condensed chromosomes were linearly organized in the proximal ovary (Fig. 4A-C, arrowheads in B indicate oocyte nuclei), and tni-1(RNAi) or tni-3(RNAi) did not affect this morphology (Fig. 4D-I). The unc-27-
null mutant with control RNAi also had normal oocytes with condensed chromosomes in a linear arrangement (Fig. 4J-L). The tni-1(RNAi)-treated unc-27 mutant had oocytes with condensed chromosomes in the proximal ovary (Fig. 4M-O). We did not detect any abnormalities in the morphology of the nucleus and chromosomes (our unpublished data). In addition, the presence of some embryos in the uterus suggests that successful ovulation and fertilization took place (our unpublished data). The tni-3(RNAi)-treated unc-27 mutant showed no detectable difference in the morphology of the gonad from the unc-27 mutant with control RNAi (Fig. 4P-R).

Further analysis of the process of ovulation in live worms revealed that ovulatory contraction of the myoepithelial sheath was greatly weakened by depletion of TNI (Fig. 5). In wild-type worms with control RNAi (supplementary material Movie 1), upon oocyte maturation, which is characterized by nuclear-envelope breakdown (Fig. 5A,D), the proximal ovary vigorously contracted and pushed the oocyte into the spermatheca (Fig. 5G). Wild-type worms with tni-1(RNAi) or tni-3(RNAi) showed normal ovulation (Table 1, supplementary material Movie 2), although one case out of nine observed tni-1(RNAi)-treated wild-type worms exhibited weak ovarian contraction and failed ovulation (Table 1). unc-27-null-mutant worms with control RNAi also executed normal ovulation (Fig. 5B,E,H,K, supplementary material Movie 3). However, tni-1(RNAi)-treated unc-27 worms showed abnormal ovulation processes (Table 1). All of the observed worms (n=11) did not exhibit vigorous ovarian contraction after oocyte maturation (Fig. 5F,I, supplementary material Movie 4). One observed worm did not ovulate at all (Table 1). In most cases, an oocyte was apparently pushed into the spermatheca by tonic contraction and weak twitching of the proximal ovary (Fig. 5F,I, supplementary material Movie 4). Furthermore, in 45% (5/11) of the observed events, ovulation was incomplete and a portion of an oocyte was pinched off and retained in the ovary (Fig. 5L), indicating that contraction and relaxation of the myoepithelial sheath and dilation of the spermatheca were uncoordinated. unc-27 worms with tni-3(RNAi) showed normal ovulation (Table 1). These live observations indicate

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that TNI is required for vigorous ovarian contraction after oocyte maturation.

Localization of a myosin heavy chain in the myoepithelial sheath revealed that the sheath was hypercontracted in TNI-depleted gonads (Fig. 6). MYO-3 is a myosin heavy chain that is expressed in the myoepithelial sheath and localizes to the contractile apparatuses (Ardiszzi and Epstein, 1987). In wild-type gonads with control RNAi, thick filaments containing MYO-3 were distributed throughout the myoepithelial sheath, which covered ~ten oocytes in the proximal ovary (Fig. 6A-C). tni-1(RNAi) or tni-3(RNAi) in wild-type gonads did not alter this morphology (Fig. 6D-I). In unc-27-null gonads with control RNAi (Fig. 6J-L), the myoepithelial sheaths were shorter than those in wild-type gonads with control RNAi (Fig. 6, compare A and J). This indicates that the myoepithelial sheath was partially contracted by deficiency of UNC-27 alone, although ovulation was apparently normal (Fig. 5, supplementary material Movie 3). More strikingly, tni-1(RNAi)-treated unc-27 gonads showed much greater hypercontraction of the myoepithelial sheath (Fig. 6M-O). The myoepithelial sheath was so contracted that it covered only one to three oocytes (our unpublished data). By contrast, tni-3(RNAi) did not induce further contraction in the unc-27 mutant (Fig. 6P-R). These results indicate that TNI-1 and UNC-27 play major roles in inhibiting contraction of the myoepithelial sheath.

To exclude the possibility that the hypercontraction phenotype of the myoepithelial sheath is due to an artifact of dissection, fixation and/or immunostaining, we observed myoepithelial sheath in living worms using a fluorescent marker (Fig. 7). GFP-tagged PAT-10 (TNC) was expressed by the promoter of pppk-1 in living worms using a fluorescent marker (Fig. 7). GFP-tagged PAT-10 (TNC) was expressed by the promoter of pppk-1 in living worms using a fluorescent marker (Fig. 7). GFP-tagged PAT-10 (TNC) was expressed by the promoter of pppk-1 in living worms using a fluorescent marker (Fig. 7). GFP-tagged PAT-10 (TNC) was expressed by the promoter of pppk-1 in living worms using a fluorescent marker (Fig. 7). GFP-tagged PAT-10 (TNC) was expressed by the promoter of pppk-1 in living worms using a fluorescent marker (Fig. 7). GFP-tagged PAT-10 (TNC) was expressed by the promoter of pppk-1 in living worms using a fluorescent marker (Fig. 7). 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Table 1. Summary of ovulation phenotypes observed by time-lapse recording

<table>
<thead>
<tr>
<th></th>
<th>Wild-type</th>
<th>tni-1(RNAi)</th>
<th>tni-3(RNAi)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Weak ovarian contraction</td>
<td>0 (0/6)</td>
<td>11 (1/9)</td>
<td>0 (0/7)</td>
</tr>
<tr>
<td>No ovulation</td>
<td>0 (0/6)</td>
<td>11 (1/9)</td>
<td>0 (0/7)</td>
</tr>
<tr>
<td>Incomplete ovulation</td>
<td>0 (0/6)</td>
<td>45 (5/11)</td>
<td>0 (0/7)</td>
</tr>
</tbody>
</table>

*aAll phenotypes were scored after oocyte maturation occurred.

*bIncomplete ovulation resulted in cleavage of an oocyte and retention of part of the oocyte in the ovary.

Data are shown as percentages (number of worms with a phenotype/total number of observed worms).
distinguishable in unc-54(s95);unc-27(e155) with tni-1(RNAi) gonads (Fig. 8J), indicating that TNI depletion did not disturb overall organization of the myoepithelial sheath. Importantly, the partial suppression of the hypercontraction phenotype in TNI-depleted gonad by a myosin mutation strongly suggests that the phenotype is due to unregulated actin-myosin interaction in the absence of inhibitory function of TNI.

Furthermore, we found that the contraction states of the myoepithelial sheath were correlated with a reduction in the number of oocytes in the proximal gonad in TNI-depleted worms. Wild-type worms with control-RNAi treatment had 9 to 11 oocytes per gonadal arm (Fig. 9A,G), and tni-1(RNAi) or tni-3(RNAi) only slightly reduced the number of oocytes (Fig. 9C,E,G). unc-27-mutant worms with control RNAi treatment had five to seven oocytes per gonadal arm (Fig. 9B,G). Thus, loss of UNC-27 is sufficient to cause a reduction of oocytes. Furthermore, unc-27 mutant worms with tni-1(RNAi) had only one to three oocytes per arm (Fig. 9D,G), whereas tni-3(RNAi) in the unc-27 background did not further reduce the number of oocytes (Fig. 9F,G). Thus, hypercontraction of the myoepithelial sheath causes a negative impact on oocyte production, suggesting that maintaining a relaxed state of the myoepithelial sheath is important for hosting an environment for proper oocyte development.

Association of TNI with thin filaments depends on tropomyosin but not on TNC

Tropomyosin mediates association of the troponin complex with actin filaments (Ebashi and Kodama, 1966). We previously showed that PAT-10 requires tropomyosin for it to localize to actin filaments in the myoepithelial sheath (Ono and Ono, 2004). However, TNI can directly bind to actin filaments in vitro (Perry, 1999), and it is not clear whether this direct binding is sufficient for TNI to localize to actin filaments in vivo. We found that tropomyosin was required for association of TNI with actin filaments in the myoepithelial sheath (Fig. 10). When tropomyosin was depleted by RNAi, TNI was detected in the diffuse cytoplasm and was no longer associated with filamentous actin (Fig. 10D-F), as compared with clear filamentous localization of TNI in control-RNAi-treated gonad (Fig. 10A-C). In addition, when PAT-10 was depleted, filamentous staining of TNI was slightly weaker than that in control-RNAi-treated gonad (Fig. 10G-I), indicating that PAT-10 is not required for association of TNI with actin filaments in vivo but might enhance or stabilize binding of TNI to the filaments. These results suggest that TNI functions as a component of the troponin complex that is anchored to actin filaments via tropomyosin.

Discussion

In this study, we identified TNI genes that control contraction and relaxation of non-striated muscle (myoepithelial sheath) in the C. elegans somatic gonad. Among the four TNI genes, two of them, tni-1 and unc-27, play major roles in the gonad. TNI was associated with non-striated networks of actin filaments in the myoepithelial sheath, and this association was dependent on tropomyosin.
Depletion of the two TNI isoforms resulted in sterility with excessive contraction of the proximal ovary. These results demonstrate that two TNI isoforms have partially redundant functions and are crucial for regulation of muscle contraction during ovulation.

Identification of TNI in the C. elegans gonad and previous studies on TNT (Myers et al., 1996) and TNC (Ono and Ono, 2004) strongly suggest that the conventional troponin complex consisting of TNT, TNI and TNC regulates contraction of the myoepithelial sheath during ovulation by an actin-linked mechanism. Each component of the troponin complex is essential for C. elegans reproduction. TNC and TNI are expressed in the myoepithelial sheath and localize to the actin thin filaments, although TNT has not been characterized at a protein level. However, the TNI-depletion phenotype was different from the TNT-mutant and TNC-RNAi phenotypes. When TNT or TNC is defective, ovulation fails, and endomitotic oocytes are accumulated in the proximal ovary (Myers et al., 1996; Ono and Ono, 2004). However, when TNI is defective, mature oocytes are often incompletely ovulated, and the myoepithelial sheath is constantly hypercontracted. TNI directly interacts with actin and inhibits its interaction with the myosin head at low Ca^{2+} concentrations, and Ca^{2+}-bound TNC releases its inhibitory action and allows actin-myosin interaction (Ehashi, 1984). Because we observed that TNI was still associated with actin filaments in TNC-depleted gonads, TNI could still inhibit actomyosin contraction in the absence of TNC. However, when TNI was depleted, actin and myosin could interact regardless of changes in Ca^{2+} concentrations, thereby allowing constant contraction of the myoepithelial sheath. RNAi does not generally deplete 100% of a target protein, and residual TNI in tni-1(RNAi)-treated unc-27 mutants might be sufficient to support cycles of contraction and relaxation during ovulation. Thus, the phenotypic consequences of depletion of troponin components are consistent with known biochemical properties of the conventional troponin complex.

Decrease in the number of oocytes in the proximal ovary by depletion of TNI was an unexpected phenotype. Because ovulation was incomplete but partially successful in TNI-depleted worms, insufficient supply of oocytes seemed to be the major cause of sterility. In wild-type worms, the proximal ovary normally holds 10-12 oocytes, which can keep up with ovulation that occurs every 20-30 minutes (McCarter et al., 1997; McCarter et al., 1999). In unc-27-null-mutant worms, the number of oocytes was decreased to five to seven, but brood size was only slightly less than that of wild type under the same culture conditions. However, tni-1(RNAi)-treated unc-27 mutants had only one to three oocytes in the ovary and, after ovulation of an oocyte, the next oocyte was often absent or immature. Interestingly, the decrease in oocytes was correlated with the extent of contraction of the myoepithelial sheath. unc-27 mutants with control RNAi or tni-3(RNAi) showed a partially contracted myoepithelial sheath and a decreased number of oocytes, and unc-27 mutants with tni-
showed further hypercontraction and decrease in the number of oocytes. These observations suggest that maintaining a relaxed state of the myoepithelial sheath is important for oocyte development. Gonadal sheath cells play important roles in germline development. In particular, distal sheath cells promote proliferation of the germline in the distal gonad (Killian and Hubbard, 2005). Contraction of the myoepithelial sheath might affect the function of distal sheath cells, which might result in a reduction in the number of germ cells. In addition, the proximal ovary is an important organ for oogenesis; it is here that oocytes gain cytoplasmic components from the distal gonad through the rachis (Wolke et al., 2007) and yolk proteins from the body cavity by endocytosis (Grant and Hirsh, 1999). Therefore, excessive contraction of the myoepithelial sheath might physically limit the number of residing oocytes and prevent these crucial processes in oogenesis.

To our knowledge, the C. elegans myoepithelial sheath is the only non-striated muscle in which the function of troponin is genetically demonstrated. Ascidian smooth muscle also expresses a functional troponin complex (Endo and Obinata, 1981), but it has not been characterized by a genetic approach. Otherwise, troponin is predominantly expressed in striated muscle and has been widely accepted as a striated-muscle-specific regulator of contraction (Ebashi, 1984; Squire and Morris, 1998). However, recently, expression of troponin in mouse vascular smooth muscle has been reported (Moran et al., 2008), although its function is still unclear. Furthermore, TNI is involved in the regulation of chromosomal stability and cell polarity in early Drosophila embryos (Sahota et al., 2009). Thus, additional cell-biological and genetic investigation on troponin in non-striated muscle and non-muscle cells might reveal unknown functions of the troponin complex or individual troponin components.

Cell contractility is also implicated in mammalian ovulation. In mammalian ovaries, each oocyte is surrounded by a follicle that contains an external layer of smooth-muscle-like cells (Espey, 1564 Journal of Cell Science 123 (9)

Fig. 9. Depletion of TNI reduces the number of oocytes in the proximal gonad. Wild-type (A, C and E) or unc-27-null (B, D and F) worms were treated with control-RNAi (A and B), tni-1(RNAi) (C and D) or tni-3(RNAi) (E and F), and Nomarski micrographs of the proximal gonads are shown. Asterisks indicate oocyte nuclei. (G) Number of oocytes per gonadal arm (n=50). A t-test was performed on control RNAi and tni-1(RNAi) or tni-3(RNAi) for each strain. Double and triple asterisks indicate 0.001<P<0.01 and P<0.001, respectively.

Fig. 10. Association of TNI with actin filaments is dependent on tropomyosin in the myoepithelial sheath. (A-I) Wild-type worms were treated with control RNAi (A-C), lev-11(RNAi) (RNAi for tropomyosin) (D-F) or pat-10(RNAi) (RNAi for TNC) (G-I), and dissected gonads were stained for TNI (A, D, and G) and F-actin (B, E, and H). Merged images are shown in C, F and I (TNI in green and F-actin in red).
1978). These cells express markers of smooth muscle, including smooth-muscle α-actin and desmin, but the function of these cells is not understood. A recent report suggests that these smooth-muscle-like cells are indeed contractile and contribute to rupture of a follicle at the late step of ovulation in the mouse ovary (Ko et al., 2006). Therefore, although the structure of the gonads and the process of ovulation are different between nematodes and mammals, there might be a similarity in the mechanisms of regulation of cell contractility during ovulation. Involvement of troponin in mammalian ovulation is unknown, but there is a possibility that cell contraction is regulated by an actin-linked mechanism during ovulation. Further analysis of the mechanisms of ovulation in different species might reveal evolutionarily conserved roles of actin-associated proteins in reproductive systems.

Materials and Methods

Nematode strains

Wild-type strain N2, an unc-27-null mutant CB155 unc-27(e155) and an unc-54 null mutant RW5008 unc-54(s95) were obtained from the Caenorhabditis Genetic Center (Minneapolis, MN). An unc-54(s95);unc-27(e155) double mutant was generated by standard crosses. A transgenic line carrying an extrachromosomal array kgEx61[pkk-1::pat-10::gfp, pRK74] (Xu et al., 2007) was provided by Myeongwoo Lee (Baylor University, Waco, TX). The transgene was placed in the unc-27(e155) background by standard crosses. Nematodes were grown under standard conditions at 20°C as described previously (Brenner, 1974).

RNAi experiments

RNAi experiments were performed by feeding with E. coli expressing double-stranded RNA as described previously (Ono and Ono, 2002). RNAi clones for mvu-1 (X-5B01) and tni-3 (V-11A06) were described previously (MRC Geneservice, Kamath et al., 2003). RNAi clones for C. elegans tropomyosin (lev-11) targeting four isotypes (previously designated as TM2) and pat-10 were described previously (Ono and Ono, 2004; Ono and Ono, 2002). L4 larvae were treated with RNAi, and phenotypes were characterized in the F1 generation.

Western blot

Western blot analysis of TNI was performed essentially as described (Ono and Ono, 2009) with modifications. Briefly, ten adult worms were lysed in SDS-lysis buffer, and subjected to SDS-PAGE (12% acrylamide gel) and blotting. Rabbit anti-C. elegans mvu-1 antibody (Nakae and Obinata, 1993) was diluted to 1:2000 to 1:8000 in Signal Enhancer HKARI (Nacalai USA) and used to detect C. elegans TNI. Mouse anti-actin monoclonal antibody (C4; MP Biomedicals) was used to detect equal loading of the samples. Horseradish-peroxidase-labeled secondary antibodies were used to detect primary antibodies. The reactivity was detected with a SuperSignal West Pico chemiluminescence reagent (Pierce Chemical).

Fluorescence microscopy

The gonads were dissected by cutting adult hermaphrodites at the level of pharynx on poly-L-lysine-coated slides as described previously (Rose et al., 1997). Immunostaining of dissected gonads was performed essentially as described (Ono et al., 2008; Ono et al., 2007) with some modifications. The samples were fixed with 4% paraformaldehyde in cytoskeleton buffer (138 mM KCl, 3 mM MgCl2, 2 mM et al., 2008; Ono et al., 2007) with some modifications. The samples were fixed with a Nikon Eclipse TE2000 inverted microscope and observed with a 40X CFI Plan Fluor objective (N.A. 1.4). Images were captured at room temperature by a SPOT Idea CMOS camera (Diagnostic Instruments) and recorded at five frames per second for 30 minutes by the SPOT Imaging Software (Diagnostic Instruments). Movie files were saved in a compressed AVI format at 20 frames per second (four times as fast as real time) and phenotypes analyzed later. Movie files were edited and converted to QuickTime files using Adobe Premiere 6.5.

Monoclonal antibody 5-6 was developed by Henry Epstein (University of Texas Medical Branch, Galveston), and was obtained from the Developmental Studies Hybridoma Bank, developed under auspices of the NICHD and maintained by the University of Iowa, Department of Biological Sciences, Iowa City, IA. C. elegans strains were provided by Caenorhabditis Genetics Center, which is supported by the National Institute of Health National Center for Research Resources. We thank Myeongwoo Lee (Baylor University, Waco, TX) for kindly providing us with the GFP-pat-10 transgenic worms. This work was supported by grants from the Ministry of Education, Science and Culture, Japan (Grant number 20570075) to T.O. and the National Institute of Health (R01 AR48615) and University Research Committee of Emory University to S.O. Deposited in PMC for release after 12 months.

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References


Time-lapse Nomarski microscopy

Worms were anesthetized in 0.1% tricaine, 0.01% tetramisole in M9 buffer for 30 minutes and mounted on 2% agarose pads (McCarter et al., 1997). Tricaine and tetramisole paralyzes body-wall movement but does not block two to three consecutive rounds of oocyte maturation and ovulation. They were set on a Nikon Eclipse TE2000 inverted microscope and observed with a 40X CFI Plan Fluor objective (N.A. 1.4). Images were captured at room temperature by a SPOT Idea CMOS camera (Diagnostic Instruments) and recorded at five frames per second for 30 minutes by the SPOT Imaging Software (Diagnostic Instruments). Movie files were saved in a compressed AVI format at 20 frames per second (four times as fast as real time) and phenotypes analyzed later. Movie files were edited and converted to QuickTime files using Adobe Premiere 6.5.


