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UNC-87 isoforms, *Caenorhabditis elegans* calponin-related proteins, interact with both actin and myosin and regulate actomyosin contractility

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**ABSTRACT** Calponin-related proteins are widely distributed among eukaryotes and involved in signaling and cytoskeletal regulation. Calponin-like (CLIK) repeat is an actin-binding motif found in the C-termini of vertebrate calpons. Although CLIK repeats stabilize actin filaments, other functions of these actin-binding motifs are unknown. The *Caenorhabditis elegans* unc-87 gene encodes actin-binding proteins with seven CLIK repeats. UNC-87 stabilizes actin filaments and is essential for maintenance of sarcomeric actin filaments in striated muscle. Here we show that two UNC-87 isoforms, UNC-87A and UNC-87B, are expressed in muscle and nonmuscle cells in a tissue-specific manner by two independent promoters and exhibit quantitatively different effects on both actin and myosin. Both UNC-87A and UNC-87B have seven CLIK repeats, but UNC-87A has an extra N-terminal extension of \(\sim 190\) amino acids. Both UNC-87 isoforms bind to actin filaments and myosin to induce ATP-resistant actomyosin bundles and inhibit actomyosin motility. UNC-87A with an N-terminal extension binds to actin and myosin more strongly than UNC-87B. UNC-87B is associated with actin filaments in nonstriated muscle in the somatic gonad, and an *unc-87* mutation causes its excessive contraction, which is dependent on myosin. These results strongly suggest that proteins with CLIK repeats function as a negative regulator of actomyosin contractility.

**INTRODUCTION** Muscle contractile apparatuses are highly organized, with properly arranged actin and myosin filaments, together with regulatory components for contractility (Clark et al., 2002). To produce efficient contractile forces, bipolar myosin filaments and actin filaments with proper orientations need to be assembled into sarcomeric patterns. Additional structural and regulatory proteins support integrity and dynamics of the contractile structures and modulate the actin–myosin interaction (Squire, 1997; Ono, 2010). Such mechanisms are important not only in muscle but also in nonmuscle cells when transient or persistent contractile apparatuses are required to support cell migration, cell division, and morphogenesis (Pollard and Cooper, 2009). However, how actomyosin-based contractile structures are assembled and regulated in vivo is not fully understood.

Calponin is a cytoskeletal protein that is expressed in both muscle and nonmuscle cells and has biochemical activities to stabilize actin filaments and regulate actomyosin contractility (Rozenblum and Gimona, 2008). Of note, calponin has been characterized as a major component of smooth muscle thin filaments in vertebrates (Takahashi et al., 1988). In vitro, calponin binds to actin filaments and myosin and inhibits actomyosin ATPase in a \(\text{Ca}^{2+}\)-calmodulin–regulated manner (Abe et al., 1990; Winder and Walsh, 1990). Consistently, in skinned smooth muscle cells, calponin is required for maintaining a relaxed state by inhibiting contractility caused by unphosphorylated myosin (Malmqvist et al., 1997). However, in vivo, calponin-knockout mice show only mild

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Abbreviations used: ADF, actin-depolymerizing factor; Ce-myosin, *Caenorhabditis elegans* myosin; CH, calponin homology; CLIK, calponin-like; GFP, green fluorescent protein; R-myosin, rabbit skeletal muscle myosin.

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alterations in smooth muscle contractility (Matthew et al., 2000; Takahashi et al., 2000; Yoshimoto et al., 2000). This may be due to the presence of functionally overlapping proteins such as caldesmon (Hodgkinson, 2000) and/or structurally related proteins such as SM22/transgelin (Pearlstone et al., 1987).

Vertebrate calponins have one calponin-homology (CH) domain in the N-terminal and three calponin-like (CLIK) repeats (or motifs) in the C-terminus (Rozenblum and Gimona, 2008). Although CH domains are found in a number of signaling or cytoskeletal proteins and function as an actin-binding domain in many cases, the CH domain of calponin does not bind to actin (Gimona and Mital, 1998; Galkin et al., 2006). Instead, the CLIK repeats in the C-terminus of calponins are essential for binding to actin filaments (Gimona and Mital, 1998). The number of CLIK repeats in naturally occurring proteins correlates with the strength of actin-binding. SM22/transgelin, which has only one CLIK repeat, binds to actin at lower affinity than calponin, which has three CLIK repeats (Shapland et al., 1993). UNC-87 in the nematode Caenorhabditis elegans is a unique protein with seven CLIK repeats and no CH domain (Goetinck and Waterston, 1994a), which binds to actin filaments with high affinity (Kranewitter et al., 2001). Of importance, proteins with multiple CLIK repeats stabilize actin filaments and prevent depolymerization in vitro (Kake et al., 1995; Yamashiro et al., 2007). Studies in mammalian cells have shown a correlation between the number of CLIK repeats and the actin-stabilizing function (Gimona et al., 2003) and as a consequence to inhibit cell migration and cytoskeleton (Lener et al., 2004).

The biochemical analysis of C. elegans UNC-87 protein unambiguously demonstrated the function of the CLIK repeat as an actin-binding motif (Kranewitter et al., 2001). The C. elegans unc-87 gene is essential for proper sarcomeric actin organization in larval and adult striated muscle but not for initial sarcomere assembly in embryos (Goetinck and Waterston, 1994a,b). UNC-87 competes with actin-depolymerizing factor (ADF)/cofilin for binding to actin filaments and protects actin filaments from severing by ADF/cofilin (Yamashiro et al., 2007). Although the actin-stabilizing role of UNC-87 is similar to that of tropomyosin (Ono and Ono, 2002; Yu and Ono, 2006), UNC-87 and tropomyosin apparently have opposite roles in sarcomere organization in striated muscle (Yamashiro et al., 2007), suggesting that UNC-87 has an additional, uncharacterized function other than actin-filament stabilization. Furthermore, the initial analysis of the unc-87 gene identified two gene products with different N-terminal sequences, but functional differences between the two isoforms are unknown. In this study, we report that the two UNC-87 isoforms are expressed in different tissues in C. elegans and bind to both actin and myosin. This results in inhibition of actomyosin motility and formation of ATP-resistant actomyosin bundles. Genetic analysis shows that UNC-87B is a negative regulator of myosin-dependent contractility of smooth muscle-like cells in the somatic gonad. These results reveal a novel function of UNC-87 isoforms as regulators of actomyosin contractility.

RESULTS
Two alternative promoters drive expression of two UNC-87 isoforms in a tissue-specific manner
The original report of the molecular characterization of the unc-87 gene demonstrated that two isoforms are expressed from the unc-87 gene by an alternative choice of the first two exons (Goetinck and Waterston, 1994a; Supplemental Figure S1, A–C). According to WormBase (www.wormbase.org), the large and small isoforms have been designated UNC-87A (565 amino acids) and UNC-87B (374 amino acids), respectively. To maintain consistency, the downstream first exon used in UNC-87A is designated exon 1A, and the upstream exon used in UNC-87B is designated exon 1B (Supplemental Figure S1A). To determine the mechanism of isoform-specific expression, we examined whether exon 1A or 1B is selectively expressed by independent promoters. We hypothesized that the flanking upstream sequence of each first exon contains promoter activity. To test this, we fused 2-kb upstream sequences from exons 1A and 1B with green fluorescent protein (GFP) as a reporter (Supplemental Figure S1, D and E) and examined their promoter activity in vivo.

The reporter analysis showed that expression of exons 1A and 1B was driven by separate promoters in a tissue-specific manner. The 2-kb upstream sequence from exon 1A, which is entirely downstream of exon 1B (Supplemental Figure S1D; Punc-87A::GFP), promoted expression of GFP in the pharynx (Figure 1, A–D), anal depressor muscle (Figure 1, A, E, and F), uterine muscle (Figure 1, A, G, and H), vulva (Figure 1, G and H), and unidentified neurons in the head and the ventral region (Figure 1, A, C, and D). By contrast, the 2-kb upstream sequence from exon 1B (Supplemental Figure S1E; Punc-87B::GFP) promoted expression of GFP in the body wall muscle (Figure 1, I–L), spermatheca (Figure 1, I, J, M, and N), and vulva (Figure 1, I–L). Because the hermaphroditic gonads were obscured by strong GFP signals in the body wall muscle, expression of GFP was further analyzed in dissected gonads by immunofluorescence staining for GFP and MYO-3, a myosin heavy chain expressed in the myoepithelial sheath (Ardizio and Epstein, 1987). The results showed predominant expression of GFP from the unc-87B promoter both in the myoepithelial sheath (MYO-3 positive) and the spermatheca (MYO-3 negative; Figure 1, U–W) but not from the unc-87A promoter (Figure 1, R–T) or in the gonad with no transgene (Figure 1, O–Q). This results suggest that separate promoters for unc-87A and unc-87B are tissue specific in a mutually exclusive manner except for the vulva and control expression of the two UNC-87 isoforms.

Previous studies reported that unc-87 mutations cause disorganized sarcomeric actin filaments in the body wall muscle (Goetinck and Waterston, 1994a,b), but the nature of mutation sites has not been characterized at a molecular level. We determined mutation sites in two unc-87 alleles: unc-87(e1216) shows mild defects in body wall muscle, and unc-87(e1459) shows very severe defects (Goetinck and Waterston, 1994a). We sequenced the genomic sequences and found that unc-87(e1216) had a point mutation (G to A) at the splice donor site in exon 2 (Supplemental Figure S1F) and unc-87(e1459) had two point mutations in exon 2, a missense mutation (A207V in unc-87A and A16V in unc-87B) and a nonsense mutation (W225stop in UNC-87A or W34stop in UNC-87B). Therefore these mutations are located in common exons for UNC-87A and UNC-87B and should affect both isoforms. unc-87(e1216) expresses reduced levels of UNC-87 proteins (Goetinck and Waterston, 1994a), suggesting that the mutation reduces efficiency of splicing but does not completely inhibit protein expression, thereby acting as a weak loss-of-function allele. By contrast, unc-87(e1459) does not express detectable UNC-87 proteins (Goetinck and Waterston, 1994a), indicating that this is a null or a strong loss-of-function mutant.

UNC-87A, a longer isoform, exhibits stronger actin-bundling activity than UNC-87B
Previous biochemical studies on UNC-87 were done using UNC-87B, a shorter isoform (Kranewitter et al., 2001; Yamashiro et al., 2007), and characterization of UNC-87A has not been reported. To determine whether the two UNC-87 isoforms have any different
FIGURE 1: Promoter analysis of unc-87A and unc-87B. (A–N) Promoter activity of unc-87A (A–H) and unc-87B (I–N) was examined using GFP as a reporter. Each fluorescence micrograph of GFP (black and white) is paired with a differential interference contrast image overlaid with green fluorescence of GFP. Images of whole worms with their heads on the left (A, B, I, and J) and magnified views of representative tissues (C–H, K–N) are shown. Note that the micrograph in C is somewhat overexposed for GFP in the pharynx to demonstrate relatively weak GFP in the neurons. (O–W) Expression of GFP in hermaphrodite gonads with no transgene (O–Q), Punc-87A::GFP (R–T), or Punc-87B::GFP (U–W) was examined by immunostaining of dissected gonads for GFP (O, R, and U) and MYO-3 as a marker of myoepithelial sheath (P, S, and V). Merged images are shown in Q, T, and W (green, GFP; red, MYO-3; blue, DNA [not shown as individual images]). Scale bars, 100 μm (whole worms; A, B, I, and J), 20 μm (magnified tissues; C–H, K–N), 50 μm (dissected gonads; O–W).
biochemical properties, we produced and purified recombinant UNC-87A protein and compared it biochemically with UNC-87B.

UNC-87B has been shown to bind to F-actin and bundle the filaments (Kranewitter et al., 2001). Similarly, UNC-87A bound to F-actin, as determined by cosedimentation assays using ultracentrifugation (Figure 2A), and bundled actin filaments, as determined by sedimentation assays using low-speed centrifugation (Figure 2B). In the cosedimentation assays using ultracentrifugation, UNC-87A bound to F-actin so strongly that nearly all UNC-87A cosedimented with F-actin at low UNC-87A concentrations (Figure 2A; 1.1 μM UNC-87A to 10 μM actin), which makes it difficult to estimate its affinity for F-actin. In the sedimentation assays using low-speed centrifugation, in which only bundled actin filaments precipitate, UNC-87A induced bundle formation of actin filaments (10 μM actin) at >1.1 μM (Figure 2C, black circles). By comparing these data with the published data on UNC-87B (Yamashiro et al., 2007), we see that UNC-87A showed much stronger actin-bundling activity than UNC-87B, in particular, at a low concentration range (<2.5 μM; Figure 2C).

**UNC-87 isoforms bind to both actin and myosin and inhibit actomyosin ATPase and motility**

To explore a novel function of UNC-87 proteins, we examined their interaction with myosin. We purified myosin from *C. elegans* (Ce-myosin) and found that Ce-myosin by itself remained highly soluble even under low-salt conditions (27 mM KCl), as determined by low-speed centrifugation (18,000 × g, 30 min; Figure 3, A and B), under which rabbit skeletal muscle myosin (R-myosin) readily precipitated (unpublished data). However, UNC-87A induced precipitation of Ce-myosin and coprecipitated with Ce-myosin (Figure 3A), although UNC-87A remained in the supernatants in the absence of Ce-myosin (Figure 3A). These results strongly suggest that UNC-87A bound to Ce-myosin and induced formation of bundles or aggregates of Ce-myosin. UNC-87B also induced precipitation of Ce-myosin (Figure 3B). Comparison of the myosin precipitation at various concentrations of UNC-87A and UNC-87B showed that UNC-87A exhibited stronger activity than UNC-87B (Figure 3C).

The myosin-binding activity of UNC-87 isoforms prompted us to test whether UNC-87 proteins can cross-link actin filaments with myosin. We tested this by observing morphology of fluorescently labeled actin filaments by microscopy in the presence or absence of UNC-87 proteins and/or Ce-myosin (Figure 4). In the absence of Ce-myosin or UNC-87, only single actin filaments were detected (Figure 4A), and ATP did not alter the morphology of actin filaments (Figure 4B). Ce-myosin induced formation of short, needle-like bundles of actin filaments in the absence of ATP (Figure 4C), but these bundles were nearly completely disassembled in the presence of ATP (Figure 4D). These are consistent with conserved properties of conventional myosin: actin-filament bundling by two-headed myosin molecules and ATP-dependent dissociation of myosin heads from actin filaments. By contrast, both UNC-87A (Figure 4E) and UNC-87B (Figure 4F) induced formation of much larger and longer actin bundles than Ce-myosin-induced bundles (Figure 4C), and UNC-87-induced actin bundles were resistant to ATP (Figure 4, F and J). Of interest, when both Ce-myosin and UNC-87 proteins were simultaneously present, more-robust formation of actin bundles was observed (Figure 4, G and K). Actin bundles were much larger in the presence of both Ce-myosin and UNC-87 than in the presence of Ce-myosin or UNC-87 alone. In addition, a number of unbundled single actin filaments remained in the presence of UNC-87A or UNC-87B alone (Figure 4, E and I), but unbundled filaments were almost undetectable in the presence of both Ce-myosin and UNC-87A or UNC-87B.
Next we examined how UNC-87 proteins affect actomyosin ATPase activity. As reported previously (Harris et al., 1977; Tani et al., 1985), Ce-myosin had much lower ATPase activity than R-myosin, and actin modestly enhanced the ATPase activity (Figure 5, A and C). Actin activation of Ce-myosin ATPase was 13–28% (Figure 5, B and D). Both UNC-87A (Figure 5, A and B) and UNC-87B (Figure 5, C and D) inhibited activation of myosin ATPase by actin. Because actin activation of Ce-myosin ATPase was modest, we also examined the effect of UNC-87 isoforms on actin-activated R-myosin ATPase, in which actin activation of R-myosin ATPase was >20-fold (Figure 5, E and F). Under these conditions, both UNC-87A (Figure 5E) and UNC-87B (Figure 5F) significantly inhibited actin-activated R-myosin ATPase. In cosedimentation assays with R-myosin, both UNC-87 isoforms only weakly interacted with R-myosin (unpublished data), suggesting that the strong inhibition of actin-activated myosin ATPase is due to strong interactions between actin and UNC-87A or UNC-87B.

The inhibitory functions of UNC-87 isoforms on actomyosin ATPase strongly suggest that they also inhibit actomyosin contractility. We examined effects of UNC-87A and UNC-87B on actomyosin motility by in vitro motility (gliding) assays (Figure 6). DyLight 549–labeled actin filaments were attached to myosin on a glass surface, incubated with a buffer with or without UNC-87A or UNC-87B for 2 min, and then exposed to a buffer containing ATP with no UNC-87 proteins to initiate motility. In Figure 6, A–C, actin filaments at time 0 and after 4 s were pseudocolored by red and green, respectively, and overlaid in single images. In control with no UNC-87 proteins, most actin filaments were displaced, and overlaps between red and green filaments were minimal (Figure 6A). By contrast, both UNC-87A (Figure 6B) and UNC-87B (Figure 6C) at 2.5 μM significantly increased populations of yellow (red plus green) filaments, indicating that actomyosin motility was inhibited by the UNC-87 isoforms. Quantitative analysis of the velocity of the filament movement showed that 2.5 μM UNC-87A or UNC-87B significantly inhibited actomyosin motility (Figure 6D). At 1.0 μM, both UNC-87A and UNC-87B had weaker effects on actomyosin motility (Figure 6D) and did not cause statistically significant changes in the mean values as estimated by one-way analysis of variance. However, it should be noted that velocity of the bottom 25th percentile was reduced to less than half by 1.0 μM UNC-87A (0.133 μm/s) or UNC-87B (0.147 μm/s) as compared with control (0.302 μm/s), suggesting that populations of slow-moving filaments were increased by 1.0 μM UNC-87A or UNC-87B. Pairwise comparisons between UNC-87A and UNC-87B at the same concentrations showed no statistically significant differences between the two isoforms. Overall this biochemical analysis strongly suggests that UNC-87 isoforms are negative regulators of actomyosin contraction.

unc-87 mutations cause excessive contraction in the somatic gonad

To determine whether the inhibitory roles of UNC-87 proteins for actomyosin ATPase and motility are functionally significant in vivo, we examined muscle contractility in unc-87 mutant worms. Previous studies showed that unc-87 mutant worms move much more slowly than wild-type worms, which is primarily due to defects in the body wall muscle (Goetinck and Waterston, 1994a). However, body wall muscle cells are firmly attached to relatively rigid cuticles (Ono, 2014), and extent of muscle contraction could not be easily quantitated. In addition, unc-87 mutation causes disorganization of to ATP (Figure 4, H and L), suggesting that UNC-87 proteins play a role in maintaining actomyosin structures when the ATP-sensitive actin–myosin association and dissociation occur.
Both wild type and *unc-87(e1459)* had six myoepithelial sheath cells, excluding the possibility that the number of cells made the difference. Actin and myosin (MYO-3) were organized into networks in both wild type (Figure 8, B–D) and *unc-87(e1459)* (Figure 8, J–L). However, myosin filaments appeared more densely aligned in *unc-87(e1459)* than in wild type (Figure 8C), which is consistent with the contraction of the actomyosin network in *unc-87(e1459)*. To determine whether this shortening is due to actomyosin contractility, we examined the effects of a mutation in *unc-54*, a gene encoding another major myosin heavy chain expressed in the myoepithelial sheath (Ardizzi and Epstein, 1987). *unc-54(s95)* is a missense mutation near the ATP-binding site in the myosin head and reduces muscle contractility without altering organization of the contractile apparatus (Moerman et al., 1982; Dibb et al., 1985). *unc-54(s95)* single mutant had slightly shorter myoepithelial sheath (Figure 8E and Table 1) and *unc-87(e1459) unc-54(s95)* double mutant had significantly longer myoepithelial sheath than *unc-87(e1459)* single mutant (compare Figure 8, I and M), indicating that the myosin mutation suppressed the Unc-87 phenotype. The *unc-54(s95)* mutation did not affect the integrity of the actomyosin networks (Figure 8, F–H), and the myosin filaments in *unc-87(e1459) unc-54(s95)* double mutant (Figure 8O) exhibited a more relaxed appearance than those in *unc-87(e1459)* (Figure 8K). Thus, these genetic evidence strongly suggests that *unc-87B* is a negative regulator of actomyosin contractility in vivo.

FIGURE 4: UNC-87 isoforms induce large actomyosin bundles. DyLight 549–labeled actin filaments (2 μM) were incubated without (A–D) or with 1.0 μM UNC-87A (E–H) or 1.0 μM UNC-87B (I–L) in the absence (left two columns) or presence of 0.19 μM Ce-myosin (right two columns) and in the absence (A, C, E, G, I, and K) or presence (B, D, F, H, J, and L) of 5 mM ATP. The actin filaments were directly observed by fluorescence microscopy. Bar, 20 μm.
both actin and myosin to inhibit actomyosin motility and form ATP-insensitive actin–myosin bundles. The inhibitory role of UNC-87 isoforms for actomyosin contractility was further demonstrated in vivo: unc-87 mutants showed excessive contraction of the myoepithelial sheath, a smooth muscle–like tissue in the somatic gonad. In addition to the previously reported function of UNC-87 to stabilize actin filaments (Kranewitter et al., 2001; Yamashiro et al., 2007), the present study revealed another function of UNC-87 to regulate actomyosin contractility. In vertebrate smooth muscle, similar dual functions have been demonstrated in vitro for calponin (Winder et al., 1998), caldesmon (Hemric and Chalovich, 1988; Velaz et al., 1989), and fesselin/synaptopodin-2 (Schroeter and Chalovich, 2005), but their in vivo functions are not clearly understood. UNC-87A and UNC-87B are the only known actin-bundling proteins in C. elegans that also inhibit actomyosin contractility. Our biochemical and genetic studies in C. elegans suggest that UNC-87 isoforms are functional homologues of these vertebrate proteins and regulate actin filament stability and actomyosin contractility in vivo.

The two UNC-87 isoforms with different N-terminal sequences are expressed in different tissues. Our biochemical analysis shows that UNC-87A, with a longer N-terminus, has more robust actin-bundling and stronger myosin-binding activities than UNC-87B, with a shorter N-terminus, suggesting that these quantitative differences in their activities are important for their tissue-specific functions. UNC-87B is predominantly expressed in the body wall muscle and the somatic gonad, where it regulates actin organization (Goetinck and Waterston, 1994a), actin filament stability (Yamashiro et al., 2007), and contractility (this study). Therefore the relatively weak activity of UNC-87B might be optimal for actin regulation in these striated and nonstriated muscles. By contrast, the in vivo function of UNC-87A is unknown because a specific mutation for unc-87A is not available. The mutations in unc-87(e1216) and unc-87(e1459) are located in exon 2, which is common to both UNC-87 isoforms (Supplemental Figure S1F). Therefore UNC-87A should also be ablated in these mutants. However, we were not able to detect a phenotype in tissues in which expression of UNC-87A was detected (unpublished data). This might be due to a weak phenotype or a phenotype other than actin organization and contractility, which we might have overlooked. Another possibility is that a functionally overlapping actin-binding protein compensates for the function of UNC-87A.

**DISCUSSION**

In this study, we found that two UNC-87 isoforms are expressed in a tissue-specific manner by two separate promoters driving two alternative first exons and that the two UNC-87 isoforms bind to

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**FIGURE 5:** UNC-87 isoforms inhibit actomyosin ATPase. (A–D) ATPase activity of Ce-myosin (0.44 μM) was determined in the absence or presence of 1.0 μM actin and 0.96 μM UNC-87A (A, B) or 3.3 μM UNC-87B (C, D). ATPase activity (A, C) and percentage activation as compared with Ce-myosin alone (B, D). (E, F) ATPase activity of R-myosin (0.35 μM) was determined in the absence or presence of 1.0 μM actin and 0.96 μM UNC-87A (E) or 3.3 μM UNC-87B (F). Data are average ± SD from three independent experiments.
that both UNC-87 isoforms can mediate formation of ATP-resistant functions of UNC-87 isoforms. Our biochemical experiments show absence of sequence similarity. (Rozenblum and Gimona, 2008), and the unique sequence of the CLIK repeats appear to be intrinsically unstructured in solution (Gimona et al., 2001) and in cultured cells (Gimona et al., 2004). An increase in actin-stabilizing functions in vitro (Kranewitter et al., 1994a), resulting in the previously characterized role of UNC-87 isoforms in the myoepithelial sheath. However, the functional difference between tropo- nin–tropomyosin system is required for contractility of myoepithelial sheet (Myers et al., 1996; Ono and Ono, 2004; Obinata et al., 2010). When the two troponin I isoforms, TN-I and UNC-27, are depleted, the myoepithelial sheet becomes excessively contracted (Obinata et al., 2010), which is similar to the phenotype of unc-87 mutants. Thus, in addition to troponin I, which is well established as an inhibitor of actomyosin interaction (Squire and Morris, 1998), UNC-87B is also important for contractile regulation.

UNC-87A and UNC-87B commonly have seven CLIK repeats but are different in their N-terminal sequences. UNC-87A (565 amino acids) has an extra ~190 amino acids with no homology to known functional protein sequences, which is absent in UNC-87B (374 amino acids; Goetinck and Waterston, 1994a). Although this N-terminal sequence of UNC-87A is the most likely responsible for the quantitative differences in the activities of the UNC-87 isoforms, a role of this unique region is unknown. If this region binds to F-actin or myosin, it may cooperate with the C-terminal CLIK repeats to enhance binding of UNC-87A to F-actin or myosin. Alternatively, the N-terminal sequence may allosterically enhance an actin- or myosin-binding function of the C-terminus. Previous functional studies on recombinant proteins containing different numbers of CLIK repeats showed that an increase in the number of CLIK repeats is correlated with an increase in actin-stabilizing functions in vitro (Kranewitter et al., 2001) and in cultured cells (Gimona et al., 2003; Lener et al., 2004). CLIK repeats appear to be intrinsically unstructured in solution (Rozenblum and Gimona, 2008), and the unique sequence of the UNC-87A may have a function similar to CLIK repeats even in the absence of sequence similarity.

Myosin binding and inhibition of actomyosin ATPase are novel functions of UNC-87 isoforms. Our biochemical experiments show that both UNC-87 isoforms can mediate formation of ATP-resistant actomyosin bundles. Therefore UNC-87 proteins can also function not only as a regulator of actomyosin contractility, but also as a stabilizer of actomyosin structural integrity. These activities of UNC-87 isoforms are indeed consistent with the previously reported unc-87 mutant phenotypes. In body wall muscle, sarcomeric actin filaments are disorganized in unc-87 mutants, but this phenotype can be suppressed by a mutation in the unc-54 myosin heavy chain (Goetinck and Waterston, 1994b), suggesting that a loss of the inhibitory function of UNC-87B for actomyosin contractility can be compensated by reduction in the contractile function of myosin itself. Alternatively, UNC-87B may normally stabilize sarcomeres against mechanical stress by actin–myosin cross-linking, and reduced contractility due to the myosin mutation can suppress mechanical damage on sarcomeres in unc-87 mutants.

The myoepithelial sheath of the C. elegans somatic gonad is nonstriated muscle and essential for oocyte maturation and ovulation (Kim et al., 2013). However, the mechanism of regulation of myoepithelial sheet contractility is poorly understood. Unlike vertebrate smooth muscle, the troponin–tropomyosin system is required for contractility of myoepithelial sheet (Myers et al., 1996; Ono and Ono, 2004; Obinata et al., 2010). When the two troponin I isoforms, TN-I and UNC-27, are depleted, the myoepithelial sheath becomes excessively contracted (Obinata et al., 2010), which is similar to the phenotype of unc-87 mutants. Thus, in addition to troponin I, which is well established as an inhibitor of actomyosin interaction (Squire and Morris, 1998), UNC-87B is also important for contractile regulation.
Promoter-reporter analysis
Promoter::GFP constructs for analysis of promoters for UNC-87 isoforms were made using fusion PCR as previously described (Hobert, 2002). For unc-87A, 2038 base pairs of genomic DNA containing 2008 base pairs upstream plus 30 base pairs downstream from the initiation codon of unc-87A was amplified by PCR and fused in-frame with the GFP-coding cassette (GFP plus let-858 3′-untranslated region) from pPD118.20 (kindly provided by A. Fire, Stanford University, Stanford, CA). For unc-87B, 2421 base pairs of genomic DNA containing 2406 base pairs upstream plus 15 base pairs downstream from the initiation codon of unc-87B was amplified by PCR and fused in-frame with the same GFP-coding cassette. The fusion-PCR constructs were injected together with pRF4, a dominant rol-6 transformation marker, into the syncytial region of the gonad of wild-type worms. GFP-positive F1 worms were isolated, and F2 worms that stably inherited the transgenes were selected to establish transgenic lines. Worms were anesthetized in M9 buffer containing 0.1% tricaine and 0.01% tetramisole for 30 min, mounted on 2% agarose pads, and observed by epifluorescence using a Nikon Eclipse TE2000 inverted microscope (Nikon Instruments, Tokyo, Japan).

MATERIALS AND METHODS

Nematode strains
The following strains were obtained from the Caenorhabditis Genetics Center (Minneapolis, MN) and used in this study: wild-type strain N2, CB1216 unc-87(e1216), CB1459 unc-87(e1459), RW1524 unc-87(e1459) unc-54(s95), and RW5008 unc-54(s95). The mutant strains have been published previously (Moerman et al., 1982; Goetinck and Waterston, 1994a,b). All mutants were used as homozygotes. Nematodes were grown under standard conditions at 20°C on Nematode Growth Medium agar plates for phenotypic analysis or at room temperature in liquid Nematode Growth Medium for myosin preparation as described (Stiernagle, 2006).

Proteins
Rabbit muscle actin was prepared from rabbit muscle acetone powder (Pel-Freeze Biologicals, Rogers, AR) as described (Pardee and Spudich, 1982). Rabbit muscle myosin was purified from rabbit back muscle as described (Perry, 1955). C. elegans myosin was purified from liquid-grown N2 strain as described (Tanii et al., 1985). Recombinant UNC-87B was expressed in Escherichia coli and purified as important for contractility of striated and nonstriated muscles. These functions have been proposed for vertebrate proteins, including calponin, caldesmon, and fesselin/synaptopodin-2, mostly from biochemical studies, but have not been clearly demonstrated in vivo. This could be partly due to functional redundancy among functionally similar proteins. The C. elegans genome does not have caldesmon or fesselin/synaptopodin-2, and its relatively simple system allowed us to determine the biological significance of the CLIK-repeat protein. Furthermore, we detected tissue-specific expression of UNC-87 isoforms in both muscle and nonmuscle cells, suggesting that UNC-87A or UNC-87B is involved in actin-dependent events in a variety of cell types. Similarly, calponins and calponin-like proteins in other organisms should also function in a similar manner to UNC-87 isoforms using CLIK repeats as actin-binding motifs.
band intensity was quantified by ImageJ (National Institutes of Health, Bethesda, MD).

Myosin sedimentation assays

Ce-myosin (0.19 μM) was incubated with various concentrations of UNC-87A or UNC-87B in a buffer containing 27 mM KCl, 1 mM MgCl$_2$, 1 mM DTT, and 20 mM imidazole-HCl, pH 7.5, for 30 min at room temperature. The reactions were centrifuged at 18,000 × g for 30 min using a Beckman Microfuge. The supernatants and pellets were adjusted to the same volumes, and analyzed by SDS–PAGE. Quantitative analysis was done in the same manner as described for actin sedimentation assays.

Actin sedimentation assays

F-actin (10 μM) was incubated with various concentrations of UNC-87A or UNC-87B in F-buffer (0.1 M KCl, 2 mM MgCl$_2$, 1 mM dithiothreitol [DTT], 20 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid−KOH, pH 7.5) for 30 min at room temperature. The reactions were centrifuged either at low speed (18,000 rpm for 10 min using a Beckman Microfuge) to examine actin bundling or at high speed (80,000 rpm for 20 min using a Beckman TL-100 ultracentrifuge with a TLA-100 rotor) to examine F-actin binding. The supernatants and pellets were fractionated, adjusted to the same volumes, and analyzed by SDS–PAGE. Quantitative analysis was done in the same manner as described for actin sedimentation assays.

Described (Kranewitter et al., 2001). To produce recombinant UNC-87A, a full-length protein coding sequence for UNC-87A was amplified by PCR from the cDNA clone yk1701a7 (kindly provided by Yuji Kohara, National Institute of Genetics, Mishima, Japan) and cloned into pET-3a (EMD Millipore, Billerica, MA). Recombinant UNC-87A with no additional tag was expressed in E. coli BL21 (DE3) and purified in the same procedure for purification of UNC-87B (Kranewitter et al., 2001). Protein concentrations were determined by the BCA Protein Assay Kit (Thermo Fisher Scientific, Waltham, MA).

![Image](image_url)

**FIGURE 8:** unc-87 mutation induces excessive actomyosin contraction in the myoepithelial sheath of the somatic gonad. Dissected gonads from wild type (A–D), unc-54(s95) (E–H), unc-87(e1459) (I–L), or unc-87(e1459) unc-54(s95) (M–P) were immunostained for MYO-3 to visualize the myoepithelial sheath at a low magnification (left column) or for both actin and MYO-3 to characterize actomyosin organization at a high magnification (right three columns). Merged images of actin (red) and MYO-3 (green) are shown in D, H, L, and P. The proximal region of the gonad is oriented to the right. Quantitative data on the length of the myoepithelial sheath are shown in Table 1. Bars, 10 μm.

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Length (μm)</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild type</td>
<td>150 ± 29</td>
<td>31</td>
</tr>
<tr>
<td>unc-54{s95}</td>
<td>131 ± 18</td>
<td>48</td>
</tr>
<tr>
<td>unc-87(e1459)</td>
<td>74.8 ± 21</td>
<td>35</td>
</tr>
<tr>
<td>unc-87(e1459) unc-54{s95}</td>
<td>125 ± 20</td>
<td>50</td>
</tr>
</tbody>
</table>

**TABLE 1:** Effects of unc-87 and unc-54 mutations on the length of the myoepithelial sheath.
Direct observation of actin bundles by fluorescence microscopy

Rabbit muscle G-actin (8 μM) was copolymerized with DyLight 549-labeled G-actin (2 μM) (Liu et al., 2010) in F-buffer for 2 h at room temperature. The labeled actin filaments (final 2 μM actin) were incubated with myosin (0.19 μM), UNC-87A (1.0 μM), and/or UNC-87B (1.0 μM) in the presence or absence of 5 mM ATP in F-buffer for 10 min at room temperature, mounted on nitrocellulose-coated coverslips, and observed by epifluorescence using a Nikon TE2000 inverted microscope.

Myosin ATPase assays

Myosin ATPase activity was determined by a colorimetric method as described previously (Obinata and Sato, 2012) in a modified buffer containing 27 mM KCl, 1 mM MgCl₂, and 20 mM imidazole-HCl, pH 7.5. Ce-myosin and R-myosin were used at 0.44 and 0.35 μM, respectively. Actin (1.0 μM), UNC-87A (0.96 μM), and/or UNC-87B (3.3 μM) were added in some reactions.

In vitro motility assays

In vitro motility of actomyosin was examined at room temperature as described by Sellers (2001) with modifications. Intact rabbit skeletal myosin muscle was diluted to 0.3 mg/ml in a buffer containing 0.3 M NaCl, 2 mM MgCl₂, and 20 mM Tris-HCl, pH 8.5, and applied to a flow cell that was made with a nitrocellulose-coated glass coverslip (No. 1, 35 × 50 mm), an 18-mm square No. 1 coverslip, and Scotch double-sided tapes as spacers. The cell was sequentially blocked by 1 mg/ml bovine serum albumin (BSA) in wash buffer (27 mM KCl, 5 mM MgCl₂, 0.1 mM ethylene glycol tetraacetic acid, 20 mM imidazole-HCl, pH 7.5) and 5 μM unlabeled F-actin and 1 mM ATP in wash buffer (for blocking nonmotile myosin). DyLight 549–labeled actin filaments (26% labeled) were diluted to 0.4 μM actin in wash buffer and incubated in the flow cell for 2 min. Wash buffer with or without UNC-87A or UNC-87B was infused in the cell and incubated for 2 min. Motility was initiated by adding wash buffer containing 1 mM ATP, 0.7% methylcellulose, and 34 mM DTT. Time-lapse images of fluorescent actin filaments were recorded for 10 s using a Nikon Eclipse TE2000 inverted microscope with a CFI Plan Fluor ELWD 40x (dry; NA 0.60) or Plan Apo 60x (oil; NA 1.40) objective and IPLab imaging software (BD Biosciences) and processed by IPLab imaging software.

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