Six Innexins Contribute to Electrical Coupling of C. elegans Body-Wall Muscle

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Abstract

C. elegans body-wall muscle cells are electrically coupled through gap junctions. Previous studies suggest that UNC-9 is an important, but not the only, innexin mediating the electrical coupling. Here we analyzed junctional current (Ij) for mutants of additional innexins to identify the remaining innexin(s) important to the coupling. The results suggest that a total of six innexins contribute to the coupling, including UNC-9, INX-1, INX-10, INX-11, INX-16, and INX-18. The Ij deficiency in each mutant was rescued completely by expressing the corresponding wild-type innexin specifically in muscle, suggesting that the innexins function cell-autonomously. Comparisons of Ij between various single, double, and triple mutants suggest that the six innexins probably form two distinct populations of gap junctions with one population consisting of UNC-9 and INX-18 and the other consisting of the remaining four innexins. Consistent with their roles in muscle electrical coupling, five of the six innexins showed punctate localization at muscle intercellular junctions when expressed as GFP- or epitope-tagged proteins, and muscle expression was detected for four of them when assessed by expressing GFP under the control of innexin promoters. The results may serve as a solid foundation for further explorations of structural and functional properties of gap junctions in C. elegans body-wall muscle.

Introduction

Gap junctions are intercellular channels connecting the cytoplasm of adjacent cells. They are formed by connexins in mammals but innexins in invertebrates. Innexins are homologues of mammalian pannexins [1-3], which may form hemichannels [1,4-11]. Deficiencies and mutations in gap junction proteins cause a variety of human disorders such as demyelination [12,13], cataracts [14,15], deafness [16,17], and heart malformations [18].

Each species generally has multiple genes encoding gap junction proteins. For example, the human genome has 21 connexins and 3 pannexins. The Caenorhabditis elegans and Drosophila melanogaster genomes have 25 and 8 innexins, respectively [19,20]. Different gap junction proteins are often expressed in the same cells, and have the potential to coassemble into functional units, which greatly increases the structural and functional diversity of gap junctions. Analyses using transfected mammalian cell lines and the Xenopus oocyte heterologous expression system have shown that many gap junction proteins may coassemble to form heterotypic or heteromeric gap junctions [21]. There is also evidence that heterotypic or heteromeric gap junctions exist in native tissues [21-25]. However, molecular identification and functional attributions for all gap junction proteins expressed in individual cell types have been hampered by a scarcity of specific antibodies and a lack of specific pharmacological inhibitors.

C. elegans is a particularly facile model system for understanding the structure and function of gap junctions for several major reasons, including their small number of somatic cells (959 cells) per animal, existence of mutants for all innexin genes, ease to assess protein expression and subcellular localization patterns, and availability of electrophysiological, cell biological and behavioral assays for gap junction functions. Studies with C. elegans have uncovered previously unknown functions of gap junctions, such as instructing defecation motor steps by propagating Ca²⁺ waves between enteric muscle cells [26], regulating asymmetric gene expression in selected
neurons [27], facilitating synchronous pharyngeal muscle contractions [28,29], promoting sperm guidance to the fertilization site [30], synchronizing body-wall muscle action potentials and Ca$^{2+}$ transients [31], and facilitating spicule thrusts by transmitting signals among male-specific sex muscles [32].

Body-wall muscle occupies a unique niche in gap junction research with C. elegans because it is the only tissue for which the dual voltage- and current-clamp techniques have been adapted to analyze physiological and biophysical properties of gap junctions. Previous studies have shown that muscle cells are electrically coupled in a highly organized pattern through low-conductance gap junctions, and that the innexin UNC-9 serves as an important component of the muscle gap junctions [31,33,34]. However, other innexin(s) must be contributing to the muscle coupling because I$_j$ between adjoining muscle cells within the same quadrant is decreased but not absent in the unc-9 null mutant [33]. To fully understand the function and molecular compositions of the muscle gap junctions, it is important to identify all the innexins functioning in the muscle. In the present study, we identified a total of six innexins contributing to electrical coupling of the muscle through electrophysiological analyses of innexin mutants. The identification of so many gap junction proteins functioning in a single type of cells is unprecedented.

Results

Six innexins are required for normal electrical coupling of body-wall muscle cells

Results of our previous study suggest that UNC-9 is not the only innexin forming gap junctions in C. elegans body-wall muscle [33]. To identify the remaining innexin(s) functioning in the muscle, we analyzed I$_j$ in additional innexin mutants. Although mutants have been isolated for all the 25 innexins of C. elegans, homozygous mutants of 3 innexins (inx-3, inx-12, inx-13) are unsuitable for electrophysiological analyses because they are either lethal or sterile. We therefore focused on mutants of the remaining 22 innexins, including inx-1(tm3524), inx-2(ok376), inx-4(che-7(ok2373), inx-5(ok1053), inx-6(tir5), inx-7(tm2378), inx-8(gk42), inx-9(ok1502), inx-10(ok2714), inx-11(ok2783), inx-14(ag17), inx-15(tm3394), inx-16(tm1589), inx-17(tm3839), inx-18(ok2454), inx-19(nsy-5(tm1896), inx-20(ok426), inx-21(ok2524), inx-22(tm1661), eat-5(ad464), unc-7(e5), and unc-9(fc16). The unc-9 mutant was also included for the purpose of comparisons. Molecular lesions of these mutants can be found at the Wormbase website (www.wormbase.org. Accessed 2013 July 1).

We chose the R1/R2 and L1/L2 pairs of ventral body-wall muscle cells (Figure 1A) for analyses because their relatively strong electrical coupling made it easier to detect mutant effects. Data from the two cell pairs were combined in subsequent statistical analyses because properties of electrical coupling are indistinguishable between these two cell pairs [33]. Mutants of 16 innexins were similar to wild type in junctional conductance (Gj) (Figure 1B), suggesting that these innexins are not important for electrical coupling. In contrast, mutants of 6 innexins, including unc-9, inx-1, inx-10, inx-11, inx-16 and inx-18, displayed a significant decrease in Gj compared with wild type (Figure 1C), suggesting that these innexins may be required for normal electrical coupling of the muscle. The observations with the unc-7 and unc-9 mutants were consistent with those of our previous studies [31,33,34]. Among the six mutants showing deficient muscle electrical coupling, unc-9(fc16) is due to a premature stop at the intracellular loop between the second and third membrane spanning domains [35] whereas the remaining mutants have either deletion or deletion plus insertion in an innexin gene (www.wormbase.org. Accessed 2013 July 1). We refer to the six mutants as if (loss-of-function) mutants in the remainder of the text.

Innexins function cell-autonomously to contribute to muscle electrical coupling

The coupling defects described above likely resulted from a deficiency of the innexins in body-wall muscle. Indeed, this has been confirmed for unc-9(If) [33,34]. For mutants of the other five innexins, however, the observed muscle coupling defects could also result from other potential causes, such as an indirect effect from non-muscle cells, or mutation of another gene. To determine whether the coupling defect resulted from innexin deficiency in body-wall muscle, we performed muscle-specific rescue experiments for each mutant by expressing the corresponding wild-type innexin under the control of the muscle-specific myo-3 promoter (Pmyo-3) [36]. In all cases, Gj was restored to wild-type level in the mutant (Figure 1C), suggesting that all the six innexins function cell-autonomously to contribute to the electrical coupling. Intriguingly, none of the rescued strains showed a significantly higher Gj than wild type, suggesting that overexpression of a single innexin in body-wall muscle is not enough to cause a significant increase in functional gap junctions.

Innexins may function as two distinct populations of gap junctions

The six innexins identified through the electrophysiological analyses unlikely form independent populations of homotypic gap junctions because the combined decrease of Gj of these mutants was far greater than 100%. To assess how many populations of gap junctions that these innexins might form in body-wall muscle, we compared Gj among various single and double mutants of the innexins. If two innexins functioned together, Gj in the double mutant would be comparable to that in either single mutant. Otherwise, Gj would be decreased to a larger degree in the double mutant than in either single mutant. We first compared Gj between single and double mutants of unc-9(If) and inx-18(If) because coupling deficiencies were observed in these two mutants well before we embarked on the project to systematically analyze essentially all innexin mutants. We found that, while unc-9(If) caused a larger decrease in Gj than inx-18(If), Gj in the double mutant was similar to that in the unc-9(If) single mutant (Figure 2A), suggesting that UNC-9 and INX-18 likely contribute to a single population of gap junctions. The lesser coupling deficiency in inx-18(If) than unc-9(If) might be because UNC-9 but not
Figure 1. Junctional conductance (G) of *C. elegans* body-wall muscle was significantly decreased in mutants of six innexins. A. Diagram showing ventral body-wall muscle cells. Ventral muscles include the left and right quadrants with each quadrant consisting of two rows of muscle cells in a monolayer. The highlighted cell pairs (L1/L2 and R1/R2) represent those used for electrophysiological analyses. B. $G_j$ was indistinguishable between wild type (WT) and mutants of 16 innexins. C. Mutants of 6 innexins showed significantly lower $G_j$ when compared with WT, and the coupling defect was rescued completely by expressing a wild-type innexin in each corresponding mutant. The asterisk (*) indicates a statistically significant difference compared with WT. The number on each bar represents the number ($N$) of cell pairs analyzed.

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INX-16 may form partially functional gap junctions in the absence of its counterpart.

We next compared \( G_i \) between single and double mutants for two other pairs of innexins, including \( inx-1/INX-10 \) and \( inx-11/INX-16 \) because \( G_i \) was essentially identical between \( inx-1\) (I) and \( inx-10\) (I), and between \( inx-11\) (I) and \( inx-16\) (I). We found that \( G_i \) was indistinguishable between \( inx-1\) (I);\( inx-10\) (I) double mutant and either \( inx-1\) (I) or \( inx-10\) (I) single mutant, and also between the \( inx-1\) (I);\( inx-16\) (I) double mutant and either \( inx-11\) (I) or \( inx-16\) (I) single mutant (Figure 2B). These observations suggest that \( INX-1 \) and \( INX-10 \) likely function together, and so do \( INX-11 \) and \( INX-16 \). The \( G_i \) was significantly lower in the single or double mutants of \( inx-11 \) and \( inx-16 \) than those of \( inx-1 \) and \( inx-10 \) (Figure 2B). To determine whether there is any functional relationship between these two pairs of innexins, we compared \( G_i \) between the double mutant \( inx-10\) (I);\( inx-11\) (I) and either single or double mutants of \( inx-1\) (I) and \( inx-16\) (I), and found that \( G_i \) was not further decreased in the \( inx-10\) (I);\( inx-11\) (I) double mutant (Figure 2B). These analyses collectively suggest that four innexins, including \( INX-1, \) \( INX-10, \) \( INX-11 \) and \( INX-16 \), likely contribute to the function of a single population of gap junctions. The lesser coupling deficiency in \( inx-1\) (I) and \( inx-10\) (I) than \( inx-11\) (I) and \( inx-16\) (I) might be because \( INX-11 \) and \( INX-16 \) but not \( INX-1 \) and \( INX-10 \) may form partially functional gap junctions in the absence of their counterparts.

We then determined whether the two populations of gap junctions described above actually belong to a single population by comparing \( G_i \) between single, double, and triple mutants of selected innexins, including \( unc-9, inx-9, inx-10, inx-11, \) and \( inx-16 \). Combining either \( inx-1\) (I), \( inx-16\) (I), or both with \( unc-9\) (I) resulted in a \( G_i \) much smaller than that in any of the single mutants (Figure 2C). Indeed, \( I_j \) was essentially indistinguishable from baseline noises in the double and triple mutants. \( G_i \) was also further decreased when combining either \( inx-1\) (I) or \( inx-10\) (I) with \( unc-9\) (I) (Figure 2C). These observations suggest that there are likely two distinct populations of gap junctions in body-wall muscle, with one population consisting of \( UNC-9 \) and \( INX-18 \) and the other consisting of \( INX-1, \) \( INX-10, \) \( INX-11 \) and \( INX-16 \).

**GFP transcriptional fusions show muscle expression of several innexins**

The electrophysiological data suggest that six innexins are expressed in body-wall muscle. To assess muscle innexin expression using an independent approach, we expressed GFP as innexin promoter::GFP transcriptional fusions and analyzed GFP signal in live worms. We first used a homologous recombination approach for all the 25 innexins to detect potential muscle expression as part of an effort to provide a high resolution map for all the 25 innexins [37]. Specifically, a plasmid containing a short promoter sequence (typically ~1 kb) fused to GFP coding sequence was made for each innexin and coinfected with a cosmids containing the corresponding innexin gene. It was expected that homologous recombination in vivo would result in a promoter::GFP transcriptional fusion containing the entire innexin promoter [38,39]. Body-wall muscle expression of several innexins, including \( unc-9, inx-8, inx-10, inx-11, inx-14, inx-18, inx-19 \) and \( inx-21 \), was detected using this approach. The muscle expression of \( inx-10, inx-11, inx-18 \) and \( unc-9 \) is shown in Figure 3. The muscle expression of the other innexins is not shown because those innexins did not appear to play a role in electrical coupling.

The experiments described above failed to confirm body-wall muscle expression for \( inx-1 \) and \( inx-16 \). One possible cause was that a full-length innexin promoter was not assembled through homologous recombination in vivo due to unexpected deletions in the cosmids or other factors. To address this possibility, we made promoter::GFP transcriptional fusions for both innexins by fusing a longer promoter sequence to GFP, and examined GFP expression patterns in transgenic worms. The length of promoter sequence used was 4925 bp for \( inx-1 \) and 1931 bp for \( inx-16 \). The length of the \( inx-16 \) promoter sequence used was relatively short because the translation initiation site of \( inx-16 \) is very close to the coding region of another upstream gene (~1.0 kb). However, GFP expression was still not observed in body-wall muscle cells (not shown).

**Innexins are localized to intercellular junctions**

Gap junctions typically appear as plaques/puncta at the plasma membranes between adjacent cells, which has been shown for \( UNC-9 \) by immunostaining using an \( UNC-9\)-specific antibody and by expressing GFP-tagged full-length \( UNC-9 \) [33,34]. We therefore analyzed subcellular localization for the remaining five innexins identified by the electrophysiological analyses. We first expressed GFP-tagged full-length innexins in body-wall muscle under the control of \( Pmyo-3 \) and analyzed subcellular localization patterns of the fusion proteins. With respect to \( INX-1 \), two different splice isoforms (\( INX-1a \) and \( INX-1c/GenBank KF137642 \)) were analyzed. GFP fluorescent puncta were observed at muscle intercellular junctions with \( INX-1c::GFP, INX-10::GFP \) and \( INX-16::GFP \) although the size and density of the fluorescent puncta varied among them and \( INX-16::GFP \) mostly appeared as intracellular aggregates (Figure 4, top). However, \( INX-1a::GFP \) showed diffuse muscle expression; and \( INX-11::GFP \) and \( INX-18::GFP \) were not localized to intercellular junctions but mainly appeared as large intracellular aggregates (not shown). We reasoned that the large intracellular aggregates might reflect a GFP-induced trafficking defect, and that a smaller epitope tag might have a lesser problem of this sort. Therefore, we expressed Myc-tagged \( INX-11 \) (Myc::\( INX-11 \)) and HA-tagged \( INX-18 \) (\( INX-18::HA \)) in muscle cells under the control of \( Pmyo-3 \) and analyzed subcellular localization of the tagged proteins by immunostaining using either a Myc- or a HA-specific antibody. \( INX-18::HA \) displayed punctate localization at muscle intercellular junctions (Figure 4, top). Myc::\( INX-11 \) displayed punctate localization at muscle intercellular junctions and inside muscle cells (likely corresponding to dense bodies) although size and density of the puncta at intercellular junctions appeared to be distinct from those of the other innexins (Figure 4, top). An image of \( UNC-9::GFP \) in body-wall muscle is also shown for the purpose of comparison.

We previously showed that \( UNC-9::GFP \) fusion protein is functional [34]. To determine whether the fusion proteins for the
Figure 2. Innexins likely assemble into two distinct populations of gap junctions in C. elegans body-wall muscle. A. Junctional conductance ($G_j$) was not further decreased in the unc-9;inx-18 double mutant compared with the unc-9 single mutant, suggesting that UNC-9 and INX-18 function together. B. $G_j$ was decreased to a similar degree in the inx-1;inx-10 and inx-11;inx-16 double mutants compared with their corresponding single mutants although $G_j$ was significantly lower in the mutants of inx-11 and inx-16 than those of inx-1 and inx-10. $G_j$ was not further decreased in the inx-10;inx-11 double mutant, suggesting that the four innexins (INX-1, INX-10, INX-11 and INX-16) likely function together. C. Combining selected mutants from A and B resulted in a further decrease of $G_j$, suggesting that there are two distinct populations of gap junctions in muscle. In A, the asterisk (*) indicates a statistically significant difference compared with the inx-18 single mutant. In both B and C, the asterisk indicates a statistically significant difference compared with the single and double mutants of inx-1 and inx-10, the pound sign (#) indicates a statistically significant difference compared with the single and double mutants of inx-11 and inx-16, and the triangle (Δ) indicates a statistically significant difference compared with inx-1;unc-9 and inx-10;unc-9. The number on each bar represents the number (N) of cell pairs analyzed.

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remaining 5 innexins are also functional, we expressed them in body-wall muscle of corresponding mutants under the control of Pmyo-3 and analyzed their effects on electrical coupling. The coupling deficiencies of 4 innexin mutants (inx-1, inx-10, inx-11 and inx-18) were rescued whereas that of inx-16 was not (Figure 4, bottom). The failure for INX-16::GFP to rescue muscle coupling might result from insufficient membrane targeting/localization and/or a nonfunctional fusion protein. Interestingly, worms expressing the fusion proteins tended to display larger Gj compared with wild type, which was different from the observations with wild-type innexins but is reminiscent of an earlier observation with UNC-9::GFP [34].

Discussion

The results of the present study suggest that six innexins likely assemble into two populations of gap junctions in C. elegans body-wall muscle. This conclusion is based on comparisons of Gj among wild type and various single and double mutants of the innexins. The fact that overexpressing any one of the six innexins in its wild form in body-wall muscle of a corresponding mutant did not result in a significantly higher Gj compared with wild type worms favors the notion that gap junctions in body-wall muscle normally consist of two or more different innexins.

The identified innexins are expected to show expression in body-wall muscle and punctate subcellular localization at muscle intercellular junctions. Indeed, muscle expression was detected for four of the six innexins when GFP was used as a reporter of innexin promoter activity, and punctate subcellular localization at muscle intercellular junctions was observed for five of the six innexins when they were expressed as either GFP- or epitope-tagged proteins. However, muscle expression was not detected for two of the six innexins; the density and size of the intercellular puncta varied considerably among the innexins when they were analyzed as fusion proteins. The failure to detect muscle expression for an innexin could be due to the absence of a regulatory element in the promoter::GFP translational fusion, such as introns [40] and distal regulatory elements [41,42]. It could also be due to weak GFP expression or GFP fusion to the initiation site of a splice isoform that is not normally expressed in muscle. The apparent variability in subcellular localization could be due to interference by GFP or the epitope tag, or the use of an innexin splice isoform that is not normally expressed in body-wall muscle. Innexin genes appear to have extensive alternative splicing because previously unknown splice variants were frequently identified during our processes of isolating innexin cDNAs by RT-PCR using primer pairs based on only one previously reported isoform for each innexin (examples described in Methods). It is very likely that some splice variants were missed, especially those with alternative 5'- and 3'-ends. Therefore, the negative expression data and variable subcellular localization patterns do not challenge our main conclusion. Convincing expression and subcellular localization results may come from immunostaining with antibodies against a common region of all splice isoforms for each innexin. However, such an antibody is currently available only for UNC-9 [34].

Two or more gap junction proteins could coassemble into homomeric/heterotypic, heteromeric/heterotypic, or heteromeric/homotypic gap junctions [21]. It is impossible to predict exact stoichiometries of the muscle gap junctions because the number of possible composition stoichiometries for a gap junction formed by only two innexins is as many as
Figure 4. Subcellular localization of innexins in *C. elegans* body-wall muscle. **Top**, Subcellular localization patterns of innexin fusion proteins in body-wall muscle. INX-1::GFP, INX-10::GFP, Myc::INX-11, and INX-18::HA mainly showed punctate localization at muscle intercellular junctions. INX-16::GFP showed some punctate localization between muscle cells but mainly intracellular aggregates. **Bottom**, Effects of the fusion proteins (expressed in corresponding mutants) on body-wall muscle electrical coupling. The asterisk (*) indicates a statistically significant difference compared with the corresponding mutant (paired t-test). The numbers inside the columns represent sample size (n). The wild-type (WT) and mutant data in Figure 1C were included for comparisons.

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196 [21]. Nevertheless, we have ventured to put together a few models based on several assumptions (Figure 5). First, each muscle cell expresses all six innexins. Second, a heteromeric hemichannel of two different innexins is more stable in structure than that of three or four different innexins because the two different innexins may each contribute three subunits and arrange in an alternate order to form a 3-fold symmetric structure. Third, two innexins that have similar effects on G_j when mutated likely exist in separate hemichannels of a gap junction but are docked to each other; and a mutation of either of them would prevent the other from contributing to functional gap junctions. These models are postulated not to imply that they represent the actual stoichiometries but to show that the six innexins could be accommodated as components of two populations of gap junctions. The models have obvious limitations or caveats. For example, a few rather than all of the six innexins might be expressed in each muscle cell. There might be other innexins contributing to muscle electrical coupling because mutants of three innexins were not analyzed. Electrical coupling between other muscle cell pairs might be mediated by gap junctions of different molecular compositions because the electrophysiological analyses were restricted to selected ventral muscle cells between the neck and vulval region. Further studies may help define how various innexins assemble stoichiometrically into gap junctions in muscle.

The identification of six innexins functioning in a single cell type was unexpected and unprecedented, although such a possibility was suggested by our earlier innexin expression map [37]. It is hard to imagine that this is a phenomenon unique to C. elegans body-wall muscle. The findings of this study suggest that molecular compositions and stoichiometries of gap junctions in a specific cell type could be highly complex. Identification of all gap junction proteins expressed in a particular cell type and determination of the molecular stoichiometries of the gap junctions should be an essential step for fully understanding how gap junctions contribute to various physiological functions.

Methods

Growth and Culture of C. elegans

C. elegans hermaphrodites were grown on nematode growth medium agar plates with a layer of OP50 Escherichia coli inside an environmental chamber (21 °C).

Electrophysiology

An adult hermaphrodite was immobilized on a glass coverslip by applying Vetbond™ Tissue Adhesive (3M Company, MN, USA). Application of the adhesive was generally restricted to the dorsal middle portion of the animal, allowing the head and tail to sway during the experiment. A longitudinal incision was made in the dorsolateral region. After clearing the viscera, the cuticle flap was folded back and glued to the coverslip, exposing the ventral nerve cord and the two adjacent muscle quadrants. A Nikon FN1 upright microscope equipped with a 40X water-immersion objective and 15X eyepieces was used for viewing the preparation. Borosilicate glass pipettes with a tip resistance of 3–5 MΩ were used as electrodes for voltage-clamp recordings in the classical whole-cell patch clamp configuration with a Multiclamp 700B amplifier (Molecular Devices, Sunnyvale, CA) and Clampex software (version 10, Molecular Devices). Series resistance was compensated to 70-80%. To record I_j, the membrane potential (V_m) of both cells was held at -30 mV, from which a series of voltage steps (-110 mV to +50 mV at 10 mV intervals, 250 ms duration) were applied to one cell (Cell 1) whereas the other cell (Cell 2) was held constant to record I_j. The voltage steps were repeated four times and automatically averaged for subsequent analyses. I_j was sampled at a rate of 10 kHz after filtering at 2 kHz. Junctional voltage (V_j) was defined as V_m of Cell 2 minus V_m of Cell 1.

The extracellular solution contained (in mM) NaCl 140, KCl 5, CaCl_2 5, MgCl_2 5, dextrose 11, and HEPES 5 (pH 7.2). The pipette solution contained (in mM) KCl 120, KOH 20, Tris 5, CaCl_2 0.25, MgCl_2 4, sucrose 36, EGTA 5, and Na_2ATP 4 (pH 7.2).

cDNA cloning

cDNAs of the innexins were cloned through RT-PCR using primer pairs complementary to the 5’- and 3’-ends of one
isoform (the longest one if more than one had been reported) for each innexin listed at the Wormbase website (www.wormbase.org. Accessed 2013 July 1). The specific cDNA isoforms used to guide the primer design were C16E9.4a for inx-1, T18H9.5a for inx-10, W04D2.3b for inx-11, R12E2.5 for inx-16, and C18H7.2a for inx-18. A cDNA of unc-9 was cloned in an earlier study [33]. Two isoforms of cDNA were cloned for inx-10, including T18H9.5a and two novel isoforms (GenBank KF137643 and KF137644). Two novel isoforms of cDNA were cloned for inx-11 (GenBank KF137645 and KF137646). The cloning of KF137646 was aided by using primers complementary to a novel exon identified through comparing genomic DNA sequences between C. elegans and C. briggsae, as we did for other genes [43]. The cDNAs cloned for inx-16 and inx-18 are identical to R12E2.5 and C18H7.2a, respectively.

Mutant Rescue

Muscle-specific rescue was performed by injecting a plasmid containing Pmyo-3 fused to the initiation site of a wild-type innexin cDNA into a corresponding innexin mutant using a standard technique [44]. A Pmyo-3::GFP plasmid (pPD118.20) was coinjected to serve as a transformation marker, which caused body-wall muscle cells to display bright green fluorescence. The specific cDNA isoforms used in these experiments were C16E9.4a for inx-1 (plasmid wp1217), T18H9.5a for inx-10 (plasmid p1219), W04D2.3c for inx-11 (plasmid wp1221), R12E2.5 for inx-16 (plasmid wp1222) and C18H7.2a for inx-18 (plasmid p724). The muscle-specific rescue strain for unc-9 was from an earlier study [33]. The plasmids used for testing the ability of fusion proteins to rescue muscle coupling was wp1227 for INX-1::GFP, wp1224 for INX-10::GFP, wp441 for Myc::INX-11, wp1235 for INX-16::GFP, and wp728 for INX-18::HA. Ij was recorded from pairs of adjacent cells showing GFP fluorescence.

Analyses of Expression Pattern

 Procedures for assessing innexin expression patterns using the homologous recombination approach are described previously [37]. To create the new promoter::GFP transcriptional fusion plasmids for inx-1 (wp1264) and inx-16 (wp1266), a promoter sequence before the translation initiation site of either gene was amplified by PCR with genomic DNA as a template, and used to replace Pmyo-3 in pPD118.20 by cutting with Pst1 and BamH1 restriction enzymes. The sequences of the PCR primers were TGTCTCTGTTGACGATT (sense) and attggatccGCGGACACAATTCTGCAAT (antisense) for Pinx-1, and atactcagATGCTGCTGTAATACCC (sense) and attggatccTTTGTTGTTACCTGAAAAT (antisense) for Pinx-16. The sense primer for Pinx-1 did not contain a Pst1 restriction site because an internal Pst1 site in the promoter was used for the cloning. The promoter::GFP transcriptional fusions were injected individually into lin-15(n765) [45]. A lin-15 rescue plasmid was coinjected to serve as a transformation marker. GFP fluorescence signal was imaged using a Nikon inverted microscope (TE2000-U) with a 40X objective, a FITC filter (41001, Chroma Technology Corp., Bellows Falls, VT, USA) and a Peltier cooled digital camera (F-view II; Olympus).

Analyses of Subcellular Localization

GFP- or epitope (Myc or HA)-tagged full-length innexins were expressed in body-wall muscle under the control of Pmyo-3. The specific cDNA isoforms used in these experiments were C16E9.4a and a novel cDNA (GenBank KF137642) for INX-1::GFP (plasmids wp1218 and wp1227), T18H9.5a for INX-10::GFP (plasmid wp1224), R12E2.5 for INX-16::GFP (plasmid wp1235) and C18H7.2a for INX-18::HA (plasmid wp728). Myc::INX-11 was made by fusing a Myc sequence to the initiation site of W04D2.3a in genomic DNA (plasmid wp441). The Pmyo-3::UNC-9::GFP strain was from an earlier study [33]. INX-11 was tagged on the amino terminal whereas the remaining innexins were tagged at the carboxyl terminal. These plasmids were injected separately into lin-15(n765) [45]. A lin-15 rescue plasmid was co-injected to serve as a transformation marker. Transgenic worms expressing a GFP-tagged innexin were imaged after immobilization whereas those expressing a Myc- or HA-tagged innexin were imaged after permeabilization with a Bouin’s fixative [46] and immunostaining first with a mouse anti-HA or a rabbit anti-Myc antibody (Invitrogen Corporation, Carlsbad, CA) and then with an Alexa Fluor® 488-conjugated goat anti-mouse or an Alexa Fluor® 594-conjugated goat anti-rabbit secondary antibody (Invitrogen Corporation). The fluorescence signals were imaged using the Nikon TE2000-U inverted microscope with a 40X or 60X objective, FITC (41001, Chroma Technology Corp.) and Texas Red (41004, Chroma Technology Corp.) filters, and the F-view II camera.

Statistical Analyses

The apparent mean steady-state current during the last 100 ms of each voltage step was determined with Clampfit (Molecular Devices) and used to plot Ij - Vj relationship. Gj was determined from the slope of the Ij - Vj relationship at the linear portion (Vj = –30 to +30 mV) and compared among the different groups by one-way analysis of variance (ANOVA) followed by Bonferroni post hoc tests for pair-wise comparisons. p < 0.05 is considered to be statistically significant. All values are shown as mean ± se. Sample traces of Ij in figures are displayed at 20-mV intervals of Vj steps after filtering at 300 Hz for clarity. Statistical analyses and data graphing were performed using Origin (version 9.0, OriginLab Corporation, Northampton, MA).

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Author Contributions

Conceived and designed the experiments: PL ZW. Performed the experiments: PL BC ZA DH. Analyzed the data: PL BC ZA DH. Wrote the manuscript: ZW PL.
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