Abstract

The stereotyped striation of myofibrils is a conserved feature of muscle organization that is critical to its function. Although most components that constitute the basic myofibrils are well-characterized biochemically and are conserved across the animal kingdom, the mechanisms leading to the precise assembly of sarcomeres, the basic units of myofibrils, are poorly understood. To gain insights into this process, we investigated the functional relationships of sarcomeric protein complexes. Specifically, we systematically analyzed, using either RNAi in primary muscle cells or available genetic mutations, the organization of myofibrils in Drosophila muscles that lack one or more sarcomeric proteins. Our study reveals that the thin and thick filaments are mutually dependent on each other for striation. Further, the tension sensor complex comprised of zipper/Zasp/α-actinin is involved in stabilizing the sarcomere but not in its initial formation. Finally, integrins appear essential for the interdigitation of thin and thick filaments that occurs prior to striation. Thus, sarcomere formation occurs by the coordinated assembly of multiple latent protein complexes, as opposed to sequential assembly.

Introduction

Muscle functionality relies on the correct assembly of myofibrils, the cylindrical organelles attached to the cell surface membrane within muscle cells that run from one end of the cell to the other end. Myofibrils are composed of tandem arrays of basic functional contractile units called the sarcomeres. Sarcomeres are highly ordered, almost crystalline-like, structures composed of thin (actin) and thick (myosin) filaments and their associated proteins (Figure 1A). Although their components have been known for many years, how the various sarcomeric proteins assemble to form these highly ordered structures is poorly understood. Understanding the process of sarcomere assembly is not only relevant to the understanding of how protein complexes interact to form complex supra-molecular structures, but is also of great significance to medicine, as many mutations in genes encoding sarcomeric proteins cause muscle diseases such as congenital myopathy and dilated cardiac hypertrophy [1,2].

The striated appearance of myofibrils is readily detectable under the polarized light microscope as alternating light and dark bands corresponding to I-bands and A-bands, respectively (Figure 1A). Thin filaments are built of actins as well as their associated tropomyosin (Tm) and troponin (Tn) complex (troponin T/TnT, troponin I/TnI and troponin C/TnC) proteins, and are anchored at the Z-line that demarcates the middle of the I-bands. The Z-line itself contains various structural proteins such as α-actinin, titin and Zasp. Thin filaments and Z lines are connected with each other and are often referred to as the “I-Z-I” complex. Thick filaments on the other hand are attached to the M-line situated at the center of A-bands and are composed of muscle myosin represented by two myosin heavy chains (MHCs) and four muscle light chains (MLCs).

A number of models have been put forward to explain how sarcomeric proteins are assembled into a highly ordered structure. One model proposes that I-Z-I complexes and bipolar myosin filaments assemble independently before joining [3]. The main observations in support of this view are that free-floating A band structures can be detected in the absence of actin, and that irregular Z-lines can be detected with attached thin filaments in muscles that lack myosin [3–5]. A second model, supported by the antibody staining of chicken cardiomycocytes fixed at different times after spreading in culture [6], proposes that premyofibrils, characterized by interdigitating barred patterns of I-Z-I complexes and non-muscle myosins, form de novo near the cell membrane and develop into mature myofibrils. This model proposes that a replacement of the non-muscle myosin filaments with muscle myosin filaments occurs during the transition from premyofibril to myofibril and that discrete aggregates of Z bodies along premyofibrils grow and fuse laterally into Z lines [6,7]. A third model emphasizes the scaffolding role of the giant protein titin during myofibrillogenesis. According to this model, the N-terminal domain of titin interacts with the I-Z-I complex and the subsequent unfolded C-terminal region associates with the M-line, thus driving the interdigitation of the I-Z-I complex and myosin. In
Author Summary

Muscle functionality relies on the correct assembly of myofibrils, which are composed of tandem arrays of basic functional contractile units called the sarcomeres. Many mutations in genes encoding sarcomeric proteins cause muscle diseases such as congenital myopathy and dilated cardiac hypertrophy. Understanding the process of sarcomere assembly is not only relevant to the understanding of how protein complexes interact to form complex supramolecular structures, but also of great significance to medicine for muscle diseases. Here, by taking advantage of our newly developed primary muscle cell culture method, we reevaluate sarcomere assembly by systematically analyzing the functional relationship of sarcomeric proteins using RNA interference or genetic ablation techniques. Our analysis leads us to propose a “two-state” model whereby sarcomeric proteins exist either in the “chaotic” state with independently assembled differential functional complexes or the “highly ordered suprastructure” state made from these complexes. Because we fail to detect any previously hypothesized sarcomere assembly intermediates in our system, our data support that sarcomere assembly is a highly coordinated process mediated by multiple latent protein complexes and does not occur in a step-wise fashion. Support of this model, using titin specific epitopes that distinguish between the N or C region, the M-band periodic pattern of titin is detected later than its Z-line pattern [8,9]. Finally, a fourth model emphasizes the role of integrin adhesion complex in sarcomere assembly as starting sites of actin/thin filaments polymerization [10]. This model is supported by genetic studies in C. elegans, Drosophila and mice showing that integrin signaling pathway components are required for sarcomere assembly [11–15].

Although not necessarily mutually exclusive, these models are based on the analysis of sarcomeric structures in either wild-type or mutant backgrounds using electron microscopy and immunohistochemistry approaches, and have been limited by the lack of a number of antibodies against key sarcomeric proteins. Thus, we decided to reexamine this process using new reagents and approaches to systematically investigate the myofilbr assembly process. In particular, we used RNAi in a primary muscle cell culture assay to systematically explore the functional relationships among sarcomeric proteins, combined with detailed in vivo analysis of available muscle mutants. Previously, we have shown that primary muscle cells can be used to study sarcomere organization using RNAi [16,17]. For example, we identified from an RNAi screen in primary muscle cells the sarcomere length short (salts) gene and showed that it is required for thin filament lengthening both in culture and in vivo [18].

Here, using antibody staining of sarcomeric proteins in mutant animals and cultured primary muscle cells, we document how protein complexes interact to form complex suprastructures, but are not required for actin/thin filament assembly as previously proposed. Altogether, our studies indicate that there are no intermediate steps for sarcomere assembly and that disruption of any protein complex leads to loss of muscle striation, thus leading to the proposal that sarcomere formation occurs by the coordinated assembly of multiple latent protein complexes.

Results/Discussion

Roles of MHC and non-muscle myosin in sarcomere formation

Previous studies have established that the first sarcomeric components to appear are the actin filaments that assemble into I-Z-I complexes [8,19,20]. This step is followed by the infiltration of muscle myosin into nascent myofilbrils, a process that is not well understood and that may be mediated by non-muscle myosin [3,6,8,21].

To clarify the process by which I-Z-I and the myosin filaments assemble, we analyzed the striation pattern of sarcomeres in Mhc1 mutants that lack thick filaments. Mhc1 contains a 101 base-pair (bp) deletion in the Mhc gene resulting in truncated MHC proteins lacking most of rod-like sub-fragment 2 that contributes to the backbone of the thick filament. Previous electron microscopic studies have shown that mutant muscles for the Mhc1 amorphic allele completely lack any discernable thick filaments [22]. Wild-type and Mhc1 mutant embryos were dissected at stage 17, when sarcomeres begin to assemble and can be detected in the wild-type body wall muscles [23] (see Methods). Muscles were then stained with various antibodies against sarcomeric proteins. While wild-type muscles displayed well-defined striated myofilbrils when visualized with phallolidin staining and anti-α-actinin antibodies, highlighting actin filaments and Z-lines respectively (upper panels in Figure 1B), homozygous Mhc1 mutant muscles no longer showed localized Z-lines (bottom panels in Figure 1B). This observation is consistent with the phenotypes of Mhc deficient primary muscles that are derived either from wild-type cells treated with Mhc dsRNAs or from cells dissociated from Mhc1 mutant embryos (Figure S1A and S1B). Altogether, these results indicate that I-Z-I proteins cannot align into striation in the absence of MHC.

Next, to test the role of non-muscle myosin in sarcomere formation, we stained Mhc1 mutant muscles with antibodies against zipper that corresponds to the only Drosophila non-muscle myosin. One prediction of the model emphasizing the role of non-muscle myosin is that removal of MHC would only have subtle effects on the zipper sarcomeric striation pattern, as non-muscle myosin would be involved in the formation of premyofilbrils. We find that zipper is strongly localized to Z lines but not to thick filaments (upper panels in Figure 1C), and that removal of MHC resulted in no obvious zipper periodic pattern (bottom panels in Figure 1C). These results suggest that, at least in Drosophila, intermediate non-muscle myosin-containing premyofilbrils do not exist during myofilbril assembly. Further, examination of other sarcomeric components (β-integrin Figure 1B, Mlp84B, titin, and Zasp [data not shown]) indicates that elimination of MHC leads to a complete disruption of the distribution of these proteins in myofilbrils.

As MHC is expressed in embryonic muscles several hours prior to sarcomere organization, one possibility is that MHC is required for the organization of proteins at later stages. To exclude this possibility, we stained embryos at stage 15–16 when many sarcomere proteins become enriched at muscle attachment sites. Removal of MHC had no effect on the distribution of integrin, α-actinin, zipper, kettin, Zasp and Mlp84B at the attachment sites.
 Strongly suggesting that formation of muscle attachments is a MHC-independent process.

To investigate the role of MHC in sarcomeric organization at later stages following sarcomere formation, we knocked down Mhc in third instar larvae by RNAi (Mhc-Gal4; UAS-Mhc-hp) and examined the distribution of sarcomeric components. Compared to control animals (Mhc-Gal4/+) the degree of loss of muscle sarcomere striation correlated well with the reduction in the MHC.

(Figures S2, S3, S4), strongly suggesting that formation of muscle attachments is a MHC-independent process.

To investigate the role of MHC in sarcomeric organization at later stages following sarcomere formation, we knocked down Mhc

Figure 1. MHC is critical for muscle striation formation. (A) Schematic organization of a myofibril, represented here with two sarcomeres. Sarcomeres are defined as the segment between two neighboring Z-lines. Thin filaments include actin filaments and their associated proteins such as troponins (Tns) (TnC, TnI, and TnT) and tropomyosin (Tm). Actin filaments are the major components of I-bands, and are cross-linked to Z-lines via α-actinin. Thick filaments are composed of myosin and are connected from the M-line to the Z-line by titin. A number of proteins important for the stability of sarcomeres, such as zipper and Zasp, are found in the Z-line. (B) Confocal fluorescent micrographs of control muscles of a stage 17 wild-type embryo (top panels) and myosin heavy chain (Mhc) amorphic mutant muscles from Mhc1 of same stage (bottom panels) stained by phalloidin (blue in merge), anti-α-actinin (red in merge) and anti-β-integrin (green in merge). Note that there is no obvious striation in Mhc null mutant muscles, and that β-integrin staining does not align with that of α-actinin. (C) Confocal images of control muscles of a stage 17 wild-type embryo (top panels) and Mhc1 of same stage (bottom panels) stained by anti-muscle MHC (blue in merge), anti-α-actinin (red in merge) and anti-zipper (green in merge). Note that in wild-type muscles, zipper colocalizes with α-actinin as shown in yellow in the merged image, but not with MHC. In addition, rat-anti-MHC was able to detect truncated MHC fragments in Mhc1 mutant muscles, and its staining overlaps with actin. This staining most likely reflects the ability of the Subfragment 1 region of MHC to bind to actin filaments. Scale bars: 10 μm.

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Figure 2. Tns-Tm complexes play an important role in sarcomere assembly. (A,B) Primary muscle cells were isolated from Oregon R embryos and treated with dsRNA against lacZ (A and B, upper panels), TnT (A, lower panels), or TnC (B, lower panels). Sarcomeric organization was evaluated...
following staining using polyclonal antibodies against MHC, α-actin, or actin. The efficacy of TnT and TnC RNAi knock-downs were evaluated using anti-TnT and anti-Tnc specific antibodies, respectively. Note that while removal of TnT has severe effects on the striated organization of the sarcomere, depletion of TnC has little effect. (C) To determine the relationships between TnT, Tnl, TnC and Tm, primary muscle cells were treated with TnT, Tnl, TnC or Tm RNAi and stained with antibodies against TnT, Tnl, TnC and Tm, respectively. Note the severe effects on the striated pattern in the absence of TnT, Tnl and Tm, but not TnC. Scale bars: 10 μm.

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expression level, as revealed by the fluorescent staining for α-actinin, Zasp and MHC (Figure S5). Altogether, these results indicate a critical role for MHC not only in sarcomere formation but also in its maintenance.

Function of the troponins-tropomyosin complex

The Tns-Tm (TnT/TnI/TnC/Tm) complex has a well-established role in acto-myosin interactions in response to changes in Ca²⁺ level. Since MHC is required for sarcomere formation, we analyzed whether this complex also participates in this process. Because complete null mutations in these genes are not available, and primary muscle RNAi has been successfully used to characterize sarcomere assembly [19], we used primary muscle cell cultures to assess the roles of Tns in sarcomereogenesis. Primary muscles were prepared from wild-type embryos, treated with various dsRNAs targeting different Tns, and stained with the three major sarcomeric proteins actin, α-actinin, and MHC. Depletion of TnT or TnI eliminated the striation pattern, indicating that both are essential for sarcomere formation (Figure 2A and Figure S6A). This is consistent with the in vivo sarcomeric disassembly phenotypes associated with loss-of-function TnT mutations described in mice and zebrafish and of Tnl in the indirect Drosophila flight muscle [24–26]. As previously proposed by Sparrow and colleagues [25], since Tnl inhibits the generation of acto-myosin force for muscle contraction in the absence of Ca²⁺ and TnT is involved in the attachment of the Tn complex to the thin filament, the defects observed in the absence of either TnT or TnI are likely caused by the upregulation of acto-myosin interactions between thin and thick filaments as they assemble.

Interestingly, we expected that knock-down of the Ca²⁺-binding protein TnC would affect sarcomere organization as loss of TnC should desensitize the response to Ca²⁺ concentration and reduce acto-myosin interaction between thin and thick filaments. However, knock-down of TnC did not lead to any significant change in MHC/α-actinin striation (Figure 2B). This non-essential role of TnC in sarcomere assembly underscores the importance of direct interactions between Tns and thin filament in sarcomere formation since, unlike TnT and TnI, TnC does not directly interact with actin and Tm [27].

Since Tns always function together with Tm to mediate actin-myosin interactions, we next examined the role of Tm in sarcomere organization. Staining of primary muscle cells treated with Tm dsRNAs with anti-actin, anti-α-actinin and anti-MHC demonstrated that Tm is another critical component participating in sarcomere assembly (Figure S6B). As TnT, TnI and Tm are all essential for sarcomere formation, we tested whether these three proteins are assembled in a sequential manner into sarcomeres, as observed in the case of other protein complexes involved in organelle assembly [28]. Thus, we depleted systematically each component of the Tns-Tm complex by RNAi and examined the overall organization of sarcomeres with antibodies against Tns and Tm. Strikingly, removal of TnT, Tnl or Tm led to random distribution of all remaining proteins, indicating that there is no loading sequence for Tns-Tm complex during sarcomere assembly (Figure 2C).

To test the possibility that removal of the Tns-Tm complex would lead to up-regulation of acto-myosin contraction by exposing the MHC binding sites on actin, and as a result causes disruption of the myofibril structure, we performed a time-course analysis of sarcomere striation. Typically, a clear striation pattern indicated by actin staining can be first observed in wild-type primary muscle cells culture at 3 days after plating at 25 °C [16], Figure S6C). In TnT RNAi treated cells, no striation could be detected at that time-point or later (Figure S6C) suggesting that the Tns-Tm complex plays an important role in the initial sarcomere assembly by allowing stable interdigitation of thin and thick filaments. Similar results were also obtained from Tnl or Tm RNAi treated primary muscles (data not shown). Interestingly, we did not observe a sequence of events in the recruitment of Tns and Tm, suggesting that the Tns-Tm complex is recruited as a complex to the sarcomere.

Relationship between MHC and I-Z-I

I-Z-I complexes are composed of thin filaments associated with nascent Z-lines and are the first identifiable structures during early myofibrillogenesis [3]. Subsequently, MHC has been hypothesized to assemble into thick filaments and to interdigitate with thin filaments to form sarcomeres.

To test directly this model, we knocked down each component of the I-Z-I complex including actin, zipper, Zasp, α-actinin and Mlp84B in primary muscle cells [29–31]. First, when actin was depleted by RNAi, primary muscles stained with anti-MHC showed much shorter and thinner striated myofibrils (Figure 3A). Since actin is essential for cell survival, the presence of striated myofibrils suggests that residual actin remains, which is consistent with the weak phalloidin staining observed (Figure 3A). We attempted to increase the severity of the RNAi phenotype by using a greater amount of dsRNAs (Figures S7 and S8), however, thin and shortened actin filaments were still present. There are six actin isoforms (Act5C, Act42A, Act57B, Act79B, Act87E and Act88F) in Drosophila, which only differ in a few amino acids. Act5C, Act42A, Act57B, Act87E and Act88F in Drosophila are muscle specific. Although the dsRNAs we used target all actin isoforms, our primary cell RNAi protocol may not allow complete depletion of actin isoforms, such as Act42A and Act5C, as they are expressed in myoblasts [17]. Since, it is possible that the initial sarcomere assembly in muscles utilizes cytoplasmic actins as building blocks, we speculate that the shortened actin filaments (Figure 3A) results from residual amount of cytoplasmic Act5C and Act42A. In further support of this model, it has been previously shown that in vertebrate muscles, the actin turn-over rate in thin filaments is over 100 fold lower than that in cytoplasmic actin filaments [33]. Thus, the extreme slow turn-over rate of these cytoplasmic actins, initially incorporated into thin filaments, could be sufficient to allow the formation of short sarcomeres [18].

Second, we investigated the role of Z-line components, such as α-actinin, zipper, Zasp, and Mlp84B, in sarcomere organization (Figure 3B–3D and data not shown). No significant changes in the striation pattern, as monitored by MHC staining, were observed in the α-actinin knock-down, indicating that α-actinin is not required for thick filament organization ([34], Figure 3B). Knock-down of
Figure 3. The I-Z-I complex is required for sarcomere organization. (A–D) DsRNAs against actin (A), α-actinin (B), zipper (C), Zasp (D) or control lacZ were applied to primary muscle cell cultures. Phalloidin, anti-α-actinin, anti-zipper and anti-Zasp antibodies revealed the knock-down
zipper led to fuzzy but discernable striation (Figure 3C and Figure S8A, S8B). Silencing of Zasp gave similar results as seen from zipper RNAi knock-down results (Figure 3D). Further, inactivation of Mlp84B appeared to have a negligible effect on the muscle sarcomere structure as determined by anti-MHC staining (data not shown). Finally, previous electron microscopic analysis of \(a\)-actinin, zip, and Mlp84B mutant muscles have shown that Z-disc structures are still present in these muscles [30,31,34], indicating that knock-down of these sarcomeric components does not affect the initial formation of sarcomeres.

Because removal of I-Z-I components individually did not lead to dramatic MHC disorganization, we tested whether these proteins act in a partially redundant manner by performing combinatorial perturbations. Simultaneous knock-downs of actin and one of Z-line proteins such as \(a\)-actinin, zip, or Zasp were associated with complete loss of striation and “stress-fiber-like” MHC phenotypes reminiscent of randomized MHC filaments (Figure 3E). These results suggest that myosin can assemble into thick filaments independently of I-Z-I complexes but that the sarcomeric structure of MHC filaments require the ordered arrangement I-Z-I complexes and vice versa.

**Zipper/Zasp/\(a\)-actinin may act as a tension sensor**

Previous studies have reported that zasp and \(a\)-actinin mutant muscles exhibit some defects in sarcomeric structures [30,34], which contrasts with our RNAi results that Zasp is not required for initial sarcomere assembly. To rule out the possibility that our results do not simply reflect a partial knock-down due to RNAi, we examined the muscle phenotypes of zasp and \(a\)-actinin amorphic mutants at the stage when myofibrils just start to form and display striated bands for MHC, \(a\)-actinin and Zasp (left panels in Figure 4A). At that stage, we could detect regularly spaced staining distribution of these proteins indicating that Zasp is not required for initial sarcomere assembly (Figure 4A). At later stages however, striation in zasp mutant embryos began to disappear, suggesting that Zasp is required for sarcomere maintenance (data not shown) [30].

Next, we analyzed \(a\)-actinin mutants using anti-\(a\)-actinin and anti-MHC antibodies and found that their Z lines were expanded, a phenotype consistent with previous electron microscopy studies (Figure 4B) [34]. Expanded Z lines were also present in \(a\)-actinin mutant muscles stained with kettin and Zasp antibodies (Figure 4C), indicating that \(a\)-actinin is required for Z-line condensation but not for sarcomere assembly or recruitment of other Z-line proteins such as titin/kettin and Zasp. In contrast, in kettin mutant muscles, no striation pattern was detected for any markers [33,36]. Collectively, these observations point out that the recruitment of Zasp, kettin and \(a\)-actinin to the Z-line does not occur in a step-wise fashion. Instead, it is more likely that they cooperate with each other in stabilizing Z-lines and are recruited concomitantly. Notably, zipper/Zasp/\(a\)-actinin are highly enriched at the muscle attachment sites and act downstream of integrin to link the muscle cytoskeleton to the cell membrane, suggesting that zipper/Zasp/\(a\)-actinin may act as a tension sensor to mediate the interaction between integrin and the internal cytoskeleton. Interestingly, single depletion of zipper, Zasp or \(a\)-actinin led to less severe phenotypes in sarcomere assembly in primary muscle cell cultures than in vivo (Figure 3A-3D), most likely reflecting the weaker stress on fibers in vitro versus in vivo.

To test this idea, we induced muscle cell stress in primary culture by increasing muscle cell intracellular calcium (Ca\(^{2+}\)) concentration. To increase contraction by augmenting the level of intracellular Ca\(^{2+}\), we knocked-down the plasma membrane calcium ATPase (PMCA) that pumps intracellular Ca\(^{2+}\) from the inside of cells [18]. We first found conditions where primary muscles were able to maintain relatively normal sarcomere structures when treated with 40 ng of dsRNAs against PMCA (Figure S10A). We next used this sensitized condition as a stress background for primary muscles in culture to examine the effect of reduction of those tension sensor components by double RNAi knock-down. Consistent with the idea that zipper/Zasp/\(a\)-actinin may act as a tension sensor, double knock-down with the mixture of dsRNAs targeting PMCA and either zipper or Zasp, or \(a\)-actinin led to significant disruption of sarcomeric organization, although primary muscles treated with either 40 ng of dsRNAs targeting PMCA, or 250 ng of dsRNAs targeting zipper or Zasp, or \(a\)-actinin still had normal sarcomere structures (Figure S10B). We further looked at the distribution of these tension-sensor proteins in muscles from paralyzed embryos caused by the mutation in immaucleate connections (inac), a gene that encodes a neuron-specific kinesin required for presynaptic maturation [37]. In agreement with our previous conclusions, these tension-sensor components were still present at muscle ends, although the sarcomere structures were disrupted, presumably due to less tension on the muscles in these animals. (Figure S11).

We then explored the functional interaction among zipper, Zasp and \(a\)-actinin, given that single knock-down of these Z-line proteins did not reveal a requirement in sarcomere assembly. Strikingly, removal of zipper, Zasp, and \(a\)-actinin together led to a complete loss of striation (Figure 4D), suggesting that they function redundantly in sarcomere assembly. Thus, single knock-down of only one of these tension sensor proteins is not sufficient to disrupt the whole complex, however, the combined removal of these components leads to a collapse of the entire sarcomere.

Finally, we examined the role of titin/kettin in sarcomere formation. Titin is a large protein localized at Z-line that contains multiple domains important for interactions among I-Z-I components. It has been proposed to act as a third filament to regulate plasticity of sarcomere. Consistent with its predominant role in the sarcomere, knock-down of titin is associated with complete disorganization of the sarcomere both in vitro and in vivo (Figure S12A and S12B). Removal of the scaffold protein titin may be equivalent to simultaneously disturbing the association of I-Z-I proteins and mimics the combinational knock-down of components from I-Z-I complex.

**Role of integrin in sarcomere assembly**

Genetic studies in C. elegans, Drosophila and mice have implicated the integrin pathway in sarcomere assembly [11–14]. Although, disruption of integrins is associated with a complete loss of...
myofibrillar striation in muscle cells [14], the exact role of integrins in sarcomere assembly has not been explored. Recently, integrin has been proposed to pave the way for sarcomere assembly by initiating the assembly of actin filaments at the muscle cell membrane [10]. A prediction of this model is that, in the absence of myosin, I-Z-I complexes should be tethered to costameres, which link muscle cell membrane to myofibrils, with integrins. To test this model, we dissected embryos from wild-type and Mhc\(^7\) mutant embryos that lack MHC and stained them with antibodies against integrin, actin and \(\alpha\)-actinin. In wild-type muscles, \(\beta\)-integrin is highly concentrated at the muscle attachment sites and relatively weakly stained at the position of the costameres with a striated pattern that appears to align with Z lines (Figure 1B). In contrast, we could not detect any striation for \(\beta\)-integrin at the location of the costameres in Mhc\(^7\) mutant muscles. No striation could be detected for actin and \(\alpha\)-actinin either. However, the pattern stained by actin and \(\alpha\)-actinin did not match that of \(\beta\)-integrin on the membrane, indicating that there is no obvious connection between Z lines and \(\beta\)-integrin in Mhc\(^7\) mutant muscles. These results suggest that integrins do not appear to be involved in the assembly of I-Z-I complexes, and that MHC is required for integrins to be aligned with Z-lines.

We further investigated the expression patterns of muscle markers in both wild-type and integrin null mutant (mys\(^{GAL4}\)) muscles of stage 17 embryos when myofibrils start to form [30]. At that stage, \(\beta\)-integrin deficient muscles contained multiple nuclei, indicating that the myoblast fusion process took place normally, and were not yet completely rounded up (Figure 5A). Further, in agreement with a previous report [14], sarcomeric structures in these muscles were totally disrupted. Immunostaining using antibodies against actin, MHC and \(\alpha\)-actinin revealed that the majority of both actin filaments and Z lines collapsed to the center of muscle cells and formed aggregates where myosin filaments were excluded (Figure 5A). This result shows that association of actin and \(\alpha\)-actinin occurs even in the absence of integrins, implying that 1-Z-I complex formation may not require integrins. Furthermore, in the absence of integrins, thin and thick filaments cannot interdigitate with each other to form sarcomeres. Altogether, our immunohistochemical study results are consistent with previous electron microscopy analysis of integrin mutant muscles, which contained disorganized thin and thick filaments distributed in the central region of myotubes [30]. Further, a recent report has elegantly dissected the dual role of integrin in muscle attachment and sarcomeric organization using an inducible targeted RNAi system [39]. This study revealed that reduction of integrin led to the dissociation of Z-disc proteins into small dots, whereas muscle attachment sites were not affected, and thus suggested a role for integrin in maintenance of sarcomere structures that may be different from its role at attachment sites. Altogether, based on the data from our study and others [14,33], integrins appear to play very important roles in maintaining the tension at the cell periphery for organization of sarcomere structures. We propose that integrins serve as anchor points for the floating I-Z-I complex and provide tensions that allow the interdigitation of thin and thick filaments.

A model for sarcomere assembly

Based on our in vitro and in vivo mutant analyses, we propose that sarcomeric proteins exist in two states: one being the chaotic but independently assembled differential functional complexes, and the other one being the highly ordered suprastructure made from these complexes (Figure 5B). We propose that functional complexes including integrin, zipper/Zasp, \(\alpha\)-actinin, I-Z-I, Tns-1m and MHC first assemble independently (as described in shadowed boxes in Figure 5C). Subsequently, we propose that integrins serve as anchor points for the floating I-Z-I complex starting from muscle ends, and provide tensions that allow the interdigitation of thin and thick filaments for de novo sarcomere assembly. This in turn may facilitate the spatial organization of integrins on cell membrane by linking the integrins to the underlying cytoskeleton and “towing” the integrin adhesion complex into periodic position, which may further strengthen newly formed myofibril structures. In the context of our loss of function studies, only those perturbations that affect the formation of the independent complexes will lead to the failure in assembly of striated sarcomeres. If depletion of any one protein has negligible effects on the localization of the complex where it belongs to, then sarcomere striation will be maintained. Altogether, our study provides a comprehensive analysis of the functional effects of perturbation of sarcomeric proteins on myofibril assembly. Because of the similarities between Drosophila and vertebrate sarcomeres the “two-state” assembly model that we have proposed should prove relevant to striated muscles.

Methods

Drosophila genetics

Oregon R was used as a wild-type strain. Mutations in genes encoding sarcomeric proteins are: Mhc\(^1\), zasp\(^{1-}\), mys\(^{GAL4}\), Actn\(^{14}\), imac\(^{F70}\) ([11,23,30,34]), and zpf\(^{22G}\). Mutant strains were balanced with twi-GAL4>UAS-GFP or Kr-GFP and mutant primary muscle cells were identified by lack of GFP expression. For RNAi experiments, Mhc-Gal4 [40] and UAS-Mhc hp [41] flies were used.

Embryonic primary cell culture

Embryonic primary cell cultures were established as previously described [17,42]. Eggs were collected on molasses plates for 2 hrs and incubated for 4 hrs at 25°C. Embryos were dechorionated in 50% bleach for 3 mins, rinsed thoroughly with 70% ethanol and sterilized water, and dissociated into a cell suspension using Dounce homogenizers (WWR Scientific, Seattle, WA) in Shields and Sang M3 medium (Sigma). Cell suspensions were spun once at 40 g for 10 mins to pellet tissue debris, large cell clumps and vitelline membranes. Supernatants were then transferred to a fresh tube and spun at 360 g for 10 mins to pellet the cells. Cells were washed once and re-suspended in primary cell medium (10% heat-

Figure 4. Zipper/Zasp/\(\alpha\)-actinin acts as a tension sensor to regulate sarcomere assembly. (A) Confocal micrographs of control muscles of a stage 17 wild-type embryo (left panels) and age-comparable zasp null mutant muscles (right panels) stained for MHC (blue in merge), \(\alpha\)-actinin (red in merge) and Zasp (green in merge). Scale bar: 10 μm. (B) Confocal micrographs of control muscles of a stage 17 wild-type embryo (left panels) and same stage \(\alpha\)-actinin null mutant muscles from Actn\(^{14}\) (right panels) stained for actin (blue in merge), \(\alpha\)-actinin (red in merge) and MHC (green in merge). Scale bar: 10 μm. (C) Fluorescent confocal micrographs of control muscles of a stage 17 wild-type embryo (top panels) and \(\alpha\)-actinin null mutant muscles from Actn\(^{14}\) of same stage (bottom panels) stained for Zasp (blue in merge), \(\alpha\)-actinin (red in merge) and kettin (green in merge). Note that \(\alpha\)-actinin null mutant muscles still have striated sarcomeres, but with expanded Z lines. Scale bar: 10 μm. (D) Primary muscle cells were treated with combinations of dsRNAs targeting components of the zipper/Zasp/\(\alpha\)-actinin complex. Muscle striation was evaluated using anti-MHC, anti-actin and anti-\(\alpha\)-actinin antibodies. Scale bar: 10 μm.

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inactivated fetal bovine serum from JRH Biosciences, 10 mU/ml bovine insulin in M3 medium from Sigma). Cells were seeded and grown in 384-well optically clear plastic plates (Costar) at 1.7–2.5 × 10^5 cells/cm^2.

**Primary cell RNAi treatment**

DsrRNAs synthesis and RNAi treatment were performed according to the DRSC protocols (http://flyRNAi.org) using amplicons to target various genes: DRSC23472 and DRSC23961.
for actin; DRSC14106 for α-actinin, DRSC03367 and DRSC25959 for Mhc; DRSC20790 for troponin T; DRSC29930 for troponin I; DRSC04916 and DRSC07564 and DRSC11307 for troponin C; DRSC16887 and DRSC26804 for troponymin; DRSC04725 and DRSC22499 and DRSC20416 for zipper; DRSC07121 for Zasp; DRSC20341 and DRSC39450 for integrin; DRSC39473 and DRSC34374 for PMCA. dsRNA against lacZ was used as a control in all the above experiments. Primary muscle cells were prepared as described above and seeded at approximately 4×10^4 cells per well in plates containing dsRNAs. After 22 hrs in serum-free M3 medium, additional serum-containing culture medium was added to bring the solution to a final concentration of 10% FCS. For multiple knock-down experiments, dsRNAs were mixed into the plate before adding the muscle cells.

**Immunofluorescence microscopy**

Flies were allowed to lay eggs for 2 hrs and embryos further developed at 25°C for 14–16 hrs to reach stage 17 of embryogenesis. The body wall muscles of staged embryos were dissected by following methods described previously [16]. Dissected muscles or primary cells were fixed with 4% formaldehyde. Cells were stained overnight at 4°C with the first antibody or phalloidin Alexa Fluor 568 (Molecular Probes; 1:200) and DAPI (Sigma, 1:5000) in PBT (PBS, 0.1% Triton X-100, 1% BSA), washed once in PBS and left in PBS containing 0.02% NaN3. For whole mount embryo stainings at stages 15–16, standard protocol was followed, including dechorionization, fixation, rehydration, and staining with antibodies.

Antibodies used in this study: rat anti-actin, anti-MHC, anti-TnT, anti-TnC, anti-Tm (1:500) (The Babraham Institute, Cambridge, UK), mouse anti-α-actinin (1:100) (from Dr J. Saide, Boston University, Boston, MA), rabbit anti-zipper (1:500) (from Dr. D. Kiehart), rabbit anti-Zasp (1:1000) (from Dr. F. Scho¨ck), anti-kettin (1:1000) (Dr. D. Andrew), anti-PS3/anti-zipper (1:500) (from Dr. D. Andrew), anti-PS3/anti-MHC (1:100) (Developmental studies hybridoma bank). The second antibodies from invitrogen include Alexa 488 and Alexa 594.

**Generation of anti-Tnl antibody**

Full-length Tnl was cloned into pGEX4T-1 and GST fusion protein was purified according to the protocols described from Pharmacia. The recombinant protein was sent to Abmart Antibody Company (Shanghai, China) and the antibody was purified using the original antigen. Tnl antibody specificity was confirmed by cell staining results of TnT RNAi-treated muscle cells.

**Real-time PCR**

qRT-PCR was performed to analyze the RNA level of actin in Drosophila S2 cells treated by dsRNA against lacZ or acta in different concentrations followed by reverse transcription using the SuperScript first-strand synthesis kit (Invitrogen). The experiment was carried out in Stratagene MX4000 thermocycler using SYBR GreenTM detection protocol. Sequencing primers for Drosophila actin, forward: GAGGCCGCTTACAGCTTCA, reverse: TCC-TTGATGTCCGGCCACA.

**Supporting Information**

**Figure S1** MHC is required for sarcomere formation. (A,B) Primary muscle culture cells were isolated from Mhc-GFP and treated with Mhc dsRNA. (A) Mhc1 mutant embryos. (B) Anti-MHC staining was used to assess the knock-down efficiency or to identify Mhc mutant primary muscle cells. Cultures were immunostained using anti-actin or anti-α-actinin to analyze sarcomeric structures. Anti-zipper antibody was used to analyze the localization of zipper in the absence of MHC. Scale bars: 10 μm. Found at: doi:10.1371/journal.pgen.1001208.s001 (2.20 MB TIF)

**Figure S2** Removal of MHC has no effect on the localization of integrin and α-actinin. (A,B) Staining of integrin and α-actinin was performed in both wild-type and Mhc1 mutant stage 15–16 embryos. MHC antibody was used to analyze Mhc1 null allele. Both MHC and other sarcomeric protein stainings were merged to check their localization relationships. Scale bar: 20 μm. Found at: doi:10.1371/journal.pgen.1001208.s002 (7.76 MB TIF)

**Figure S3** Localization of zipper and kettin at muscle attachment sites in the absence of MHC. (A,B) Zipper and kettin were stained with antibodies to show their muscle attachment site localization in the presence and absence of MHC in stage 15–16 embryos. Scale bar: 20 μm. Found at: doi:10.1371/journal.pgen.1001208.s003 (6.09 MB TIF)

**Figure S4** Zasp and Mlp84B localize at muscle attachment site in a MHC-independent manner. (A,B) Zasp and Mlp84B distribution were assessed by their antibody stainings and merged with MHC staining to check their localization relationships in 15–16 stage embryos. Scale bars: 20 μm. Found at: doi:10.1371/journal.pgen.1001208.s004 (6.41 MB TIF)

**Figure S5** Localization of sarcomeric components in larval muscles is disrupted upon MHC reduction. Confocal micrographs of second instar larval body wall muscles from control animal (top panels) and age comparable muscles from a larva carrying transgenes of Mhc-Gal4;UAS-Mhc hp (bottom panels) stained for MHC (green in merge), β-actinin (blue in merge) and Zasp (red in merge). Scale bar: 50 μm. Note that the presence of striated organization of these sarcomeric components correlated well with the presence of MHC expression (arrowheads at bottom panels), while loss of MHC expression led to disruption of sarcomere striation and distribution of these sarcomeric proteins. Found at: doi:10.1371/journal.pgen.1001208.s005 (2.73 MB TIF)

**Figure S6** The Tn-Tm complex is essential for sarcomere assembly. (A,B) DrsRNAs against Tnl or Tm were applied to primary muscle cells, and anti-Tnl and anti-Tm antibodies were used to document the knock-down effectiveness. The sarcomeric organization of treated muscles was analyzed using anti-actin and anti-α-actinin antibodies. (C) Tn-T knock-down time course experiment. No striation was observed in Tnl RNAi-treated primary muscle cells, even at 3 days after plating when the sarcomeres begin to form in the lacZ RNAi control. Muscle cultures were stained with anti-MHC antibody. Scale bars: 10 μm. Found at: doi:10.1371/journal.pgen.1001208.s006 (3.84 MB TIF)

**Figure S7** Persistent arrest of residual actin protein in myofibril after dsRNA treatment. Different amounts of actin dsRNA were added to primary muscle cell culture from 50 ng to 1 mg. lacZ dsRNA was used as a control. Anti-actin antibody was applied for analysis of residual actin signal and anti-MHC for muscle structure. Scale bars: 10 μm. Found at: doi:10.1371/journal.pgen.1001208.s007 (3.38 MB TIF)

**Figure S8** Quantitative RT-PCR analysis of actin RNAi efficiency. 250 ng or 800 ng of dsRNA against actin were applied to Drosophila S2 cells in comparison of treatment with 250 ng of lacZ dsRNA. Quantitative RT-PCR analysis was performed to assess the actin knock-down effectiveness. The amount of actin mRNA from lacZ dsRNA treatment was used as a normalization control. Found at: doi:10.1371/journal.pgen.1001208.s008 (0.27 MB TIF)
Figure S9 Zipper is not exclusively required for sarcomere striation. (A) Primary muscle cells treated with zipper dsRNA and stained using anti-MHC, anti-α-actinin. Anti-β-actin antibody was used as an anti-MHC control. Developmental stage: 18–19 h. (B) Primary muscle cells were isolated from 18–19 h/zipper and zipper mutant embryos identified by the lack of GFP expression. Cultures were stained with anti-MHC, anti-α-actinin, and anti-β-actinin. Scale bars: 10 μm. Found at: doi:10.1371/journal.pgen.1001208.s010 (7.08 MB TIF)

Figure S10 Zipper/Zasp/α-actinin senses Ca²⁺ stress in vitro. (A) Various amounts of dsRNAs against PMCA were added to primary muscle cells. Muscle striation was monitored by anti-PMCA-actinin. Scale bars: 10 μm. Note that the tension sensor components Zasp and α-actinin were still localized at muscle attachment sites (arrows in the bottom panels) even though the sarcomere structures were disrupted in imac null mutant muscles. Found at: doi:10.1371/journal.pgen.1001208.s011 (3.61 MB TIF)

Figure S11 Tension sensor components are still localized at muscle ends in the paralyzed animals. Confocal micrographs of late embryonic body wall muscles from a control animal (top panels), and age comparable muscles from imac null mutant (bottom panels) stained for MHC (green in merge), α-actinin (blue in merge) and Zasp (red in merge). Scale bar: 10 μm. Found at: doi:10.1371/journal.pgen.1001208.s010 (7.09 MB TIF)

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Author Contributions

Conceived and designed the experiments: YR. Performed the experiments: YR. Analyzed the data: YR. Contributed reagents/materials/analysis tools: YR. Wrote the paper: YR.

References


