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ACTIN SCAFFOLDING BY CLATHRIN HEAVY CHAIN IS REQUIRED FOR SKELETAL MUSCLE SARCOMERE ORGANIZATION

Stéphane Vassilopoulos^{1,2,3,4} #, Christel Gentil^{1,2,3,4}, Jeanne Lainé^{1,2,3,4,5}, Pierre-Olivier Buclez^{1,2,3,4,7}, Agathe Franck^{1,2,3,4}, Arnaud Ferry^{1,2,3,4,6}, Guillaume Précigout^{1,2,3,4,7}, Robyn Roth⁸, John E. Heuser⁸, Frances M. Brodsky⁹, Luis Garcia^{1,2,3,4,7}, Gisèle Bonne^{1,2,3,4}, Thomas Voit^{1,2,3,4}, France Piétri-Rouxel^{1,2,3,4} * and Marc Bitoun^{1,2,3,4} *.

¹ Université Pierre et Marie Curie-Paris 6, UM 76, Paris, F-75013, France.

² INSERM U974, Paris F-75013, France.

³ CNRS UMR 7215, Paris F-75013, France.

⁴ Institut de Myologie, Paris, France.

⁵ Université Pierre et Marie Curie-Paris 6, Site Pitié-Salpêtrière, Département de Physiologie, Paris F-75013 France.

⁶ Université Paris Descartes-Paris 5, Paris, France.

⁷ Université de Versailles – Saint-Quentin-en-Yvelines, Versailles, France.

⁸ Department of Cell Biology and Physiology, Washington University, St. Louis. USA

⁹ Departments of Bioengineering and Therapeutic Sciences, Pharmaceutical Chemistry, and Microbiology and Immunology, UCSF, San Francisco, USA.

* These authors have contributed equally to this work

To whom correspondence should be addressed

Stéphane Vassilopoulos
Inserm U974, Institut de Myologie, Paris, France.
E-mail: s.vassilopoulos@institut-myologie.org
Tel: 33 (0) 1.42.16.57.12
Fax: 33 (0) 1.42.16.57.00

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Abstract

The ubiquitous clathrin heavy chain (CHC), the main component of clathrin coated vesicles, is well characterized for its role in intracellular membrane traffic and endocytosis from the plasma membrane (PM). Here, we demonstrate that in skeletal muscle CHC regulates the formation and maintenance of plasma membrane-sarcomere attachment sites also known as costameres. We show that clathrin forms large coated lattices associated with actin filaments and the muscle-specific isoform of α -actinin at the PM of differentiated myotubes. Depletion of CHC in myotubes induced a loss of actin and α -actinin sarcomeric organization, whereas CHC depletion *in vivo* induced a loss of contractile force due to the detachment of sarcomeres from the PM. Our results suggest that CHC contributes to the formation and maintenance of the contractile apparatus through interactions with costameric proteins and highlight an unconventional role for CHC in skeletal muscle which may be relevant to pathophysiology of neuromuscular disorders.

Introduction

Clathrin is composed of trimerized heavy chains on top of which associate three light chains (CLC) and assembles to form a membrane coat. The clathrin lattice formation on cellular membranes is initiated by clathrin adaptors which are drawn into the lattice and trigger clathrin-coated vesicle (CCV) budding at various subcellular compartments (Kirchhausen, 1999; Pearse and Robinson, 1990) reviewed in (Brodsky et al., 2001). Once the invagination process has initiated, dynamin family members associate with the neck of the forming pit, recruit additional proteins including actin and allow the scission of the coated vesicle. The role of dynamin 2 (DNM2) in skeletal muscle has received considerable attention because the autosomal dominant form of centronuclear myopathy (CNM) has been linked to mutations in the *DNM2* gene (Bitoun et al., 2005). DNM2 is involved in various cellular processes but to date it remains unclear whether its role in CCV fission is implicated in centronuclear myopathy.

Several recent studies have proposed a role for clathrin and dynamin in actin organization that is distinct from coated vesicle formation (Bonazzi et al., 2012; Bonazzi et al., 2011; Saffarian et al., 2009; Schafer et al., 2002; Veiga et al., 2007). Intriguingly, a role for clathrin in myofibril assembly was previously proposed (Kaufman et al., 1990) and several teams have reported direct protein-protein interaction between clathrin heavy chain, α -actinin and vinculin, two focal adhesion proteins crucial for muscle function (Burrige et al., 1980; Fausser et al., 1993; Merisko, 1985; Merisko et al., 1988; Schook et al., 1979). Alpha-actinin and vinculin are expressed in both skeletal and cardiac muscles where they are localized to a specialized compartment called the « costamere ».

Costameres, muscle-specific adhesion sites, are sub-sarcolemmal protein assemblies circumferentially aligned in register with Z-disks of peripheral myofibrils. They were

originally described as electron-dense plaques rich in the focal adhesion protein vinculin and function as attachment complexes (Pardo et al., 1983a; Shear and Bloch, 1985). They are located between the plasma membrane and myofibrils and play both a mechanical and a signalling role during contraction, transmitting force from the contractile apparatus to the extracellular matrix in order to stabilize skeletal muscle fibers. They are considered as the "Achilles' Heel" of skeletal muscle because, when disrupted, they directly contribute to the development of several distinct myopathies (Blake et al., 2002; Ervasti, 2003). In the present work, we have undertaken the study of the role played by CHC in skeletal muscle costamereogenesis and costamere maintenance. We show that clathrin associates with α -actinin at the surface of myotubes and forms large plaques which are associated with actin filaments. Depletion of CHC induces severe defects in α -actinin distribution and subsequently leads to defective costamere formation *in vitro* and induces impairment of contractile properties associated with structural abnormalities including sarcomere disorganization as well as detachment of the sarcomeres from the sarcolemma *in vivo*.

Results

CHC is a component of the costameric complex in skeletal muscle.

In adult skeletal muscle, DNM2 and CHC are both localized at the level of the I-band and present a striated pattern which overlaps with the muscle-specific α -actinin (isoform 2) staining (Fig. 1, A and B) (Butler et al., 1997; Cowling et al., 2011; Durieux et al., 2010; Kaufman et al., 1990; Towler et al., 2004). Using confocal optical sectioning and 3D reconstruction, we demonstrated that this striated labelling was mainly present at the surface of dissociated fibers (Fig. 1, C and D). Analysis of the compartment labelled by anti-CHC antibodies on adult skeletal muscle using electron microscopy revealed that clathrin is present along the sarcolemma and localizes at sites which correspond to contacts between sarcolemma and Z-discs which are also densely stained with tannic acid (Fig. S1, A and B). As several studies have described a direct association between α -actinin and vinculin with clathrin (Burrige et al., 1980; Fausser et al., 1993; Merisko, 1985; Merisko et al., 1988; Schook et al., 1979), we then assessed the association between clathrin and the α -actinin-vinculin-talin complex in adult mouse skeletal muscle (Fig. 1 E). We observed co-immunoprecipitation of CHC and clathrin light chain upon α -actinin immunoprecipitation but only minor amounts of vinculin and talin. Additionally, minor amounts of α -actinin are detected upon clathrin immunoprecipitation while vinculin co-immunoprecipitates talin efficiently and some α -actinin, but not clathrin. This suggests that part of the α -actinin pool is associated with clathrin in a distinct complex than the vinculin-talin- α -actinin complex.

We further characterized the sites of interaction between CHC and α -actinin during the early steps of differentiation in primary myotubes and showed that these proteins partially colocalize along the PM after 4 days of differentiation (Fig. 2 A). By sub-cellular fractionation, we observed an association between α -actinin and CHC in the PM fraction of fibroblasts, myoblasts and differentiated myotubes. This association was stronger in

differentiated myotubes as the levels of α -actinin expression increase (Fig. 2, B and C). In addition, we detected α -actin in both CHC and α -actinin immunoprecipitates from muscle cells. Several adaptor proteins are required for targeting clathrin to specific intracellular compartments, among which AP2 involved in recruiting clathrin to the PM. In differentiated myotubes, AP2 was localized at the surface of myotubes and displayed a predominant PM localization (Fig. 2 D and E). These results show that clathrin is a costameric protein which is recruited by AP2 and associates with actin and α -actinin at the PM of differentiated myotubes.

Costameric clathrin forms large plaques where α -actinin and actin are localized.

Myofibril assembly and subsequent sarcomere formation initiate at the PM. Following the formation of a sub-cortical actin lattice, additional nascent sarcomeres keep adding to the initial actin network (Dlugosz et al., 1984; Epstein and Fischman, 1991; Quach and Rando, 2006; Sanger et al., 2002). We reasoned that α -actinin and actin rely on a clathrin scaffold at the PM to start recruiting additional myofibrils and explored this possibility using immunogold cytochemistry coupled with quick-freeze, deep-etch rotary replication and electron microscopy (Heuser, 1980). We showed that clathrin forms two distinct coated structures; i.e. large plaques which contain much more polymerised clathrin that would be required to form a single vesicle (pseudo-colored in red in Fig. 3, D and F) and coated-vesicles found in regions devoid of plaques (Fig. 3, D and F, Fig. S1, I-M). Using clathrin antibodies we were able to confirm that CHC was indeed the proteinaceous material composing these structures (Fig. S1). As expected, α -actinin strongly labelled dense longitudinally organized actin filament bundles (arrows in Fig. 3 D and Fig. S2, A and B) which are reminiscent of the contractile apparatus. In addition, α -actinin was associated with actin filaments which surround or overlap flat clathrin lattices (Fig. 3, E and G and Fig. S2, A,

C and D) highly enriched with the adhesion marker $\beta 5$ integrin (Fig. S2, E-H) as previously described on rat myotubes (De Deyne et al., 1998; Pumplin, 1989; Pumplin and Bloch, 1990) while vinculin was not localized at these clathrin-coated plaques (our unpublished data). Our experiments on primary myotubes show that CHC forms large plaques that may serve to anchor the actin cytoskeleton and α -actinin.

CHC is required for the formation of costameres.

We next tested the effect of CHC knock-down on α -actinin sub-cellular organization. We achieved high knock-down efficiency in C2C12 myotubes differentiated for 6 days (Fig. 4, A and B) with two previously published siRNA sequences “siRNA1 and siRNA2” (Ezratty et al., 2009). Additionally, we showed that levels of CLC were also reduced because of the instability of the light chain upon heavy chain depletion as previously reported in non muscle cells (Brodsky, 1985; Esk et al., 2010). Expression level of α -actinin, vinculin and talin were unaffected in myotubes treated with CHC siRNA (Fig. 4 A).

As expected, control myotubes present a striated α -actinin distribution after 6 days of differentiation. This striated pattern is first visible at the periphery of the myotube and gradually fills the entire cytoplasm (Quach and Rando, 2006). In CHC depleted myotubes, the α -actinin striated pattern was strongly disrupted and instead, clusters of α -actinin were randomly dispersed in the intracellular space (Fig. 4 C). Three dimensional projections of confocal sections confirmed a dramatic reduction of the striated pattern between controls and CHC depleted myotubes (Fig. 4 D). Upon CHC depletion, a similar phenotype arises in mouse primary myotubes (Fig. 4, E and F) in which organization of actin filaments is also affected. To determine the role played by clathrin during the early steps of costamere formation, we performed siRNA depletion experiments on C2C12 myotubes following 4 days of differentiation, before α -actinin and actin could reach a striated organization. In control

myotubes, α -actinin was mainly localized along the PM. In stark contrast, myotubes which had been depleted of CHC presented a strong increase in the amount of intracellular α -actinin (Fig. 5 A). These results demonstrate that CHC is required for proper α -actinin distribution during early myotube differentiation.

AP2 and DNM2 are required for α -actinin scaffold formation.

We next tested the effect of AP2 knock-down on α -actinin sub-cellular localization. AP2 (α -subunit) depletion by siRNA (Fig. S3 C) induced a strong decrease in the amount of CHC recruited at the PM. More importantly, AP2 knock-down phenocopied the effect of CHC depletion on α -actinin distribution and dramatically increased the intracellular α -actinin staining (Fig. 5). We have also depleted the AP1 (γ -subunit) and AP3 (δ -subunit) adaptors responsible for CHC targeting at the Golgi apparatus and endosomal systems. As expected, AP1 and AP3 were localized at the perinuclear region and in intracellular structures dispersed throughout the myotube (Fig. S3 A). siRNA-depletion of AP1 and AP3 drastically reduced the intracellular CHC staining without affecting CHC localization at the PM (our unpublished data). Moreover, knock-down of AP1 and AP3 had no effect on α -actinin distribution (Fig. 5) suggesting that the impact of CHC depletion on α -actinin organization is not due to perturbations of the secretory pathway or trafficking between the endo/lysosomal system.

DNM2 is also present at or near the plasma membrane of differentiating myotubes where its staining partially overlaps with that of α -actinin (Fig. S4 C). AP2 depletion strongly reduced the amount of DNM2 at the plasma membrane suggesting that part of the PM staining corresponds to an association with clathrin lattices (Fig. S4 D). Efficient DNM2 depletion (Fig. S3 A) induced an increase in the amount of intracellular α -actinin staining (Fig. 5 and Fig. S4 A). Of interest, these intracellular aggregates observed upon CHC, AP2 and DNM2 depletion also contain actin aggregates (Fig. S4 A).

We performed the quantification of the intracellular α -actinin-positive area for all siRNA (Fig. 5 B). CHC, AP2 and DNM2 had a strong impact on the intracellular α -actinin area while AP1 and AP3 values were not different from controls. Altogether, these results confirm that the primary impact observed upon CHC depletion on α -actinin organization is related to its cytoskeleton scaffolding role at the PM through a process also involving AP2 and DNM2.

Hip1R depletion stabilizes α -actinin and actin on clathrin-coated structures.

Actin has been shown to be present at clathrin plaques (Collins et al., 2011) and required for their formation (Saffarian et al., 2009). The interaction between actin and clathrin may be mediated by Hip1R and the clathrin light chain (CLC) (Chen and Brodsky, 2005; Engqvist-Goldstein et al., 2001; Engqvist-Goldstein et al., 2004; Legendre-Guillemain et al., 2005). In myotubes, Hip1R colocalizes with AP2-positive puncta in differentiated myotubes (Fig. S3 D). We tested the effect of both CHC and AP2 depletion on Hip1R (Fig. S3 D) and showed a dramatic reduction of the Hip1R staining at the PM. These results confirm that CHC and AP2 are required for recruiting the clathrin-associated actin binding protein Hip1R to the PM. It has been recently shown that Hip1R depletion stabilizes the actin filaments on clathrin coated plaques in HeLa cells (Engqvist-Goldstein et al., 2004). We decided to test the contribution of Hip1R on the distribution of actin and α -actinin in differentiated myotubes. siRNA-depletion of Hip1R drastically reduced Hip1R protein levels (Fig. S3 A). Hip1R depleted myotubes presented a disorganized actin cytoskeleton and α -actinin distribution. The Hip1R depleted myotubes accumulated unusual actin structures near the cell cortex (Fig. 6). These structures, which are very similar to the ones observed by Engqvist-Goldstein et al., upon Hip1R depletion in HeLa cells (Engqvist-Goldstein et al., 2004) took a variety of forms including tails and rings at the bottom and the top of differentiated myotubes (Fig. 6 and Fig. S4 B). These results demonstrate that when Hip1R expression is reduced, actin polymerization at clathrin-

coated structures is no longer properly regulated and results in accumulation of actin and α -actinin on clathrin-coated plaques.

CHC is required for sarcomere maintenance in mature isolated muscle fibers.

We then assessed the function of CHC in mature skeletal muscle fibers. For this purpose, we developed an AAV strategy combined to the use of a short hairpin RNA (shRNA) against CHC (AAV-shCHC) for *ex vivo* transduction of isolated fibers from Flexor Digitorum Brevis (FDB) muscle. In AAV-shCHC infected fibers, a strong decrease in CHC expression was confirmed, both at the RNA and protein level (Fig. 7, A-C), and α -actinin was no longer correctly localized, with very few striations noticeable and displaying a diffuse or disorganized staining pattern (Fig. 7 D). We also tested the effect of CHC depletion on DNM2 distribution. In control fibers, DNM2 displayed a striated pattern at the surface, with each striation being centered on the I-band, and in the core of the fiber, DNM2 was mainly present at the PM (Cowling et al., 2011; Durieux et al., 2010). In fibers depleted of CHC, the DNM2 signal at the core of the fiber was greatly reduced at the PM (Fig. 7 E). In order to assess the effect of CHC depletion on the integrity of the actin cytoskeleton we used phalloidin staining. In control fibers, F-actin displayed a regularly spaced striated distribution. However, CHC depleted fibers presented a diffuse actin distribution (Fig. 7 F) and in some fibers detachment of entire sarcomeric bundles (Fig. 7 G). In addition to its role in costamereogenesis, these data suggest that CHC is also involved in their preservation in mature fibers.

***In vivo* CHC depletion impairs force and causes muscle degeneration.**

We used AAV-shCHC in order to induce an *in vivo* depletion of CHC in Tibialis anterior (TA) muscle. By performing Hematoxylin and Eosin staining on transverse TA sections (Fig. 8 A) no histological change was visible in muscles injected with control AAV or with AAV-

shCHC until day 18 post-injection (PI). However, fibers displaying a rounded shape with a clearly disorganized intracellular space were noticed at day 21 PI. Between days 21 and 23 PI, some degenerating fibers appeared and by day 25 PI, the vast majority of the fibers had a rounded shape with a clear degenerating phenotype. As a general feature of muscle degeneration, an immune infiltration was visible at day 21 and was extremely pronounced by days 23 and 25 PI. At day 25, small myotubes appeared and their presence reached a peak by day 38 where the majority of the muscle was centrally nucleated, the signature of a muscle undergoing massive regeneration.

Quantitative RT-PCR and Western blot analysis confirmed that at day 18 PI, a strong decrease in CHC had indeed occurred (Fig. 8, B-D). When we measured CHC RNA and protein levels at later time points, we noticed that CHC levels increased from 40% back to 55% of control levels at day 21 PI and were back to control levels by day 23. Surprisingly, at day 25 PI, CHC levels were higher in the AAV-shCHC injected muscle compared to the contralateral control muscle (Fig. 8, B and C). A possible explanation is that the infiltration by macrophages contaminates our analysis with exogenous clathrin from non transduced immune cells. Indeed, at days 21, 23 and 25 PI, a strong increase in CD11b protein levels was evident (Fig. 8 C and Fig. S5, A and B). Immunofluorescent staining confirmed that CD11b-positive macrophages had massively infiltrated the CHC depleted muscle. In fact, CD11b protein levels correlated perfectly with the sudden increase in CHC levels. In order to circumvent the immune infiltration and to show that a clathrin knock-down had occurred, we performed immunoprecipitation experiments using antibodies against the muscle specific α -actinin isoform. This assay enabled us to show that in muscles injected with AAV-shCHC, there was a strong decrease in the amount of CHC which co-immunoprecipitated with α -actinin (Fig. 8 E).

Direct muscle stimulation was performed in order to measure the maximal isometric strength

of the TA muscle in CHC-depleted muscles. At day 18 PI, when no morphological alteration could be observed, CHC depletion induced a 57% decrease in specific maximal force, *i.e.* absolute maximal force divided by muscle weight (Fig. 8 F). At day 21 PI, CHC depletion had an even more robust effect since muscle depleted of CHC displayed a 74% decrease in specific maximal force (Fig. 8 F). No significant muscle mass loss was observed at both early time-points (Fig. 8 G). These results show that AAV-shCHC efficiently knock-down CHC *in vivo* with dramatic consequences in terms of muscle fiber homeostasis.

***In vivo* CHC depletion perturbs α -actinin, γ -actin and DNM2 distribution.**

We then sought to analyze the distribution of α -actinin, DNM2 and the costameric actin isoform γ -actin (Sonnemann et al., 2006) in CHC-depleted skeletal muscle by immunocytochemistry. At 18 days PI, the α -actinin labelling was not different from the control while a clear reduction in sarcolemmal clathrin was apparent (Fig. 9 A). However, at day 21 and at further time points (day 23 and day 25), α -actinin and γ -actin both displayed a diffuse distribution compared to control muscle and areas where α -actinin was no longer adjacent to the sarcolemma were also visible (Fig. 9, B-D). In addition, in control fibers DNM2 presented a strong PM staining and a weak intracellular staining. At day 21 and at further time points (day 23 and day 25) the DNM2 intracellular staining was greatly enhanced (Fig. 9 B and D). We also tested the effect of CHC depletion on the distribution of other costameric proteins such as dystrophin, β -dystroglycan, and Caveolin-3 (Fig. S5, B and C). Even though at days 23 and 25 the muscle presented a severe dystrophic phenotype, no change in the distribution of these proteins was noticeable. We therefore conclude that CHC depletion induces a strong effect on the localization of intracellular costameric proteins related to the actin cytoskeleton including γ -actin, actin-associated proteins, and DNM2, without affecting the distribution of dystrophin and PM integral components such as members

of the dystrophin-associated glycoprotein complex.

***In vivo* CHC depletion induces disorganization of the contractile apparatus and detachment between the sarcolemma and myofibrils.**

We performed electron microscopy analysis of CHC-depleted muscle at day 18 PI (without phenotype upon HE analysis) and at day 25 PI (peak of degeneration). At day 18 PI, focal regions displayed sarcomere abnormalities such as Z-band streaming and disorganization of the contractile apparatus especially in the I-band region (Fig. 10, A-C). It is of interest that these focal sarcomeric disorganizations were usually found adjoining the sarcolemma. In addition, at day 18 PI, occasional detachments of the sarcomeric apparatus from the sarcolemma were visible (Fig. 10, E-G). Both these phenotypes were consistent with the phenotypes produced upon clathrin depletion on isolated fibers. Of note, the T-tubule and sarcoplasmic reticulum system did not seem affected in the disorganized regions (Fig. 10 D). At day 25 PI, both transverse (Fig. 10, H-J) and longitudinal sections (Fig. 10, K and L) presented a particularly exacerbated detachment phenotype, with fibers displaying large areas of the sarcolemma which were no longer associated with sub-sarcolemmal sarcomeres.

Discussion

In this study, we identify a new role for clathrin heavy chain in costamere formation and maintenance. Our results raise the possibility that CHC, AP2 and DNM2 contribute, early during myotube differentiation, to the formation of the costameric scaffold that is necessary for anchoring the actin cytoskeleton through recruitment of actin binding protein Hip1R and actin cross-linking protein α -actinin. Consequently, depletion of CHC, by perturbing the actin cytoskeleton and redistributing α -actinin, induces a strong effect on the formation and maintenance of the contractile apparatus. The possibility that we are observing a defective α -

actinin trafficking inside the cell towards the PM is improbable because α -actinin is not a membrane associated protein, does not co-localize with clathrin in intracellular compartments and because AP1 and AP3 knock-down does not induce an α -actinin redistribution phenotype. In fact, our results clearly point out towards a defect in the attachment of α -actinin to the cytoplasmic side of the PM and raise the possibility that clathrin is involved in actin scaffolding thereby performing a task similar to what has been described for bacterial uptake, adherens junctions and immune synapses (Bonazzi et al., 2012; Bonazzi et al., 2011; Calabia-Linares et al., 2011). In addition, our results demonstrate a predominant role for AP2 in CHC targeting to the PM of differentiated myotubes.

At the plasma membrane, clathrin is present at classical coated pits and more large coated plaques, forming two distinct endocytically active structures (Saffarian et al., 2009). The assembly of clathrin in plaques has been previously observed in several cell types including muscle cells, but their physiological importance has proven controversial (Akisaka et al., 2003; Bellve et al., 2006; De Deyne et al., 1998; Pumplin, 1989; Pumplin and Bloch, 1990; Saffarian et al., 2009) and their functional role is still not clear. They were thought to represent artifacts of cell adhesion of cultured cells. However, large clathrin lattices have also been observed in non adherent floating cultured adipocytes (Bellve et al., 2006) and we observe them on non adherent regions such as the top or the sides of myotubes (Fig. 2 D and Fig S4 B). Because one putative role for clathrin-coated membrane domains is to maintain the stable attachment to the substrate, we suggest that clathrin plays the same role at the costamere, which is the muscle-specific adhesion site.

These large clathrin coated plaques are also endocytically active and invaginated pits are observed at their edges. Several groups have reported the implication of both CHC and DNM2 in focal adhesion turnover (Ezratty et al., 2009; Ezratty et al., 2005). Therefore one may hypothesize that this protein complex responsible for formation and maintenance of the

costameres is also involved in the turnover of these structures by endocytosis. In addition, we cannot exclude that the defects we are observing both *in vitro* and *in vivo* are at least partially due to defective endocytosis of costamere components. However, because i) plaques in myotubes contain much more polymerized clathrin than would be required to form a single vesicle, ii) adhesion inhibits clathrin-mediated endocytosis (Batchelder and Yarar, 2010) and iii) endocytosis is reduced during differentiation (Kaisto et al., 1999), the defects we are observing are probably not mainly due to endocytosis defects but could relate to the scaffolding role of CHC and DNM2 for the actin cytoskeleton (Schafer et al., 2002).

Our experiments show that these large clathrin-coated structures are intimately associated with the actin cytoskeleton and actin-binding proteins such as α -actinin. Because vinculin and talin were not detected in our co-immunoprecipitation experiments and because vinculin did not localize on clathrin-coated plaques (our unpublished data), we believe that the coated domain corresponds to a different entity from classical focal adhesion contacts. The detection of clathrin plaques *in situ* in adult skeletal muscle is of great interest but remains technically challenging at the moment. It is of interest that costameres have been recurrently described as sub-sarcolemmal dense plaques (Chiesi et al., 1981; Hijikata et al., 2003; Hijikata et al., 2008) and therefore it is tempting to speculate that part of the density which characterizes these structures could be attributed to clathrin plaques.

Early during *in vivo* clathrin depletion appear signs which point towards defects in costamere function. The first phenotype attributed to clathrin depletion is the presence of small detachments between the sarcolemma and the underlying myofibrils. CHC knock-down induces sarcomeric disorganization with streaming of the Z-line which leads to a drastic and rapid muscle degeneration. Our experiments confirm the results obtained on differentiated myotubes and again point towards a role for CHC in the formation of stable plasma membrane compartments which contributes to the attachment of sarcomeric lattices. Our

results also point towards differences between three different costamere complexes. The first is centered around the DGC complex, dystrophin, ankyrins and plectin (Ayalon et al., 2008; Randazzo et al., 2013), the second is centered around integrins, vinculin, α -actinin and talin (Belkin et al., 1986; Pardo et al., 1983a; Pardo et al., 1983b) and the third around integrins, α -actinin and clathrin plaques (this study) while all contain actin. Also the strong maximal force-loss seen in the clathrin depleted muscle before any significant histological sign testifies of the vital role played by clathrin. The fact that more than 40% of the direct maximal force is lost before any change in total muscle mass or before the appearance of histological changes at the photon microscopy level could be attributed to the loss of both longitudinal and lateral force transmission in these myofibers and can be a direct consequence of both the large areas where disorganization of sarcomeres is evident and an uncoupling between the sarcolemma and the underlying contractile apparatus. While the contribution of a trafficking dysfunction cannot be excluded, we believe that clathrin plays a structural role at the plasma membrane and that first appear signs related to this structural function.

Autosomal dominant Centronuclear myopathy (CNM) is a rare neuromuscular disorder due to mutations in the *DNM2* gene (Bitoun et al., 2005). The results reported here and previous data suggest that impairment of CHC and DNM2 function in the formation and maintenance of the contractile apparatus via costamere organization may be relevant in CNM pathophysiology. Similarly to clathrin, DNM2 is mainly localized at the PM and the Z-line where it colocalizes with α -actinin in both mouse and human skeletal muscle (Cowling et al., 2011), and we show that DNM2 is required for proper α -actinin organization. Recently a CNM-related *DNM2* mutation was associated with an accumulation of electron-dense material located between disorganized myofibrils as well as DNM2 and CHC mislocalization within the atrophied myofiber (Kierdaszuk et al., 2013). In addition, overexpression of *DNM2* mutants in mouse skeletal muscle leads to α -actinin structural defects with prominent misalignment of the Z-

line *in vivo*, which may explain at least in part the reduced specific maximal force observed in these mice (Cowling et al., 2011). All these data associated with the undeniable role of DNM2 in regulation of actin cytoskeleton dynamics and focal adhesion maintenance (Ezratty et al., 2009; Ezratty et al., 2005) highlight a new putative pathomechanism of *DNM2*-related CNM which merits further investigation.

Overall, our experiments clearly demonstrate the involvement of the clathrin heavy chain, the AP2 adaptor protein, Hip1R and DNM2 in the formation of structures which, through association with α -actinin and actin filaments, allow the organization of the contractile apparatus in close association with the sarcolemma. The present study sheds light on a long-standing question, namely, the role of clathrin in skeletal muscle sarcomeric structure organization. Altogether, our results highlight an unconventional, novel role for clathrin in striated muscle which may be relevant to muscle physiology and whose dysfunction could be associated with the physiopathology of centronuclear myopathy linked to *DNM2* mutations.

Materials and Methods

Antibodies

Anti-CHC mouse monoclonals X22 and TD.1, rabbit polyclonal against clathrin light chains (anti-consensus) and mouse monoclonal AP6 against AP2 (alpha subunit) antibodies used in this study were produced by Dr. Brodsky's laboratory. The rabbit polyclonal antibody against CHC is from Abcam. For CHC detection in immunohistochemistry, immunocytochemistry and electron microscopy we used the mouse monoclonal X22 antibody. The rabbit polyclonal was used for double immunofluorescent labelling. Both X22 and the rabbit polyclonal anti-CHC completely co-localize. For Western blot experiments, both mouse monoclonal TD.1 and rabbit polyclonal (Abcam) antibodies were used. Other commercial sources of antibodies were as follows: mouse monoclonals against α -actinin 2, smooth muscle-actin, γ -actin, vinculin and talin (SIGMA), mouse monoclonal SA4 against AP3 (delta subunit DSHB), rabbit polyclonal against AP1 (gamma subunit), rabbit polyclonal anti-DNM2, rabbit polyclonal anti-integrin β 5, rabbit polyclonal anti-GAPDH and rabbit polyclonal anti-CD11b (Abcam), goat polyclonal anti-caveolin3 (Santa Cruz), rabbit polyclonal against dystrophin (Novocastra), rabbit polyclonal against Hip1R (Millipore). Secondary antibodies for immunofluorescence were from Life Technologies (Alexa-488, Alexa-568 and Alexa-633 conjugates). Secondary antibodies coupled to horseradish peroxidase were from Jackson Laboratories.

Muscle tissue lysate

Lysates of whole tissue were prepared from freshly dissected mouse skeletal muscle. Tissue was homogenized by dounce in lysis buffer (1g/3ml 50 mM Tris-HCl pH 7.5, 0.15 M NaCl, 1 mM EDTA, 1% NP40 and 1 protease inhibitor cocktail tablet (1/10ml buffer, Roche)).

Homogenate was centrifuged (10 min, 14,000 x g) and the pellet discarded to obtain a post-nuclear supernatant. Protein concentration of the lysate was determined by Bradford assay (Biorad).

Immunoblot analysis

Protein samples were separated by electrophoresis (4-12% bis-acrylamide gel, Life Technologies), then electrophoretically transferred to 0.45 μ m nitrocellulose membranes (Life Technologies) and labelled with primary antibodies and secondary antibodies coupled to horseradish peroxidase. The presence of proteins in samples was detected using Western Lightning Chemiluminescence Reagent (GE Healthcare Life Sciences). Quantification was performed using Quantity One software (Biorad).

Subcellular fractionation

3T3 fibroblasts, C2C12 myoblasts and differentiated C2C12 myotubes were homogenized in HES buffer at pH 7.4 (20 mM Hepes, 1 mM EDTA, 225 mM sucrose) by passing 10 \times through a 27-gauge needle. Homogenates were centrifuged (20 min, 19,000 x g). The resulting pellets were resuspended in HES buffer, layered on top of a 1.12 M sucrose cushion and centrifuged (101,000 x g) for 60 min at 4°C. The PM fraction was collected from the interface of the two solutions. Centrifugation of the initial supernatant (212,000 x g) for 60 min at 4°C allowed separation of the intracellular membrane fraction (pellet) and cytosol (supernatant).

Immunoprecipitation

For 3T3 and C2C12 subcellular fractions (PM and cytosol), 250 μ l of each fraction were diluted to half using lysis buffer (50 mM Tris-HCl pH 7.5, 0.15 M NaCl, 1 mM EDTA, 1% NP40 and 1 protease inhibitor cocktail tablet (1/10ml buffer, Roche)). Mouse skeletal muscle

tissue lysate (post-nuclear supernatant) was diluted to 3 mg/ml in lysis buffer, precleared with 30 μ l washed Protein-G-Sepharose (PGS 4 fast flow, GE Healthcare Life Sciences) per 400 μ l of diluted lysate, and incubated with 20 μ g of specific antibody overnight (4°C). Then 30 μ l washed PGS was added and incubated for 1 hr at 4°C. Pelleted PGS was taken up in sample buffer and subjected to electrophoresis and immunoblotting. For all immunoprecipitation experiments, HRP-conjugated rabbit and mouse IgG TrueBlot secondary antibodies (eBioscience) were used.

Dissociated fiber cultures

Myofibers were isolated from the dissected *flexor digitorum brevis* muscle of 6-wk old mice by digestion with collagenase 1a (Worthington Biochemical Corp.) and mechanical dissociation. Isolated fibers were cultured (37°C, 5% CO₂) in Dulbecco's Modified Eagle medium (DMEM) with high glucose, 1% horse serum (GIBCO), and penicillin/streptomycin on coverslips coated with Matrigel (Becton Dickinson Labware, Bedford, MA). Myofibers were transduced with 70 μ l of AAV vector (7.10^{10} viral genomes /well) for 12 days.

Mouse myoblast cultures

Primary skeletal muscle cells and C2C12 cells were maintained in tissue culture dishes coated with rat-tail collagen in basal medium with 20% fetal bovine serum (FBS), 50 U/ml penicillin, 50 mg/ml streptomycin (growth medium). Differentiation was induced when cells were ~80% confluent by culturing in differentiation medium (basal medium with 2% horse serum). For siRNA treatment, cells (differentiated for either 4 or 6 days) were transfected using 10 nM siRNA and transfection reagent (JetPrime, PolyPlus Transfection) according to manufacturer's instructions. Targeting and GAPDH-specific control siRNAs were synthesized

by Eurogentec. For CHC, sequences targeted were similar to those used in (Ezratty et al., 2009).

5'-AACAUUGGCUUCAGUACCUUG-3' for CHC-1, 5'-
AAUGGAUCUCUUUGAAUACGG-3' for CHC-2, 5'-
GAGCAUGUGCACGCUGGCCAGCU-3' for the α subunit of AP2. The GAPDH siRNA
sequence was 5'-AAAGUUGUCAUGGAUGACC-3' and was used as a negative control in
all siRNA experiments. The sequence of the siRNA specific for the γ subunit of the AP-1
complex (γ -adaptin) was 5'-AGCUAUGAAUGAUUAUUA-3', which is similar to the
sequence used in (Dugast et al., 2005) (Braun et al., 2007). The sequence of the siRNA
specific for the δ subunit of the AP-3 complex (δ -adaptin) was 5'-
CAUCAAGAUCAUCAAGCUG-3' which is similar to the sequence used in (Braun et al.,
2007). The sequence of the siRNA specific for DNMT2 was 5'-
ACCUACAUCAGGGAGCGAGAA-3'. For CHC depletion, cells were transfected twice for
48h. For all other siRNA constructs, cells were transfected once for 48 to 72h.

Immunofluorescence microscopy

Adult mouse skeletal muscle was embedded in Tissue-Tek OCT compound (Miles Inc.),
frozen, and stored at -80°C. Cryosections (10 μ m thick) were fixed (15 min, 4%
paraformaldehyde in PBS or 10 min, 95% ethanol (Fig. 1D) at room temperature (RT)),
permeabilized (5 min, 0.5% Triton X-100 in PBS, RT) and blocked (30 min, PBS with 0.1%
Triton X-100, 5% bovine serum albumin (BSA)). Sections were incubated with primary
antibodies (overnight, 4°C, in PBS with 0.1% Triton X-100, 5% BSA) and washed in PBS
with 0.1% Triton X-100. Sections were then incubated with secondary antibodies (60 min,
RT), washed in PBS with 0.1% Triton X-100, and mounted with anti-fading solution
(DABCO) containing DAPI (300 nM). For double labelling, the two primary antibodies (from

different species) or the two secondary antibodies were added simultaneously at the appropriate step. Secondary antibodies were labelled with either Alexa-488 or Alexa-568.

For myofibers isolated from mouse muscle, immunolabelling was performed directly in the 24-well culture plate. Cells were fixed (15 min, 4% paraformaldehyde in PBS, RT), permeabilized (10 min in PBS with 0.5% Triton X-100), and immunolabelled using the same procedure described for mouse muscle sections.

For mouse cells grown on coverslips, cells were washed in warm PBS, fixed in paraformaldehyde (4% in PBS, 15 min), then washed in PBS, permeabilized (10 min, 0.5% Triton X-100 in PBS) and blocked in blocking solution (5% BSA in PBS with 0.1% Triton X-100, 30 min). Antibody labelling was performed by addition of 200 μ l blocking solution with primary or secondary antibodies (1-5 μ g/ml) and washing with PBS with 0.1% Triton X-100. Samples were mounted in Vectashield containing DAPI (Vector Laboratories).

Muscle sections, mouse cells, and myofibers from mouse muscle were analyzed by confocal laser scanning microscopy using either an inverted LEICA SP2 operating system equipped with HCX Plan-Apo CS 40x, 1.20 NA and 63x, 1.40 NA oil immersion objective lenses. Images presented in Figs. 1 A; Fig. S3, and Fig. S4, were acquired using an upright OLYMPUS FV-1000 confocal laser scanning microscope equipped with UPlanS-Apo 60x, 1.35 NA oil immersion objective lenses. DAPI, Alexa-488 and Alexa-568 fluorescence was sequentially excited using lasers with wavelengths of 405 for DAPI, 488 (LEICA) or 473 (OLYMPUS) for Alexa-488 and 543 nm for Alexa-568. Z-series from the top to the bottom of fibers were sequentially collected for each channel with a step of 0.9-1 μ m between each frame. Imaging was performed at room temperature using Leica Type F immersion oil. Images (1024 \times 1024 pixels) were saved as TIFF files in OLYMPUS FV-1000 software and Leica Confocal Software, and input levels were adjusted in Adobe Photoshop. Image

quantification was performed using National Institutes of Health's ImageJ (<http://rsb.info.nih.gov/ij>).

Image analysis

Striated surface measurement.

The α -actinin striated surface per myotube was quantified by manually drawing a region of interest in ImageJ around the striated or total cell surface in a single Z-stack from the middle of the cell. The striated surface was obtained by performing the ratio between striated vs total myotube surface.

Alpha-actinin aggregate size analysis.

The “Analyze particles” ImageJ plugin (version 1.46) was used to count intracellular particles on binary confocal images of C2C12 myotubes in a single image from the middle of the cell and to automatically measure the number and area of these particles.

shRNA constructs.

The siRNA against CHC (CHC-1) used for *in vitro* transfection experiments was synthesized to be directly cloned in pSUPER under the control of the H1 promoter. ShRNA consisted of a 21-nt sense sequence followed by a 9-nt loop (TTCAAGAGA), a 19-nt reverse sequence, and a RNA pol III terminator (TTTTT). The H1 cassette was then introduced into an AAV1-based vector between the two ITRs using the blunted *SpeI* and *Sall* sites on pSUPER-shRNA plasmid and *XbaI* site on the pSMD2 AAV2 vector backbones, which were type1 pseudotyped (Généthon, Evry, France).

Virus production and titration.

AAV2/1 pseudotyped vectors were prepared by transfection in 293 cells as described earlier (Riviere et al., 2006) using the pSMD2-shRNA plasmid, the pXX6 plasmid (Généthon) coding for the adenoviral sequences essential for AAV production, and the pRepCAp plasmid (Généthon) coding for AAV1 capsid. Vector particles were purified on iodixanol gradients and concentrated on Amicon Ultra-15 100K columns. The final viral preparations were kept in PBS solution at -80°C . The particle titer (number of viral genomes) was determined by quantitative PCR. Titres for AAV shCHC were 4.10^{12} vector genomes (vg)/ ml.

***In vivo* gene transfer**

Experiments were performed on adult 8 week old C57/Bl6 mice. Anaesthesia was achieved with a mix of 100 mg/kg ketamine and 10 mg/kg xylazine or using isoflurane. Two intramuscular injections (40 μl /TA), 24 h apart of AAV-shCHC was carried out in TA of the right hindlimb; the contralateral muscles were injected using the same procedure with control AAV vector (AAV-CTRL) which expresses the muSEAP protein (murine secreted embryonic alkaline phosphatase). Mice were sacrificed at different intervals following the injection.

Gene expression analysis

Total RNA was prepared from 400 μm cryostat sections of TA using Nucleospin RNAII kit (Macherey-Nagel). Complementary DNA generated with Invitrogen Superscript II Plus reverse transcriptase kit (Life Technologies) was analyzed by real-time qPCR performed on Opticon2 system (Bio-Rad) using iTaq SyberGreen Supermix with ROX (Bio-Rad). In all samples, we quantified transcript of the *PO* gene encoding mouse acidic ribosomal phosphoprotein ubiquitously expressed as endogenous RNA control and each sample was normalized on the basis of its PO content. Primers used for CHC: Forward Exon6: 5'-GAGTCAACAGAAAGGGACA-3' and Reverse Exon7: 5'-

CATTCTCAGAGCCAAGTCAG-3'. Primers used for PO: Forward: 5'-CTCCAAGCAGATGCAGCAGA-3' and Reverse: 5'-ATAGCCTTGCGCATCATGGT-3'.

Histomorphological and ultrastructural analyses

Samples were frozen in liquid nitrogen-cooled isopentane. Transverse sections of tibialis anterior (TA) muscle (8 μ m thick) were stained with HE by standard methods. Light microscopy was performed using an upright microscope (DMR, Leica) and 40 \times NA 0.85 HCX Plan Apo objective (Leica). Images were captured using a monochrome camera (DS-Ri1, Nikon) and NIS-Elements BR software (Nikon). For all imaging, exposure settings were identical between compared samples and viewed at room temperature.

For morphological electron microscopy, muscles were fixed by intra-aortic perfusion with 2% paraformaldehyde, 2% glutaraldehyde in 0.1M phosphate buffer (pH 7.4). Tibialis anterior samples were postfixed with 2% OsO₄, in 0.1 M phosphate buffer (pH 7.4) for 1 h then dehydrated in a graded series of acetone including a 1% uranyl acetate staining step in 70% acetone and finally embedded in epon resin. For pre-embedding immunogold clathrin labellings, mice were fixed with 2% paraformaldehyde and tibialis anterior longitudinal 100 μ m thick sections were cut with a vibratome. Then a standard free-floating immunocytochemical procedure was followed, using 0.1M saline phosphate buffer as diluent and rinsing liquids. After preincubation in 5% normal goat serum, 5% BSA, sections were overnight incubated at 4°C in 1/500 anti-clathrin antibody. A further 4 hour incubation in ultra-small gold conjugate of goat anti-rabbit IgG (1/20; Aurion, Netherlands) was followed by extensive washings, 10 min post-fixation in 2% glutaraldehyde and 0.70 nm gold beads were then silver enhanced (HQ silver; Nanoprobes, Stony Brook, NY). After 15 min post-fixation in 1% OsO₄, sections were dehydrated in graded acetone and embedded in Epon resin.

Thin (70 nm) sections were stained with uranyl acetate and lead citrate and in some cases (Fig.1F), a 1% tannic acid (1 min) staining step preceded uranyl staining. Observations were made on a Philips CM120 electron microscope (Philips, Eindhoven, The Netherlands) and images were recorded with a Morada digital camera (Olympus Soft Imaging Solutions GmbH, Münster, Germany).

Electron microscopy of unroofed myotubes

Adherent PM from myotubes cultured as described above were prepared for rapid-freeze, deep-etch electron microscopy as described previously (Heuser, 2000). Briefly, cells grown on small oriented pieces of glass coverslip were disrupted by sonication, fixed in paraformaldehyde, and labelled with anti-clathrin (X22), anti- α -actinin, anti-vinculin or anti- β 5 integrin and then 18-nm gold-conjugated anti-mouse or anti-rabbit antibody before flash freezing with a homemade liquid-helium cooled, copper cold block “slammer,” freeze dried *in vacuo* at -80°C for 15 min, replicated with a ~2 nm film of platinum vacuum-evaporated from an electron beam gun mounted 15-20° above the horizontal, and supported by vacuum-evaporating ~10 nm of pure carbon onto the platinum, using a standard carbon-arc supply mounted some 10° off the vertical. The resultant platinum replicas were floated off the glass by angled immersion into full strength (47%) hydrofluoric acid, washed several times by floatation on distilled water, picked up on 75 mesh formvar-coated EM grids, and imaged while mounted in a eucentric side-entry goniometer stage of a JEOL 1400 electron microscope equipped with a high-resolution digital camera.

Contractile properties

The isometric contractile properties of TA muscles were studied *in situ* on mice anesthetized with 60 mg/kg pentobarbital. The distal tendon of the TA muscle was attached to a lever arm

of a servomotor system (305B Dual-Mode Lever, Aurora Scientific). Muscles were directly stimulated by a bipolar silver electrode using a supramaximal (80 V) square wave pulse of 0.1 ms duration. Absolute maximal force (P_0) was measured during isometric contractions in response to electrical stimulation (frequency of 25–150 Hz; train of stimulation of 500 ms). All isometric contraction measurements were made at optimal muscle length (L_0) at which P_0 was obtained. TA muscles were weighted and specific maximal force (sP_0) was calculated by dividing P_0 by muscle weight.

Statistics and institutional guideline compliance

Statistical analysis was performed using Student's t-test except as otherwise stated. All animal studies were performed in compliance with the French guidelines.

Online supplemental material

Fig. S1 shows ultrastructural localization of clathrin using X22 monoclonal antibody in clathrin coated pits and plaques of mouse primary myotubes. Fig. S2 displays additional images of α -actinin localization and localization of β 5-Integrin in clathrin plaques of mouse primary myotubes. Fig. S3. shows efficiency of siRNA-mediated AP1, AP2, AP3, DNM2 and Hip1R depletion, the effect of siRNA-mediated AP2 depletion on CHC distribution and localization of Hip1R in control and CHC or AP2-depleted myotubes. Fig. S4 compares the effect of siRNA-mediated depletion of CHC, AP1, AP2, Hip1R and DNM2 on actin distribution, characterizes localization of DNM2 in control or CHC and AP2-depleted myotubes. Fig. S5 shows that *in vivo* CHC depletion induces a strong inflammatory response but has no effect of β -dystroglycan and dystrophin distribution.

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Abbreviations

AP adaptor protein

CCV clathrin-coated vesicle

CHC clathrin heavy chain

CLC clathrin light chain

CNM centronuclear myopathy

DNM2 dynamin 2

FDB flexor digitorum brevis muscle

Hip1R huntingtin-interacting protein 1related

PI post-injection

PM plasma membrane

TA tibialis anterior

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Figure 1

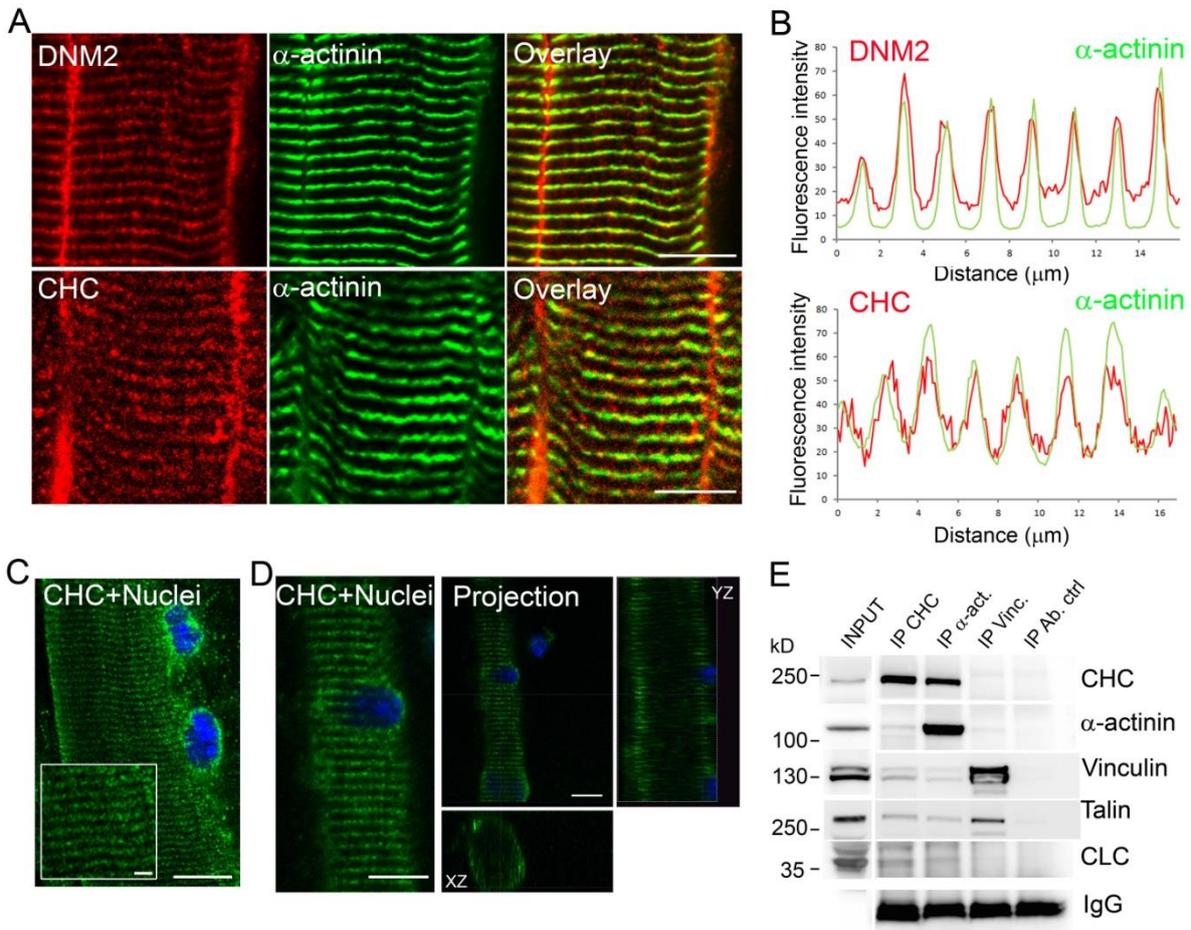


Fig. 1: CHC association with α -actinin2 in adult skeletal muscle. (A) Confocal sections of mouse skeletal muscle were processed for double immunofluorescent labelling of DNM2 (red) and α -actinin (green) or CHC (red) and α -actinin (green) (scale=10 μ m). (B) The fluorescence intensity as a function of distance on the fiber was plotted for eight successive striations, and was reported on the graph for each labelling. The green curve corresponds to the green immunolabelling, whereas the red curve corresponds to the red labelling. (C-D) CHC localization in mouse dissociated FDB fibers. (C) Confocal sections of fibers processed for immunofluorescent labelling with the monoclonal antibody against CHC (X22, green) (scale=10 μ m and 2 μ m for the inset). (D) XZ and YZ projections of confocal optical sections illustrated in D. (E) Immunoblot of proteins associated with CHC, α -actinin (isoform 2), vinculin or control (CTRL) immunoprecipitates from mouse muscle lysates and 1 to 5% lysate input. Bands for co-immunoprecipitated CHC, α -actinin (isoform 2), vinculin, talin or CLC (clathrin light chain) are indicated at the right.

Figure 2

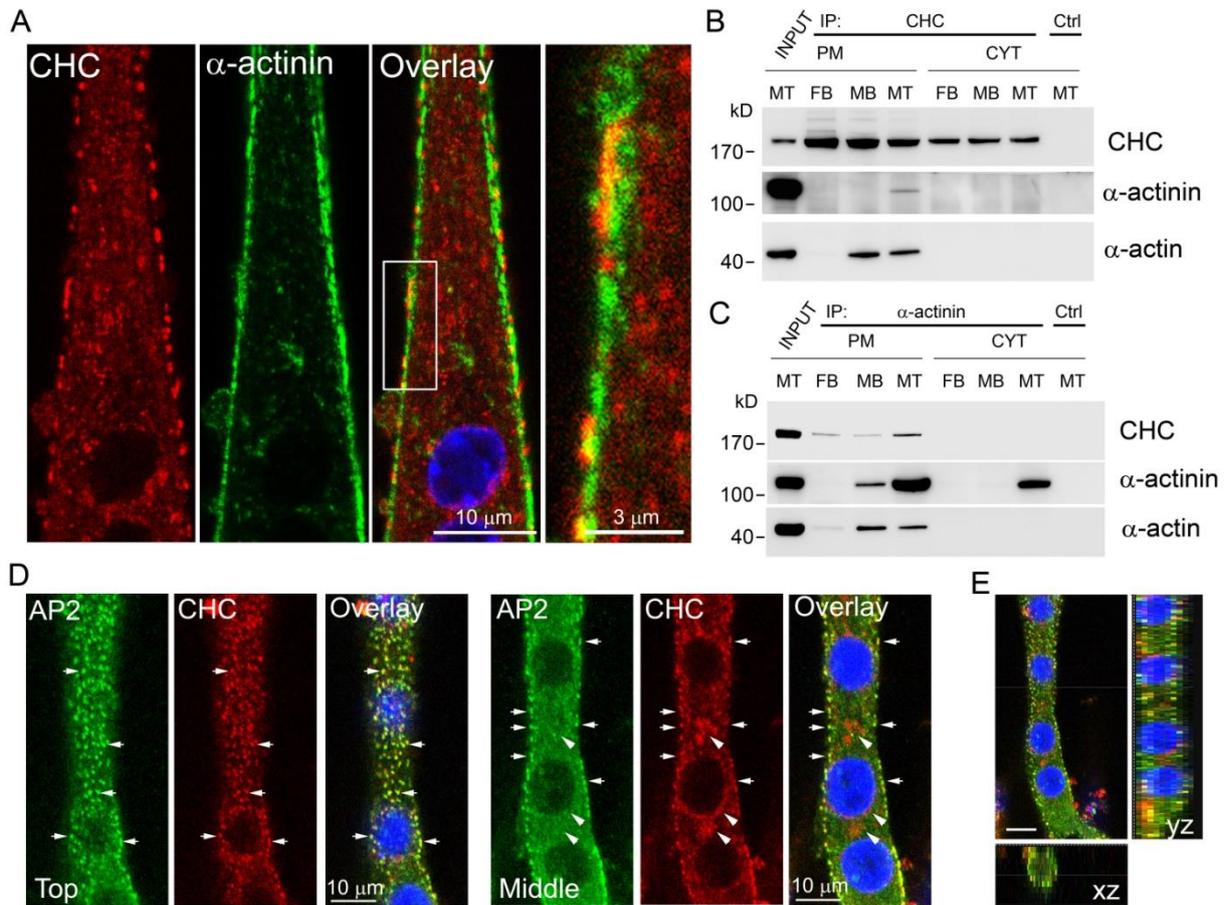


Fig. 2: CHC association with α -actinin2 and actin at the PM of primary myotubes. (A) Localization of α -actinin (green) and CHC (red) in mouse primary myotubes differentiated for 4 days (scale=20 μ m). (B) Immunoblot analysis of proteins associated with CHC or CTRL immunoprecipitates from subcellular fractions (plasma membrane, PM and cytosol, CYT) of 3T3 fibroblasts (FB), undifferentiated C2C12 (MB) and differentiated C2C12 cells (MT). (C) Immunoblot analysis of proteins associated with α -actinin or CTRL immunoprecipitates from subcellular fractions (PM and CYT) of 3T3 fibroblasts (FB), undifferentiated C2C12 (MB) and differentiated C2C12 cells (MT). (D) Confocal microscopy of AP2 (green) and CHC (red) in differentiated C2C12 skeletal muscle cells. The left panel displays images from the top of the myotube and the right panel displays images from the middle of the myotube. Arrows indicate large clusters of colocalization between AP2 and CHC and arrowheads indicate intracellular clusters near the nuclei positive for CHC and negative for AP2. (E) XZ and YZ projections of serial confocal sections are shown on the overlay (scale=10 μ m).

Figure 3

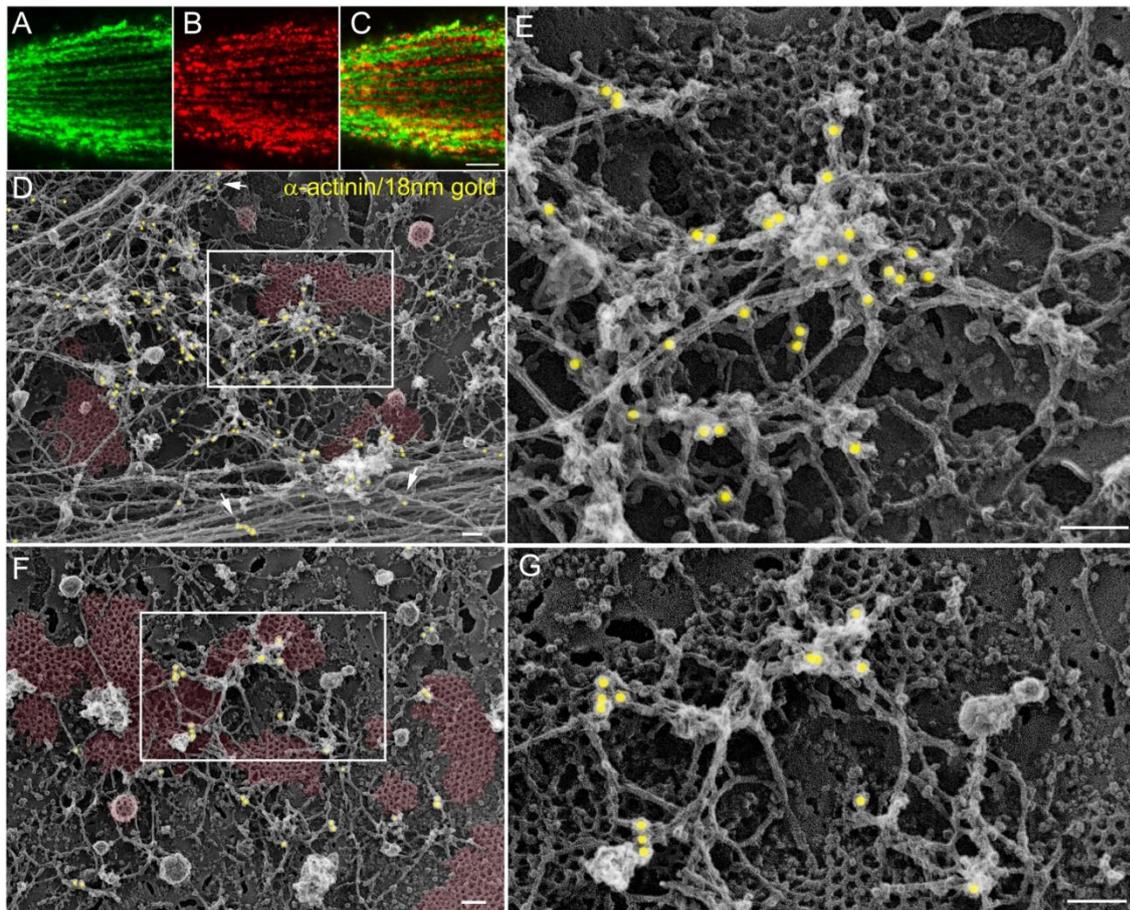


Fig. 3: α -actinin and actin are localized on large flat clathrin-coated plaques. (A–C) Immunofluorescent staining of α -actinin (A), CHC (B), and merged images (C) in untreated differentiated primary mouse myotubes visualized using confocal microscopy (scale=5 μ m). (D–F) Adherent plasmalemmal sheets prepared from control primary myotubes differentiated for 4 days and labelled with α -actinin antibodies followed by secondary antibodies conjugated to 18 nm colloidal-gold particles (gold particles are pseudo-colored yellow (D–G) while clathrin lattices are pseudo-colored pale red in D and F). The boxed region in D and F is magnified in E and G respectively. Note that α -actinin is found on both actin cables (arrows in D) and branched filaments laying on top of flat clathrin coated lattices (D–G) (scale=100 nm).

Figure 4

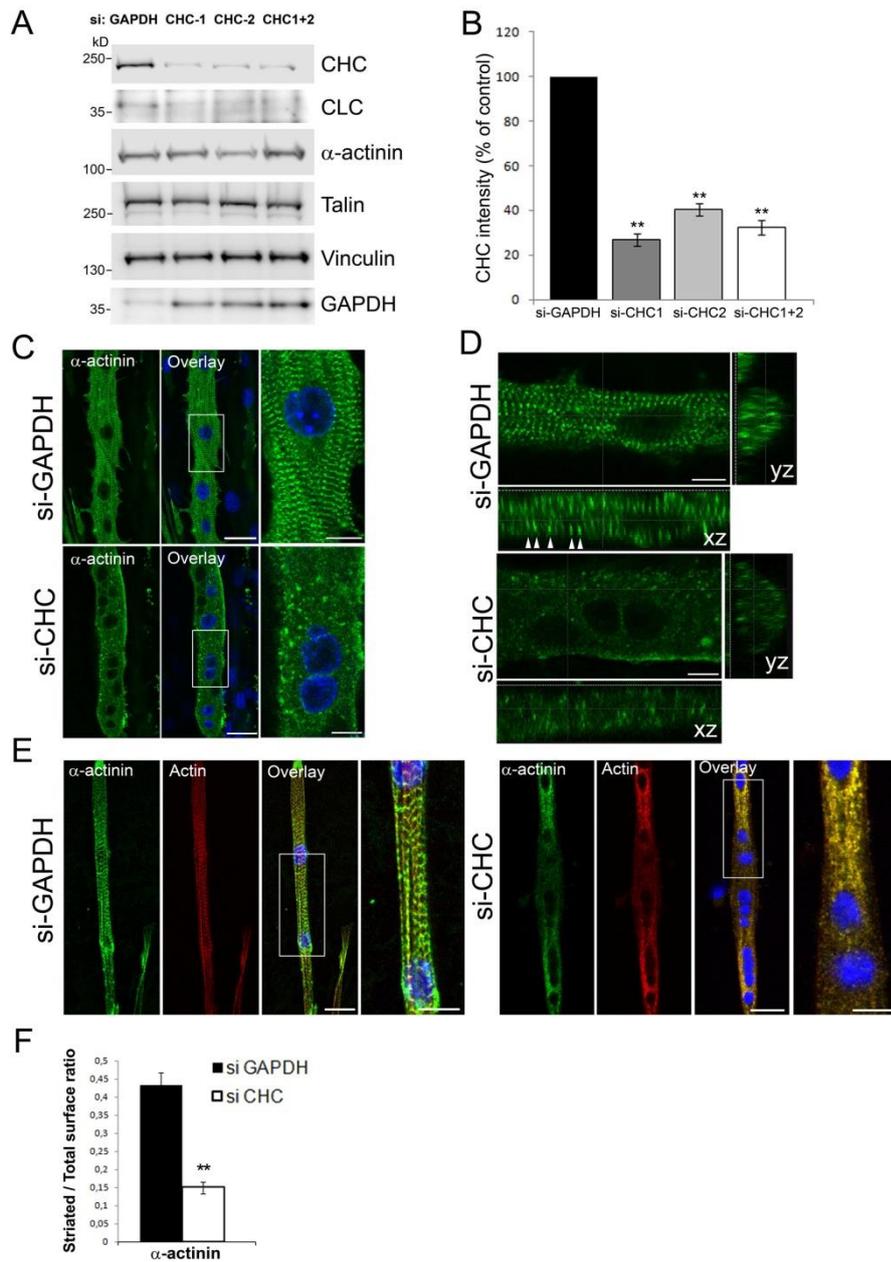


Fig. 4: CHC regulates formation and organization of sarcomeres in cultured myotubes.

(A) Differentiated C2C12 cells were treated with control GAPDH siRNA, two different siRNA targeting CHC or a mixture of both (CHC-1, CHC-2, CHC-1+2, indicated at the top) and cell lysates immunoblotted for proteins (indicated at the right). (B) CHC expression in cultured C2C12 myotubes treated with GAPDH or CHC siRNA. The graph depicts quantification of CHC band intensity ($n=8$ for CHC-1 and $n=4$ for CHC-2 and CHC1+2, data are presented as mean \pm s.e.m, $**P < 0.01$). (C) α -actinin staining in differentiated C2C12 skeletal muscle cells treated with control GAPDH siRNA or CHC siRNA. DNA staining (DAPI blue) identifies multinucleated myotubes (scale=30 μ m). The boxed region in the merged images is magnified in the insets (scale=10 μ m). (D) XZ and YZ projections of serial confocal sections (scale=10 μ m). (E) CHC depletion on cultured primary mouse myotubes. Immunofluorescent staining of α -actinin (green) and actin (red) in differentiated primary mouse myotubes treated with control GAPDH siRNA or CHC siRNA. DNA staining (DAPI blue) identifies differentiated, multinucleated myotubes (scale=20 μ m). The boxed region in the merged images is magnified in the insets (scale=10 μ m). (F) Quantification of striated vs non-striated surface in control (siRNA against GAPDH) or in myotubes treated with siRNA against CHC ($n=30$ to 60 myotubes, data are presented as mean \pm s.e.m, $**P < 0.01$).

Figure 5

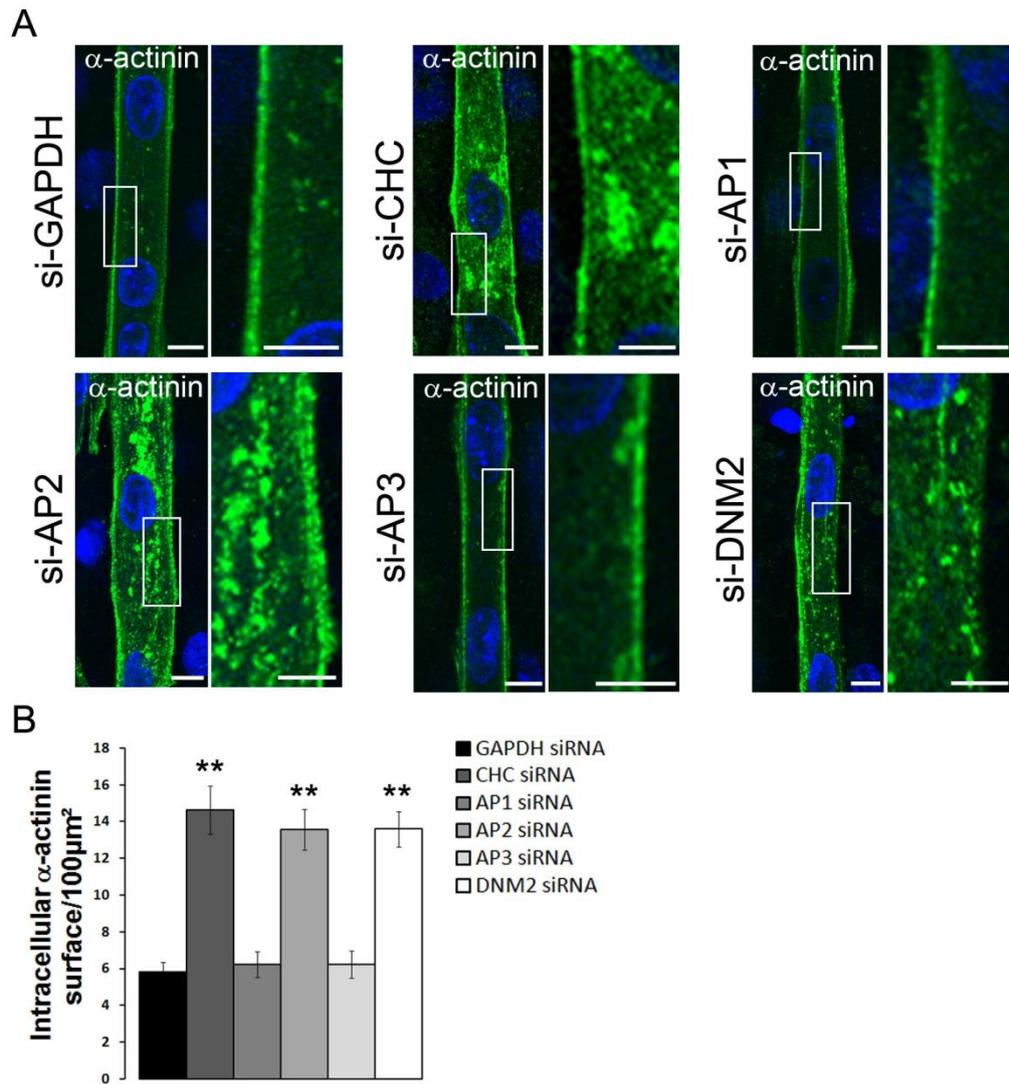


Fig. 5: CHC, AP2 and DNM2 but not AP1 or AP3 depletion in cultured mouse myotubes perturbs α -actinin distribution before striations appear. (A) Differentiated C2C12 myotubes were treated with siRNA targeting proteins (indicated at the left). Immunofluorescent staining of α -actinin in myotubes treated with control siRNA against GAPDH or siRNA targeting either CHC, AP1, AP2, AP3, or DNM2. DNA staining (DAPI blue) identifies differentiated, multinucleated myotubes (scale=10 μm). The boxed region in the merged images is magnified in the insets (scale=5 μm). (B) Quantification of α -actinin fluorescence surface as a function of total myotube surface in 4-day differentiated myotubes treated with siRNA against GAPDH, CHC, AP1, AP2, AP3 or DNM2 (n=30 to 50 myotubes, data are presented as mean \pm s.e.m, **P < 0.01).

Figure 6

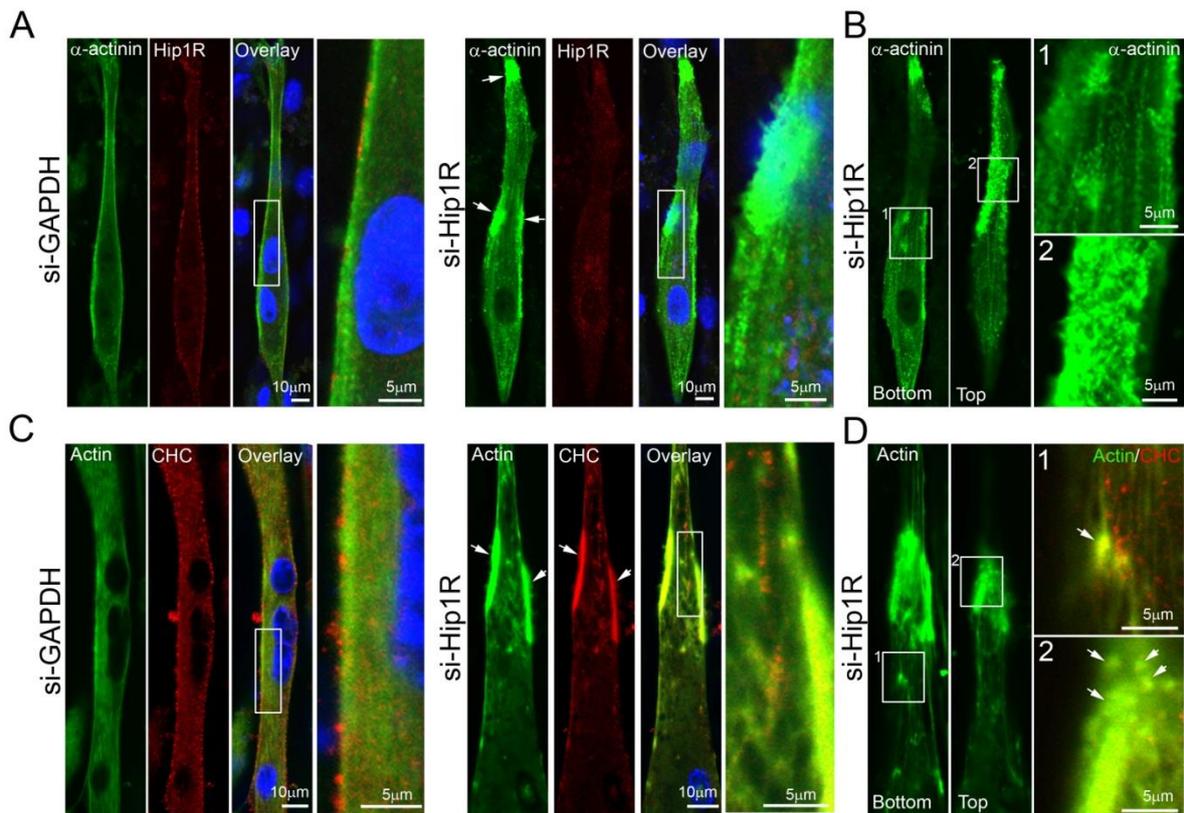


Fig. 6: Hip1R depletion in cultured myotubes stabilizes actin and α -actinin on clathrin-coated structures. (A) Differentiated C2C12 myotubes were treated with siRNA targeting proteins (indicated at the left). Immunofluorescent staining of α -actinin (green) and Hip1R (red) in myotubes treated with control siRNA against GAPDH or siRNA targeting Hip1R. DNA staining (DAPI blue) identifies differentiated, multinucleated myotubes (scale=10 μ m). Arrows indicate large α -actinin aggregates. The boxed region in the merged images is magnified in the insets (scale=5 μ m). (B) α -actinin localization in differentiated C2C12 skeletal muscle cells. The left panel displays an image from the bottom of the myotube and the right panel displays an image from the top of the myotube. The boxed regions in the images are magnified in the insets 1 and 2 (scale=5 μ m). (C) Differentiated C2C12 myotubes were treated with siRNA targeting proteins (indicated at the left). Immunofluorescent staining of actin and CHC in myotubes treated with control siRNA against GAPDH or siRNA targeting Hip1R. DNA staining (DAPI blue) identifies differentiated, multinucleated myotubes (scale=10 μ m). The boxed region in the merged images is magnified in the insets (scale=5 μ m). (D) Actin localization in differentiated C2C12 skeletal muscle cells. The left panel displays an image from the bottom of the myotube and the right panel displays an image from the top of the myotube. The boxed regions in the images are magnified in the insets (1 and 2) and include CHC staining (red) (scale=5 μ m). Arrows indicate large clusters of colocalization between actin and CHC.

Figure 7

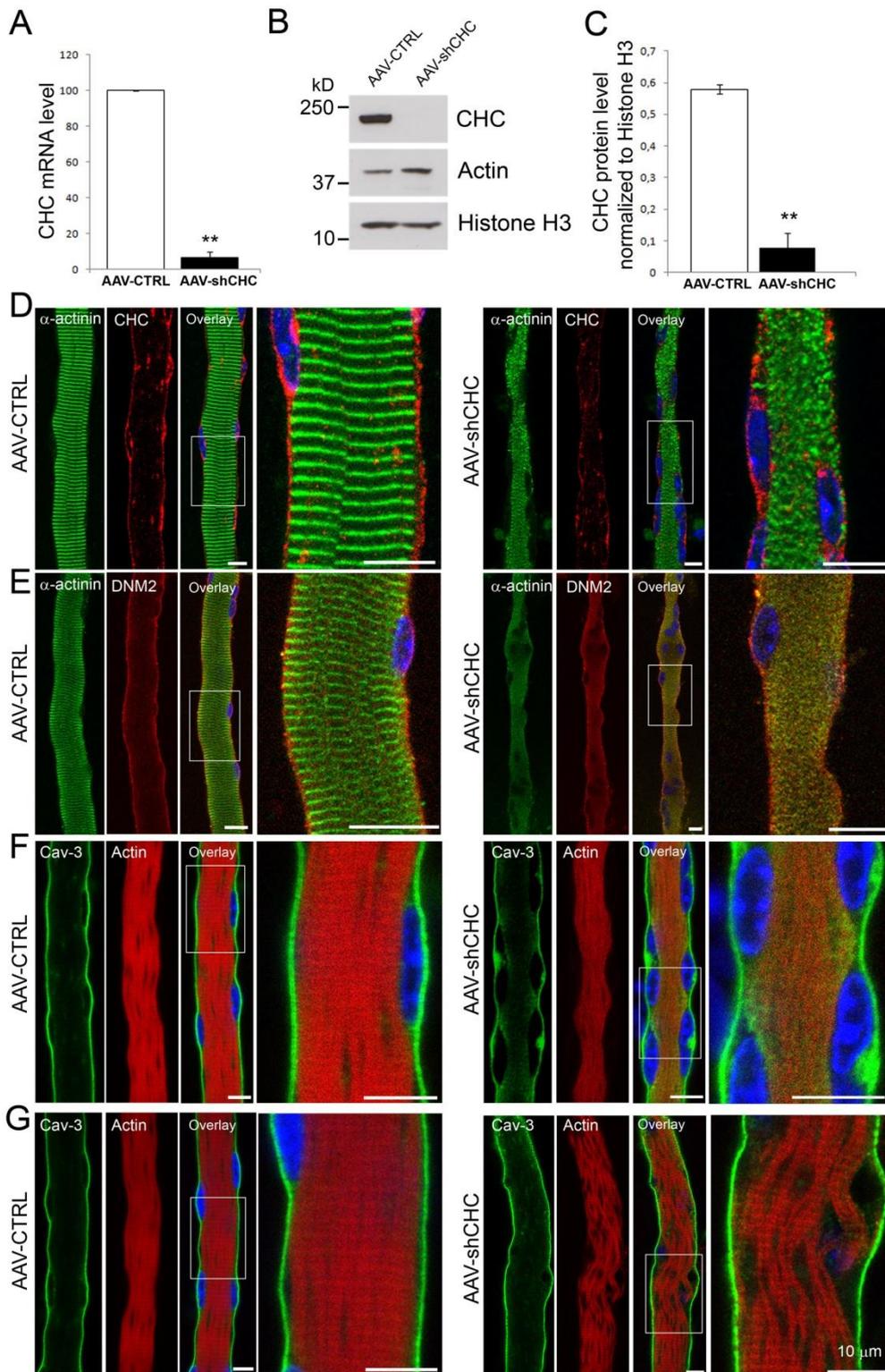


Fig. 7: CHC is required for sarcomere maintenance in adult fibers. (A) Quantitative RT-PCR and (B) Western blot analysis of CHC levels and proteins indicated at the right in FDB fibers infected with AAV-CTRL or AAV-shCHC for 12 days *in vitro*. (C) Quantification of Western blot CHC band intensity (n=3, data are presented as mean \pm s.e.m, **P < 0.01). (D-G) Confocal microscopy of dissociated FDB fibers infected with AAV-CTRL or AAV-shCHC for 12 days and co-stained with antibodies against either α -actinin and CHC (D), α -actinin and DNM2 (E), or F-actin (phalloidin staining) and Caveolin-3 (F and G).

Figure 8

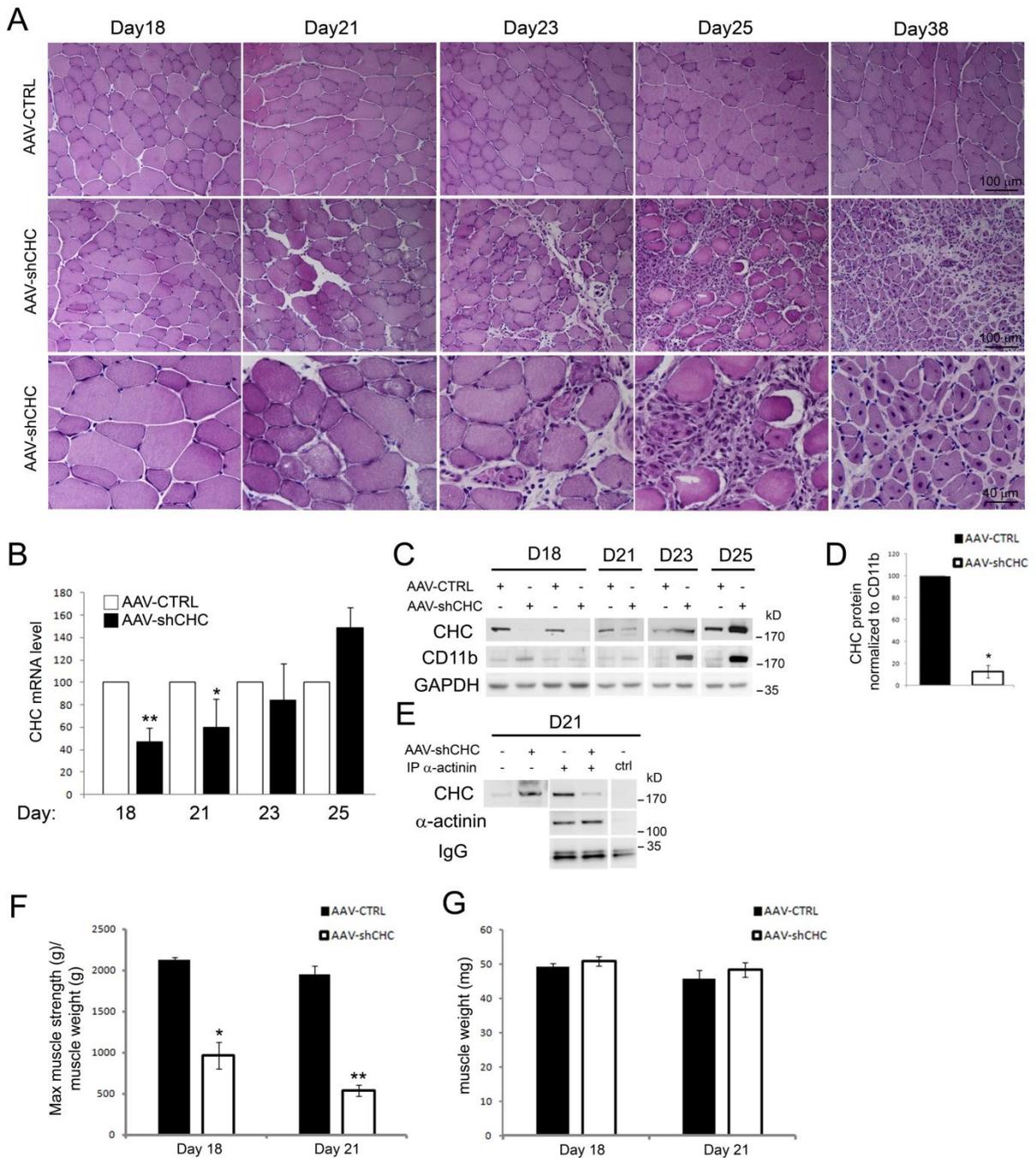


Fig. 8: CHC depletion using AAV-shCHC *in vivo* impairs force and causes muscle degeneration. (A) Hematoxylin and eosin staining of AAV-CTRL or AAV-shCHC injected muscle at day 18, 21, 23, 25, and 38 following virus injection). (B) RT-qPCR of CHC mRNA levels at different time points following AAV-CTRL or shCHC AAV injection. (n=2 to 9 mice), data are presented as mean \pm s.d., *P < 0.05, **P < 0.01. (C) CHC protein levels at different time points following AAV-CTRL or shCHC AAV injection. (D) Quantification of Western blot CHC band intensity (n=4 mice at day 18, data are presented as mean \pm s.e.m., *P < 0.05, Mann-Whitney U test). (E) Immunoblot analysis of proteins associated with α -actinin or CTRL immunoprecipitates of TA muscle lysates injected with AAV expressing a control construct or AAV expressing shCHC at 21 days following injection. (F-G) Measures of the specific maximal force (F) and of the muscle weight (G) of isolated TA muscle injected with AAV expressing a control construct or AAV expressing the shCHC construct (n=3 to 5 mice) at either 18 or 21 days following injection. Data are presented as means \pm s.e.m., *P < 0.05, **P < 0.01.

Figure 9

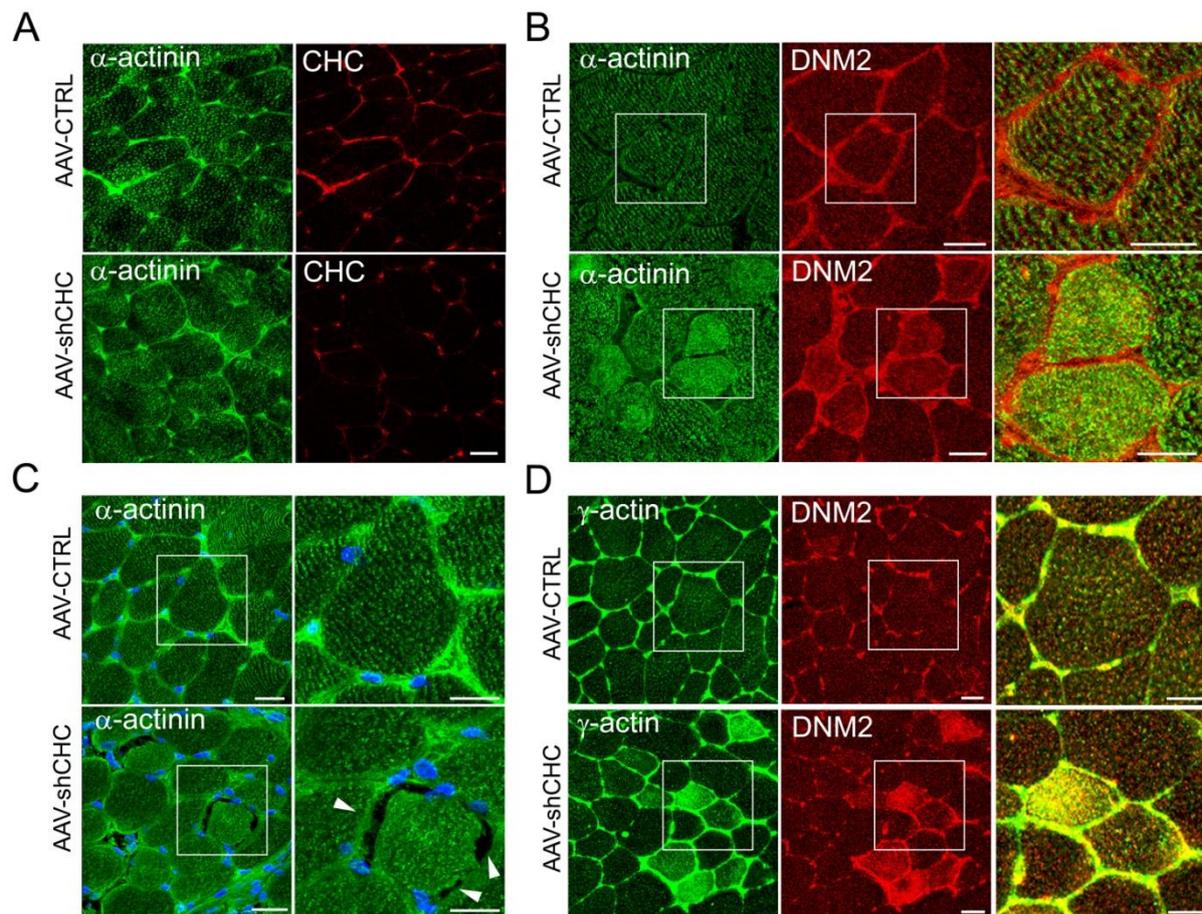


Fig. 9: α -actinin, γ -actin and DNM2 distribution is perturbed upon *in vivo* CHC depletion. (A-D) Confocal microscopy on transverse sections of skeletal muscle processed for double immunofluorescent labelling using antibodies against CHC (red) and α -actinin (green) at day 18 PI (A), or against α -actinin (green) and DNM2 (red) (B), α -actinin (green) and DAPI (blue) (C) and γ -actin (green) and DNM2 (red) (D) at day 21 PI (scale=30 μ m). The boxed region in the merged images is magnified in the insets (scale=20 μ m).

Figure 10

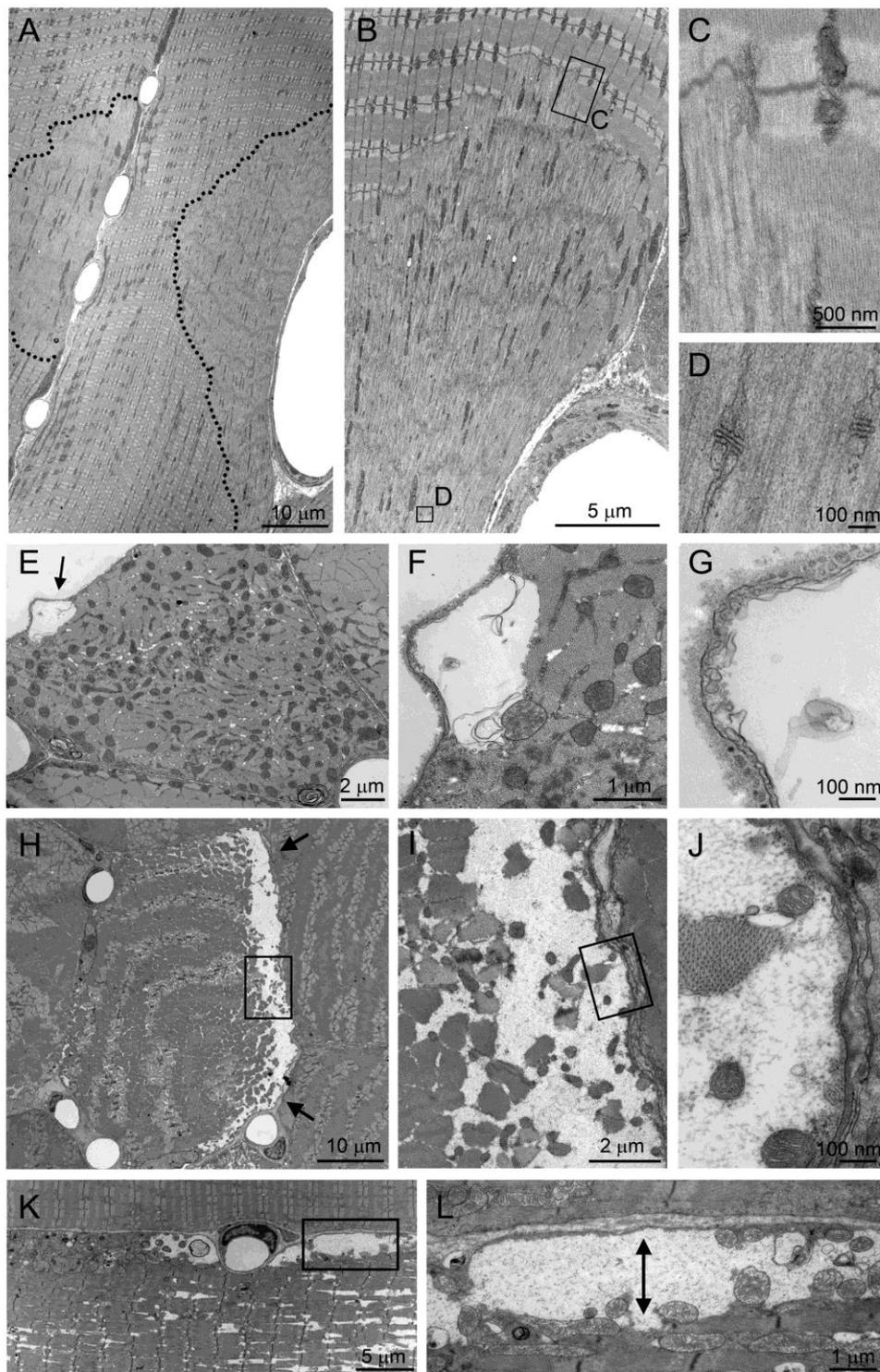


Fig. 10: Sarcomeric disorganization and detachment between sarcolemma and myofibrils upon *in vivo* CHC depletion. Transmission electron microscopy of longitudinal (A-D, K-L) and transverse (E-J) muscle sections from mice injected with AAV-shCHC constructs at day 18 (A-G) and day 25 PI (H-L). In A-D the sarcomeric apparatus is disorganized in focal regions adjoining the sarcolemma (within the dotted lines in A) and presents streaming and disassembly of the Z-line. C and D are insets of the respective boxed regions in B. In E-G, small detachments between the sarcolemma and the contractile apparatus are visible at day 18 PI. At day 25 PI, large detachments are seen in both transverse (H-J) and longitudinal orientation (K-L). Arrows in E, H and L indicate detachments.

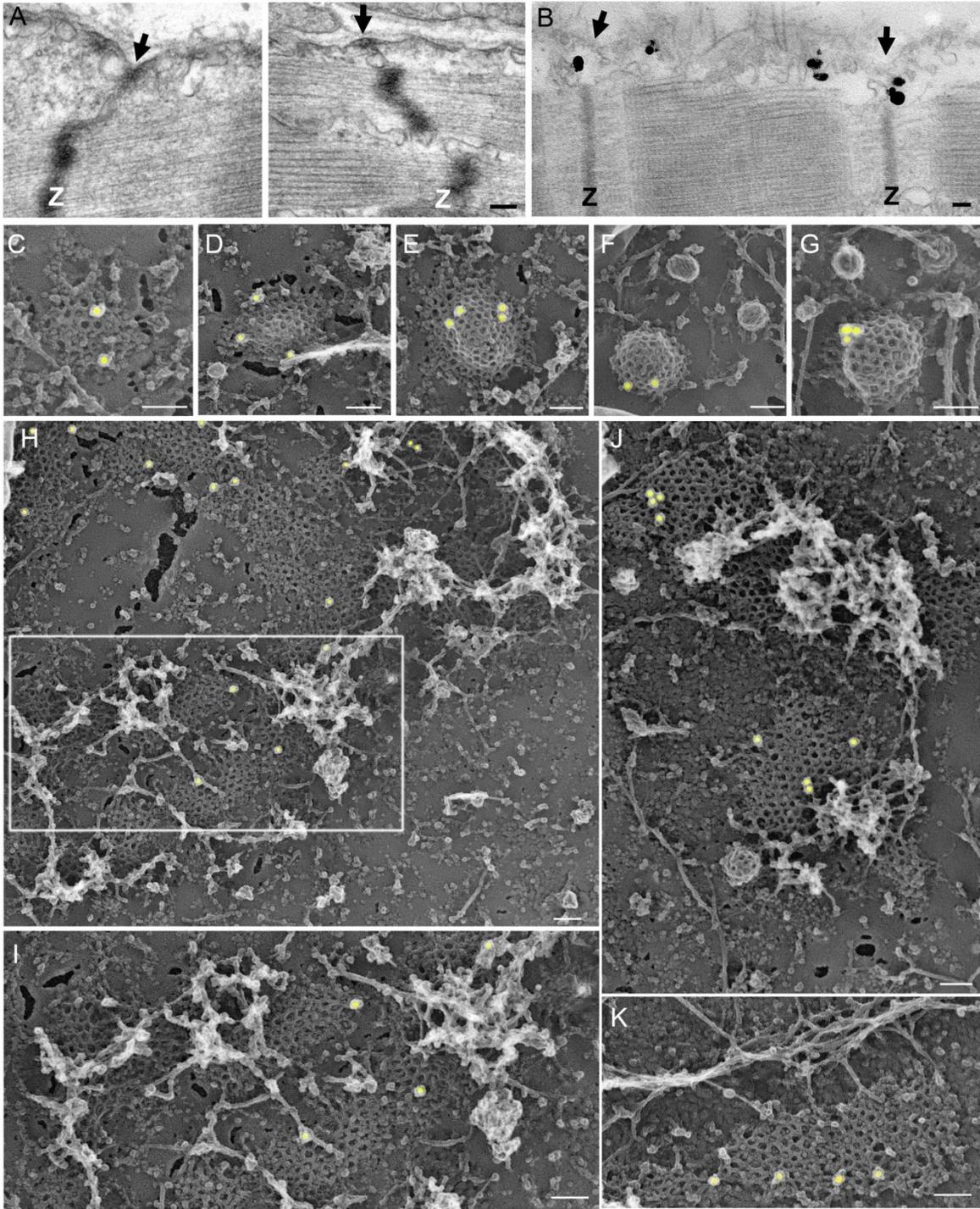


Figure S1: Ultrastructural localization of clathrin in adult skeletal muscle and mouse primary myotubes. (A) Thin (70 nm) longitudinal muscle sections were stained with 1% tannic acid followed by uranyl acetate and lead citrate. An arrow points at electron dense regions corresponding to costameres. (B) Thin longitudinal muscle sections immune-labelled with CHC polyclonal antibody and processed for EM. (C–K) Fixed PM sheets prepared from control primary myotubes were labelled with CHC antibodies, and then with secondary antibodies conjugated to 18-nm colloidal-gold particles. Representative freeze-etch images show the distribution of clathrin on small, flat or invaginated coated pits (C–G) or on large, flat, actin-associated coated plaques (H–K). (scales=100 nm)

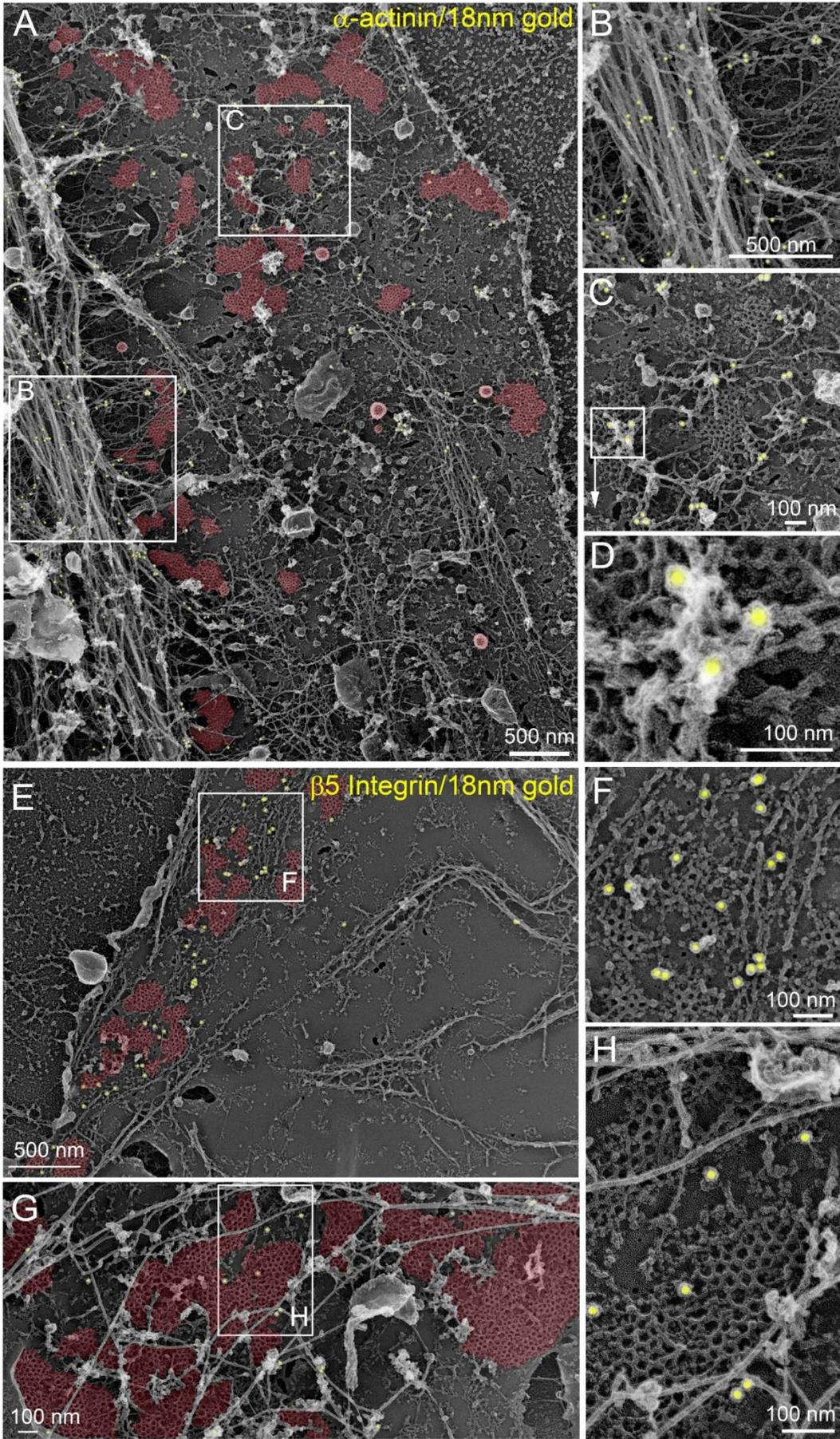


Figure S2: Additional images of α -actinin localization and localization of β 5-Integrin in clathrin plaques. (A–D) Adherent plasmalemmal sheets prepared from control primary mouse myotubes were labelled with α -actinin antibodies followed by secondary antibodies conjugated to 18 nm colloidal-gold particles (pseudo-colored yellow). Note that α -actinin is found on both the contractile apparatus (A and B) and on large and flat clathrin coated lattices (A, C, and D). (E–H) Fixed PM sheets prepared from control myotubes were labelled with β 5-integrin antibodies, and then with secondary antibodies conjugated to 18-nm colloidal-gold particles. Representative freeze-etch images show the spatial distribution of endogenous β 5-integrin enriched on large clathrin coated structures. In A and E clathrin lattices are pseudo-colored pale red.

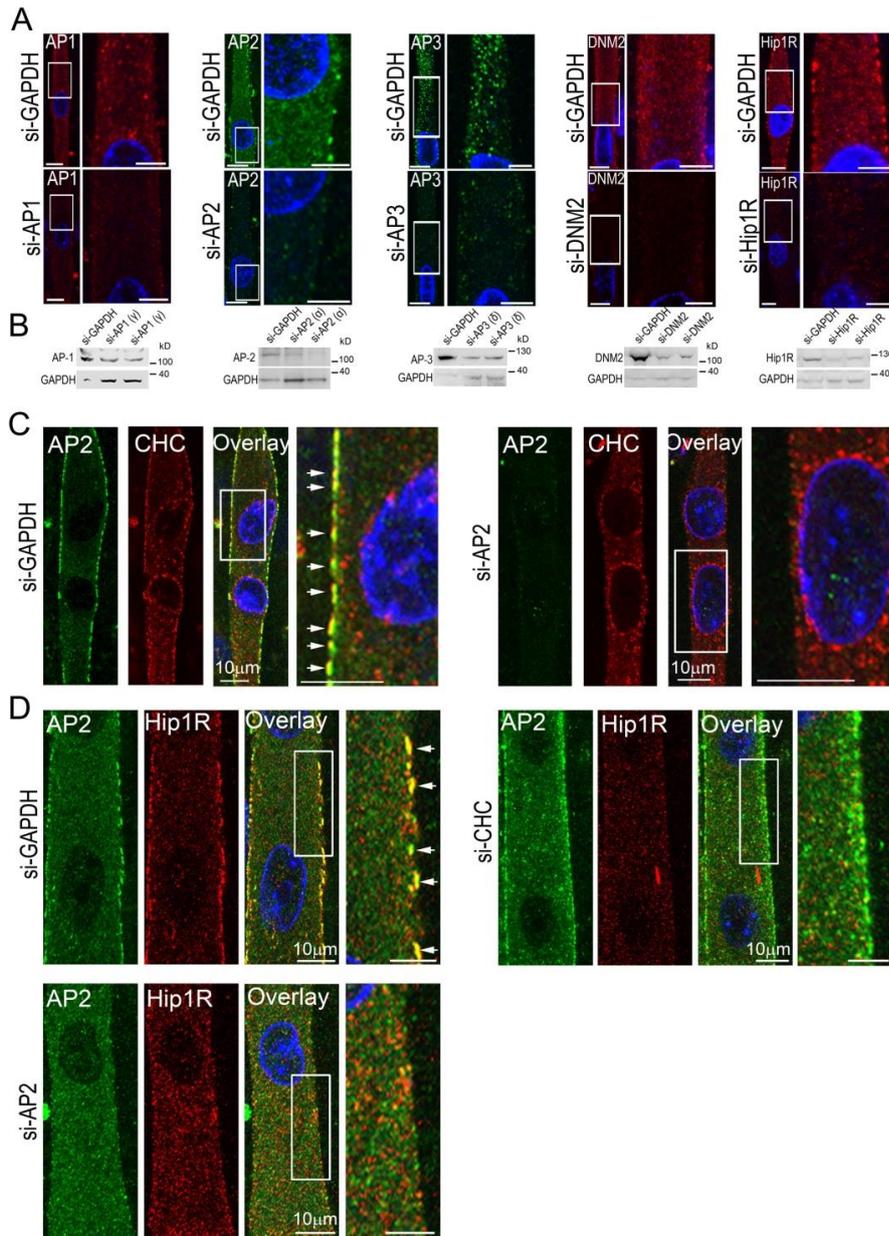


Figure S3: siRNA-mediated depletion of AP1, AP2, AP3, DNM2 and Hip1R. (A) Differentiated C2C12 myotubes were treated with either siRNA targeting GAPDH, the γ subunit of AP1, the α subunit of AP2, the δ subunit of AP3, DNM2 or Hip1R. Immunofluorescent staining (DNA staining with DAPI is shown in blue (scale=10 μ m) and the boxed region in the merged images is magnified in the insets (scale=5 μ m)). (B) Differentiated C2C12 myotubes were treated with either siRNA targeting GAPDH, the γ subunit of AP1, the α subunit of AP2, the δ subunit of AP3, DNM2 or Hip1R and subjected to Western blot detection of the γ subunit of AP1, the α subunit of AP2, the δ subunit of AP3, DNM2, or Hip1R. (C) Differentiated C2C12 skeletal muscle cells were treated with control GAPDH siRNA or siRNA against AP2 and processed for immunofluorescent staining with antibodies against the α subunit of AP2 (green) and CHC (red). DNA staining (DAPI, blue) identifies differentiated, multinucleated myotubes (scale=10 μ m). The boxed region in the merged images is magnified in the insets (scale=10 μ m). (D) Differentiated C2C12 myotubes were treated with either siRNA targeting GAPDH, CHC or the α subunit of AP2 and processed for immunofluorescent staining with antibodies against the α subunit of AP2 (green) and Hip1R (red). DNA staining (DAPI, blue) identifies differentiated, multinucleated myotubes (scale=10 μ m). The boxed region in the merged images is magnified in the insets and arrows in C and D indicate co-localization between AP2 and CHC or AP2 and Hip1R respectively (scale=5 μ m).

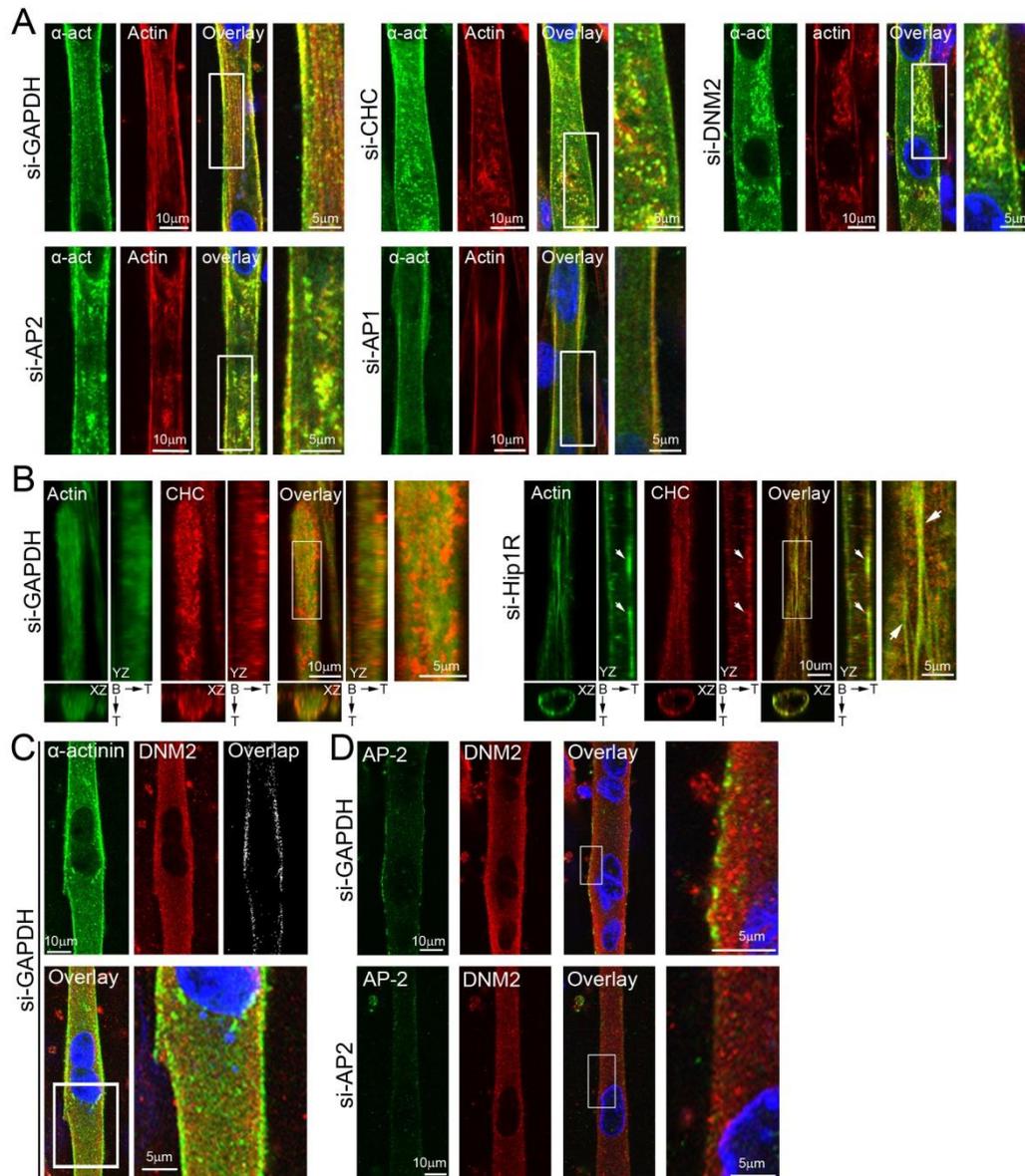


Fig. S4: Effect of siRNA-mediated depletion of CHC, AP1, AP2 and DNM2 on actin distribution and localization of DNM2 in control and AP2-depleted C2C12 myotubes. (A) Differentiated C2C12 myotubes were treated with either siRNA targeting GAPDH, CHC, the γ subunit of AP1, the α subunit of AP2 or DNM2 and processed for immunofluorescent staining with antibodies against α -actinin (green) and treated with TRITC-phalloidin in order to label actin filaments (red). DNA staining (DAPI, blue) identifies differentiated, multinucleated myotubes. The boxed region in the merged image is magnified in the inset. Arrows indicate regions with intracellular α -actinin accumulations. (B) Differentiated C2C12 myotubes were treated with either siRNA targeting GAPDH or Hip1R and processed for immunofluorescent staining with FITC-phalloidin (green) in order to label actin filaments and antibodies against CHC (red) (scale=10 μ m). XZ and YZ projections of serial confocal sections are shown for each individual channel and the overlay image. The boxed region in the overlay is magnified in the insets (scale=5 μ m). Arrows indicate co-localization between actin filaments and CHC. (C) Immunofluorescent staining of α -actinin (green) and DNM2 (red) in differentiated C2C12 myotubes. DNA staining (DAPI, blue) identifies differentiated, multinucleated myotubes. The boxed region in the merged images is magnified in the insets. A black and white composite image depicting co-localized pixels between the α -actinin and DNM2 staining is shown. (D) Differentiated C2C12 myotubes were treated with either siRNA targeting GAPDH or the α subunit of AP2 and processed for immunofluorescent staining with antibodies against the α subunit of AP2 (green) and DNM2 (red). DNA staining (DAPI, blue) identifies differentiated, multinucleated myotubes. The boxed region in the merged images is magnified in the insets.

