Analysis of skeletal muscle function in the C57BL6/SV129 syncoilin knockout mouse

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Abstract Syncoilin is a 64-kDa intermediate filament protein expressed in skeletal muscle and enriched at the perinucleus, sarcolemma, and myotendinous and neuromuscular junctions. Due to its pattern of cellular localization and binding partners, syncoilin is an ideal candidate to be both an important structural component of myocytes and a potential mediator of inherited myopathies. Here we present a report of a knockout mouse model for syncoilin and the results of an investigation into the effect of a syncoilin null state on striated muscle function in 6–8-week-old mice. An analysis of proteins known to associate with syncoilin showed that ablation of syncoilin had no effect on absolute expression or spatial localization of desmin or alpha dystrobrevin. Our syncoilin-null animal exhibited no differences in cardiotoxin-induced muscle regeneration, voluntary wheel running, or enforced treadmill exercise capacity, relative to wild-type controls. Finally, a mechanical investigation of isolated soleus and extensor digitorum longus indicated a potential differential reduction in muscle strength and resilience. We are the first to present data identifying an increased susceptibility to muscle damage in response to an extended forced exercise regime in syncoilin-deficient muscle. This study establishes a second viable syncoilin knockout model and highlights the importance of further investigations to determine the role of syncoilin in skeletal muscle.

Introduction

Syncoilin is a recently described member of the intermediate filament (IF) family of proteins (Newey et al. 2001). So named for its enrichment at the neuromuscular synapse and its coiled-coil-forming rod domains, the syncoilin gene has been mapped to chromosome 1 in humans and to chromosome 4 in mice. The syncoilin gene comprises five coding exons that generate a 64-kDa protein product (Newey et al. 2001). Following its discovery, subsequent yeast two-hybrid screening analysis showed that the IF protein desmin and the dystrophin-related protein \( \alpha \)-dystrobrevin were syncoilin binding partners in striated muscle (Newey et al. 2001; Poon et al. 2002). Histologic studies utilizing immunofluorescent and confocal microscopy have confirmed that syncoilin colocalizes under normal conditions with both desmin and \( \alpha \)-dystrobrevin (McCullagh et al. 2007; Newey et al. 2001; Poon et al. 2002).

One distinctive feature of syncoilin is its apparent departure from spontaneous dimer-forming behavior. Syncoilin does not appear to form a filamentous network by itself or with other IF proteins, although it may form filaments in certain myopathies (Brown et al. 2005; Poon et al. 2002).

Syncoilin is highly expressed in skeletal and cardiac muscle, and in the former it is concentrated at the sarcolemma, Z-lines, perinuclear space, myotendinous junction, and neuromuscular junction (McCullagh et al. 2007;
Within the skeletal muscle fiber population, syncoilin is enriched at the sarcolemma and sarcoplasm of oxidative muscle fibers (McCullagh et al. 2007). Thus, the localization of syncoilin parallels that of desmin, to which it binds directly.

Desmin is the dominant IF protein in mature muscle and has been predicted to play a major role in protecting muscle from mechanical stress and is important for muscle integrity (Capetanaki et al. 2007; Costa et al. 2004). An analysis of the tensile properties of single desmin filaments has shown that it exhibits a nonlinear response to strain (notably above 50% extension) and a tensile strength of at least 240 mPa (Kreplak et al. 2008). These properties add further weight to the theory that desmin acts as both a mediator of myocyte spatial organization and provides a physical link between adjacent contractile units.

Desmin knockout (des−/−) mice show a skeletal myopathy as well as a cardiomyopathy, and desmin aberrations have been linked to disease in humans (Li et al. 1996; Milner et al. 1996; Olive et al. 2004). A study of exercise ability showed a reduction in the capacity of des−/− mice to perform both voluntary wheel running and enforced treadmill exercise. Most interestingly, the study showed that an absence of desmin was not sufficient to increase serum creatine kinase levels (a marker of sarcolemmal damage) (Haubold et al. 2003).

We have previously shown that the levels of syncoilin are reduced by 60% in these animals and that the subcellular location of syncoilin is altered. In the des−/− mice, syncoilin is lost from the Z-lines and reduced at the sarcolemma (McCullagh et al. 2007). Alterations in syncoilin expression have been reported for other myopathies, including elevated levels in dystrophic muscles of mdx mice and in atrophic muscle fibers of patients with neurogenic disorders (Brown et al. 2005; Newey et al. 2001).

Although there is clear evidence showing that syncoilin binds directly to both α-dystrobrevin and desmin, whether syncoilin has the ability to bind to these proteins simultaneously and thus form a link between the cytoskeletal IF network and the sarcolemma via the dystrophin-associated protein complex (DAPC) remains to be investigated. This potential linking of the DAPC to the IF network via syncoilin is of interest since the lateral transmission of the cytoskeleton and contractile apparatus of myocytes (via an interaction of dystrophin with F-actin and indirectly with the IF network via α-dystrobrevin) (Blake et al. 2002).

In view of the findings that the loss of either desmin or dystrobrevin results in the development of muscular dystrophy in association with changes in the distribution of syncoilin, we decided to test the hypothesis that loss of syncoilin might have an adverse effect on myocyte function. We generated a syncoilin knockout mouse by standard techniques of homologous recombination in embryonic stem (ES) cells and a Cre-loxP-mediated conditional gene-targeting approach. Such null mutants would allow the investigation of the role of syncoilin in skeletal and cardiac muscle. This strategy allowed us to produce two lines of mice: conditional (sync-floxed) and constitutive syncoilin (sync−/−) knockout mice. Constitutive deletion of syncoilin resulted in a viable mouse, precluding the need for tissue-specific excision of the syncoilin gene. In this article we present the results of a molecular and phenotypic characterization of striated muscle from a novel sync−/− mouse.

Materials and methods

Animals and antibodies

All mice were bred in-house and genotyped by standard PCR methods. All animal experiments were performed according to guidelines and protocols approved by the Home Office. Antibodies used were anti-syncoilin-FP 81, anti-desmin (Abcam8592), anti-desmin monoclonal antibody (DE-U-10; Sigma), anti-dystrobrevin-1 and -2 antibody β1CTFP (Blake et al. 1998), anti-dysbindin antibody (Benson et al. 2001; Newey et al. 2001), anti-α-actinin mouse (N40; Santa Cruz), anti-α-actinin antibody (EA-53; Sigma) and anti-dystrophin antibody (Dys-2; Novacastro). A synemin antibody (anti-Syn-3) that detects both α- and β-synemin was generated as previously described (Mizuno et al. 2004).
Gene targeting and generation of syncoilin knockout mice

The targeting construct is shown in Fig. 1. Genomic regions of the syncoilin gene were amplified by PCR from a 129/Sv bacterial artificial plasmid (Wellcome Trust Sanger Institute, Cambridge, UK) which contained the syncoilin locus. The targeting construct was assembled in the pBluescript II KS+ vector. The 5' arm of homology consisted of a 1.6-kb fragment upstream of syncoilin exon 2 followed by a loxp site and neomycin flanked by frt recombinase sites. The 3' arm of homology was a 5.9-kb fragment containing exon 2 followed by a loxp site. The targeting vector was verified by sequencing and linearized with NcoI before electroporation into E129/sv ES cells. Four hundred G418-resistant ES clones were selected for screening for homologous recombination by PCR and Southern blotting. Genomic DNA isolated from each clone was digested with NcoI and analyzed by Southern blotting. A 5' internal probe and 3' external probe (indicated in Fig. 1) were generated from mouse E129 genomic DNA.

Southern blots were hybridized with radiolabeled probes and visualized by autoradiography. The 5' internal probe was used to confirm recombination of the 5' arm of the targeting vector, where the wild-type allele is indicated by a band at 2.3 kb, and a 3.5-kb band represents the correctly targeted allele. Hybridization of blots with the 3' external probe was used to verify recombination of the 3' arm of the targeting vector, where the wild-type allele is represented by a band at 8.2 kb, and a 6.3-kb band represents the correctly targeted allele.

Generation and genotyping of mice

Two independent correctly targeted ES clones were microinjected into blastocysts from C57BL/6 mice. Chimeric mice were bred with transgenic mice ubiquitously expressing Cre recombinase under the cytomegalovirus promoter (CMV), to generate germline-transmitted syncoilin heterozygous (sync+/−) mice. Sync+/− mice were intercrossed to generate homozygous syncoilin null (sync−/−) mice. Offspring from intercrosses were genotyped by PCR analysis of sync−/+ and sync−/− muscles. Standard exonic GAPDH primers were run on duplicate samples as a control for the cDNA template.

Protein preparation and immunoblotting

Skeletal or cardiac muscles were snap-frozen immediately after dissection in liquid nitrogen and stored at −80°C until use. The tissue was homogenized in 300 μl cold Newcastle buffer (75 mM Tris pH 6.8, 3.8% SDS, 4 M urea, and 20% glycerol) with protease inhibitor cocktail (Sigma). Protein quantification was performed using the Bio-Rad Dc Protein Assay according to the manufacturer’s protocol. Total protein (40 μg) was separated on 8 or 10% SDS polyacrylamide gels (Bio-Rad) and electrophoretically transferred to nitrocellulose membranes. After blocking membranes overnight at 4°C in Tris-buffered saline containing 0.1% (v/v) Tween20 (TBS-T) and 5% nonfat dry milk, blots were incubated for 1 h with primary antibody and washed three times in TBS-T prior to incubation with horseradish peroxidase conjugated IgG in blocking buffer. Membranes were washed as previously described and developed using ECL detection reagents (Amersham) as per the manufacturer’s instructions. Blots were stripped at 50°C for reprobing with different antibodies. Exposed film was scanned with a HP Scanjet 5400c.

Single myofiber preparations

Mouse flexor digitorum brevis muscles (FDBs) were enzymatically dissociated using an established protocol (Brown and Schneider 2002; Woods et al. 2004). The FDB was dissected free and placed into 1 mg/ml of collagenase (Type 1, Sigma) in Ringer buffer (119 mM NaCl, 5 mM KCl, 1.25 mM CaCl2, 1 mM MgSO4, 1 mM KH2PO4, 10 mM glucose) at 37°C for 1.5 h. Collagenase activity was stopped by washing the muscle with warmed Ringer’s solution and the muscle was gently triturated using a Pasteur pipette until single fibers were isolated. One hundred fifty microliters of Ringer’s solution containing FDB single fibers was spread across laminin-coated glass bottom culture dishes (Matek Corp., Part No P35G-1.0) and incubated in 5% CO2 at 37°C. After 2 h, Ringer’s solution was removed and the dish was
flooded with 4% PFA/PBS for 5 min to fix the fibers. This method yielded approximately 100-200 fibers per dish. The fibers were then permeabilized in 0.1% Triton/PBS and then processed for immunolabeling.

Primary antibodies were diluted in PBS containing 10% FCS and applied to the fiber preparations. After an overnight incubation at 4°C, fiber preparations were rinsed in PBS followed by incubation with secondary antibodies (Molecular Probes); Alexa fluor 594 and/or Alexa fluor 488 conjugated IgG diluted (1:2000) in PBS, 10% FCS for 2 h at room temperature. Finally, fiber preparations were washed thoroughly in PBS and mounted in an aqueous medium (VectorShield) with DAPI. Finished preparations were viewed under a Zeiss fluorescent photomicroscope or Zeiss 410 confocal laser-scanning microscope. Digital images were extracted using Axiovision LE 4.3 software.

Measurement of contractile properties

Mechanical measurements of skeletal muscle performance in vitro were performed using a previously described protocol (with minor adaptations) (Moens et al. 1993). Briefly, extensor digitorum longus (EDL) and soleus muscles were dissected from 8-week-old male mice killed by cervical dislocation. The muscles were pinned out in oxygenated Ringer’s solution at room temperature to a length approximating that seen in situ. Oxygenated (95% O2/5% CO2) mouse Ringer’s solution (pH 7.4) was made up to include 118 mM NaCl, 24.8 mM NaHCO3, 4.75 mM KCl, 2.54 mM CaCl2, 11.8 mM MgSO4, 11.8 mM KH2PO4, and 10 mM glucose. Muscles were experimented on within 4 h of excision. The freshly dissected muscles were mounted onto the muscle lever of the Intact Muscle Measurement System (Aurora Scientific Inc.) using 5-0 silk suture, and maintained throughout the protocol at room temperature in oxygenated Ringer’s solution. The muscles were stimulated using platinum electrodes connected to a Aurora 701B stimulator. The optimum length was identified, and then the maximum isometric tetanic force was measured (EDL, 120 Hz, 500 ms; soleus, 80 Hz, 800 ms). Force measurements were normalized to muscle cross-sectional area. The wet weight of the muscles was recorded at the end of the experiment. Data were captured and analyzed using the DMC/DMA Dynamic Muscle Control and Analysis software (Aurora Scientific Inc.).

Cardiotoxin-induced damage of hind limb muscle

Sex-matched adult mice were induced and maintained under isoflurane inhalation anesthetic as discussed above. A small incision was made to expose the tibialis anterior. Using a 100-μl insulin syringe, 50 μl of a 10 μM solution of Cardiotoxin 1 (Sigma Aldrich, Dorset, UK) dissolved in sterile PBS was introduced into the muscle belly. The overlying skin was sutured to close the wound. The contralateral limb was treated with an identical volume of PBS to serve as a control. Three mice of each genotype were treated for each timepoint. Following treatment, mice were aged for 1, 3, 5, 7, or 21 days. Treated and control tibialis anterior muscle for histologic analysis was dissected, embedded in OCT, and snap-frozen in liquid nitrogen-cooled isopentane. Treated and control tibialis anterior muscle for RNA analysis was dissected and snap-frozen in liquid nitrogen.

Morphologic analysis of myoregeneration following cardiotoxin treatment

OCT-embedded cardiotoxin-treated muscle and PBS-treated tibialis anterior muscle were sectioned on a cryostat to a thickness of 9-12 μm. Sections were air-dried at room temperature, rapidly fixed in absolute ethanol, then stained with hemotoxilin and eosin and mounted in an aqueous mounting media using standard histologic techniques.

Analysis of myoregeneration marker induction

Total RNA extraction and generation of cDNA from excised cardiotoxin- and PBS-treated tibialis anterior...
tissue were performed as described above. Semiquantitative PCR analysis of the cell cycle-dependent kinase inhibitors p21CIP1, p27KIP1, p57KIP2 and myogenin was performed as described previously (Koo et al. 2007).

Measurement of voluntary wheel running activity

To investigate the effect of syncoilin ablation on activity levels, we monitored voluntary wheel running in two groups (n = 6) of 6-week-old male sync+/+ and sync−/−
mice. Mice were placed in individual cages supplemented with a monitored running wheel. Following an acclimatization period of 1 week, the total number of wheel revolutions performed by each animal was electronically recorded for an additional 7-day period using the ClockLab software suite (Actimetrics Software).

Measurement of enforced treadmill exercise capacity and resultant muscle damage

Treadmill exercise is a well-established means of analyzing muscle function in rodents (Armand et al. 2003). Two groups of 6-week-old male sync+/- and sync-/- mice were exercised on a motorized treadmill (Exer6-M, Columbus Instruments). The regimen consisted of 5 days of familiarization running (10 m/min for 15 min), 9 days of endurance running (17 m/min for 45 min), and 1 day of exhaustion running (17 m/min for 45 min followed by an increase in speed of 1 m/min for 15 min). Each of the endurance/exhaustion running stages began with a 5-min warm-up period at a speed of 10 m/min. Animals were rested for 2 days after each 5-day block of running. The weight of each animal was recorded before and after each running session.

Animals were killed immediately following the final running session and approximately 100 µl of venous blood was collected from the exposed jugular vein. Following a 3-h incubation on wet ice, serum was isolated by centrifugation in a benchtop centrifuge for 5 min at 14,000 rpm. Serum creatine kinase (CK) analysis was performed using a kinetic UV serum analysis according to manufacturer’s instructions (OSR6179, Olympus R&D, Ireland). Muscle was dissected for hemotoxylin and eosin staining as described previously. Evans Blue staining of isolated muscle was performed in 1 ml of mouse Ringer’s solution supplemented with 0.1% (w/v) Evans Blue dye. Muscle was incubated in solution for 2.5 h at 37°C before being washed in mouse Ringer’s solution and embedded in OCT as described previously.

Results

Generation of gene-targeted syncoilin-null mice

We generated homozygous syncoilin knockout (sync-/-) mice by deleting the gene by in vivo Cre-loxP-mediated excision of exon 2, which encodes 85% of the amino acid sequence (Fig. 1a). This strategy was predicted to ensure the ablation of functional syncoilin expression in knockout animals. The targeting vector was electroporated into E129/sv embryonic stem (ES) cells, and 167 clones were screened. Twenty-four clones were identified that had undergone homologous recombination (example shown in Fig. 1a). Two independent clones were each injected into blastocysts from C57BL/6 mice and then reimplanted to generate chimeric mice, bearing the floxed targeted syncoilin allele. Chimeric mice were bred with female CMV-Cre-expressing transgenic mice to excise exon 2 of syncoilin in vivo and give rise to germline-transmitted heterozygous (sync-/-) mice. In vivo mediated excision of exon 2 was verified by PCR genotyping. Heterozygous mice were interbred to generate homozygous syncoilin knockout (sync-/-) mice, which were confirmed by PCR genotyping (Fig. 1c) and RT-PCR analysis for exons 2–5 of RNA from tibialis anterior, soleus, and heart muscles (Fig. 1d). Different genotypes (sync+/-, sync +/-, and sync -/-) were obtained in the expected Mendelian ratio. Syncoilin-null mice were viable and were indistinguishable from their wild-type littermates in appearance, behavior, and body weight.

Syncoilin protein analysis in sync-/- mice

Western blotting analysis of muscle protein from sync-/- mice with a polyclonal-antibody recognizing full-length (FL) syncoilin protein confirmed that we had ablated syncoilin expression (Fig. 2). Immunoblotting of the above muscle samples with an antibody recognizing an amino terminal (N) epitope (amino acids 1–146; encoded by exons 1 and 2), which includes the translation initiation site, gave the same results as the full-length syncoilin antibody, indicating that no aberrant syncoilin protein could be detected in sync-/- tissues.
Analysis of syncoilin-binding proteins and other muscle proteins in sync−/− mice

DAPC-associated and syncoilin-binding partner proteins were quantified by Western blotting from total skeletal and cardiac muscle protein prepared from sync+/+, sync+/−, and sync−/− mice. Binding partners of syncoilin, desmin and α-dystrobrevin were unaffected in sync+/+ and sync−/− mouse skeletal or cardiac muscle (Fig. 2). An antibody recognizing synemin isoforms H and M revealed no variation in muscles from the different syncoilin mouse genotypes (Fig. 2). Equal protein loading was demonstrated by similar α-actinin levels between samples.

To assess if the absence of syncoilin affects the subcellular localization of syncoilin-binding proteins, the sarcomeric Z-lines, or other cytoskeletal proteins, we performed immunofluorescence on isolated myofibers. Fibers immunostained for desmin revealed no change in their normal location, with fibers from both sync+/+ and sync−/− mice showing normal localization to Z-lines and myotendinous junctions (ends of myofibers) (Fig. 3). α-Dystrobrevin immunostaining (data not shown) also exhibited a normal distribution in both sync+/+ and sync−/− fiber preparations. Thus, the subcellular localization of known syncoilin-binding partners is not dependent on the presence of syncoilin in muscle. Immunostaining of fibers for the Z-line protein α-actinin gave the expected Z-disc striated pattern staining in both sync+/+ and sync−/− myofibers (Fig. 3).

Morphology of skeletal and cardiac muscle in sync−/− mice

We examined the morphology of the more highly active muscles in 8-week-old mice by light microscopy of hematoxylin and eosin-stained sections of soleus, diaphragm, and heart from sync+/+ and sync−/− mice (Fig. 4). Histologic analysis revealed no gross differences between genotypes, and there was no sign of any dystrophic features or abnormal morphology in skeletal and cardiac muscle from sync−/− mice.

Absence of syncoilin reduces contractile function in skeletal muscles

To investigate the effect of syncoilin ablation on maximum isometric force generation, we measured force production in isolated soleus and extensor digitorum (EDL) muscles from 8-week-old sync+/+ and sync−/− mice. Four EDL muscles were assayed for each genotype. For the soleus analysis, eight sync+/+ and six sync−/− muscles were assayed. Isometric contractile testing of EDL muscles from sync−/− mice revealed a strongly significant decrease in normalized maximum force generation of −27.9% compared with sync+/+ mice (25.86 N/cm² vs. 35.89 N/cm²; p < 0.05) (Fig. 6a), whereas there was no significant difference observed between the soleus muscles of sync+/+ and sync−/− mice (Fig. 5).

Ablation of syncoilin does not elicit a morphologic change in the rate of myoregeneration

To investigate any changes in muscle regeneration due to the absence of syncoilin expression, we assessed regeneration at five time points, 1, 3, 5, 7, and 21 days, following the intramuscular administration of cardiotoxin to the tibialis anterior muscle in sync+/+ and sync−/− mice. Figure 6a shows representative images of muscle regeneration from the sync+/+ and sync−/− groups (n = 3 for each genotype) across the five time points. No obvious differences in degeneration, infiltration, or regeneration (including the number of centralized nuclei at 21 days) were observed between sync+/+ and sync−/− animals.

Ablation of syncoilin does not elicit an alteration in markers of myogenic differentiation

To investigate any potential difference in the kinetics of skeletal myocyte regeneration between sync+/+ and sync−/− animals, we performed a semiquantitative assessment of the relative RNA levels of the cell cycle-dependent kinase inhibitors p21CIP1, p27KIP1, p57KIP2 and myogenin in cardiotoxin-treated tibialis anterior muscle

Fig. 3 Immunostaining of single-fiber preparations from flexor digitorum brevis in sync+/+ and sync−/− mice. Localization of syncoilin to the Z-lines, sarcolemma, and MTJ is evident in the sync+/+ animal and lost in the sync−/− animal. Desmin and α-actinin organization shows no evidence of disruption as a result of syncoilin ablation. (a) Syncoilin staining across whole fiber. (b) Desmin staining across whole fiber. (c) Syncoilin staining at Z-lines. (d) Desmin staining at Z-lines. (e) α-Actinin staining at Z-lines.
at 1, 3, 5, and 7 days after administration. An increase in cell cycle-dependent kinase inhibitors can be correlated with the initiation of myogenic differentiation at the expense of cell cycling. As such, these transcript levels can be used as markers of myoregeneration kinetics (Koo et al. 2007). No obvious difference in transcript levels was detected between the sync+/+ and sync−/− animals, indicating that the regeneration kinetics of skeletal muscle regeneration are not altered by the ablation of syncoilin (Fig. 6b).

Ablation of syncoilin does not produce a statistically significant difference in voluntary wheel running

A strong interanimal variation in running levels was identified in both groups (sync+/+ and sync−/−) of age- and sex-matched mice. Total individual (A) and group average (B) revolution counts are shown in Fig. 7. Individual cage positions are presented in paired form to compensate for any potential differential effect in

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environment. Two-tailed analysis using Student’s t test did not identify a statistically significant difference between the averaged activity scores (p = 0.57) of the sync+/+ and sync−/− groups.

Ablation of syncoilin induces increased muscle damage without producing an observable difference in enforced treadmill running ability

An analysis of the enforced exercise capacity of sync+/+ and sync−/− mice failed to demonstrate a reduction in the ability of the sync−/− group to complete the prescribed exercise protocols. Eighty-three percent of sync+/+ and 100% of sync−/− mice successfully completed the final 60-min exhaustion test, reaching a maximum speed of 32 m/min after 60 min of running. All animals continued to gain weight normally over the course of the study (data not shown). Figure 8 shows the results of a histologic examination of muscle morphology following 15 running sessions. No obvious differences in the gross morphology of sync+/+ and sync−/− transverse sections were detected. A very small number of centralized nuclei were detected in both sync+/+ and sync−/− tibialis anterior sections. Evans Blue staining of isolated muscle did not indicate increased dye penetration (and therefore sarcolemmal damage) in sync−/− muscle.

CK analysis of serum taken from sync+/+ and sync−/− animals (n = 6 per genotype) immediately following the completion of their running protocol showed a significant (p < 0.05, two-tailed Student’s t test) increase in serum CK levels in the sync−/− population (Table 1).

Discussion

Syncoilin is a 64-kDa intermediate filament-like protein expressed in skeletal and cardiac muscle that is found to be enriched at the perinucleus, the Z-lines, and at the sarcolemma and neuromuscular junction (Newey et al. 2001). Previous studies have shown that syncoilin interacts with both α-dystrobrevin (a DAPC component) and desmin (the major intermediate filament expressed in mature myocytes) (Newey et al. 2001; Poon et al. 2002). By ablating in excess of 80% of the syncoilin-coding sequence, we have generated a mouse model that is null (−/−) for syncoilin expression.

Developmentally, sync+/+ and sync−/− mice are indistinguishable. Protein expression analysis of the syncoilin-binding partners desmin and α-dystrobrevin failed to demonstrate a difference in their net concentrations in muscle derived from sync+/+ and sync−/− mice. Immunofluorescent staining of single-fiber preparations of flexor digitalis brevis muscle demonstrated no difference in desmin or α-dystrobrevin localization between sync+/+ and sync−/− mice. Together, these data indicate that syncoilin is not essential for either the correct cellular localization or expression of these proteins (Figs. 2 and 3).

The treatment of skeletal muscle with myonecrotic cardiotoxin induces widespread necrosis followed by a well characterized cascade of satellite cell activation, proliferation, differentiation, and maturation (Shi and Garry 2006). Our histologic analysis of cardiotoxin-treated tibialis anterior muscle yielded no detectable difference in regenerating muscle at any of the time points between sync+/+ and sync−/− mice. Semiquantitative PCR analysis of myogenin, p21, p27, and p57 expression in cardiotoxin-treated tissue also failed to elicit a noticeable difference in transcript levels. When their expression is upregulated, these proteins act to initiate myogenic differentiation and prevent the cell from re-entering the cell replication cycle (Koo et al. 2007). These findings indicate that a lack of syncoilin does not obviously retard muscle tissue regeneration and that thus syncoilin is unlikely to play an essential role in this process. The results of both our voluntary running and enforced treadmill exercise studies indicate that phenotypically syncoilin is dispensable for the normal mechanical function of striated muscle under exercise conditions.
Interestingly, an investigation of contractile function in isolated soleus and EDL demonstrated a reduction of nearly 30% in the maximum force generated by sync−/− EDL compared with sync+/+ EDL (although no difference in the soleus analysis) (Fig. 5). EDL comprises mostly fast (type II) myofibers, in contrast to soleus muscle which is populated mainly by slow (type I) myofibers. The reduction in maximum force generation seen in EDL may indicate

**Fig. 6** (a) Regeneration of tibialis anterior muscle following treatment with cardiotoxin. Representative images (n = 3 for each genotype at each time point) of hemotoxylin and eosin staining of 9–12-μm cryosections taken from animals at 1, 3, 5, 7, and 21 days after toxin administration. No obvious difference in degeneration, basophilic infiltration, or regeneration was noticed between sync+/+ and sync−/− mice. (b) Myogenic differentiation analysis. Semiquantitative PCR analysis of p21CIP1, p27KIP1, p57KIP2, and myogenin RNA transcript levels in tibialis anterior at 1, 3, 5, and 7 days following cardiotoxin administration. Images (which are representative of n = 3 for each genotype at each time point) indicate no difference in regeneration kinetics between sync+/+ and sync−/− animals.
that syncoilin plays varying roles or has a variable functional redundancy in different fiber types. The identification of a force reduction in sync-/− EDL is in accordance with a similar analysis performed by Zhang et al. (2008). Using an alternative syncoilin-null model (generated by the introduction of a point mutation resulting in the aberrant mRNA splicing of the second syncoilin exon), this study reinforces the findings presented in our own analysis, namely, that syncoilin appears to be largely redundant for striated muscle development and regeneration (Zhang et al. 2008).

Our analysis of serum CK levels following 3 weeks of enforced treadmill exercise is the first report of a functionally induced myopathy deriving from an absence of syncoilin (Table 1). Elevated serum CK levels are widely used as a marker of sarcolemmal damage. This finding is in contrast to earlier studies performed on des−/− mice, which failed to exhibit an increase in serum CK levels after a bout of enforced treadmill exercise (Haulbold et al. 2003).

Fig. 7 Seven-day analysis of voluntary wheel running in sync+/+ and sync−/− mice. No significant difference was detected in the total activity levels of the two groups

An analysis of serum CK levels in des−/− mice was performed immediately following and 24 h after a brief period of low-speed downhill running as described by Vilquin et al. (1998) for use in the dystrophin mdx mouse model (Haulbold et al. 2003; Vilquin et al. 1998). The results of this study did not show a difference in CK levels between wild-type and des−/− animals either immediately after or 1 day after exercise.

The mdx mouse is used as a model of dystrophin ablation-induced muscular dystrophy and exhibits a severe myopathic phenotype (Vilquin et al. 1998). The phenotype of a des−/− mouse is far milder than that exhibited by the mdx mouse, although des−/− mice still exhibit a reduction in running ability when compared with wild-type animals (Haulbold et al. 2003). At a molecular level, a loss of desmin does not appear to induce disruption of the essential DAPC as is seen with a loss of dystrophin. Although it is surprising that a loss of syncoilin induces an increase in serum CK levels whereas a loss of desmin does not (given the proposed role of syncoilin and the abundance of desmin in mature myocytes), it is a possibility that the exercise regimen applied to the des−/− mice was not sufficient in either duration or intensity to induce sufficient damage to cause mechanical stress.

The precise function of syncoilin remains undelineated. However, we do possess several pieces of evidence that can assist us in generating hypotheses about the role(s) of syncoilin in muscle. Given that syncoilin is a member of the IF protein family (at least at a sequence homology level), we can reasonably expect that it plays a role in line with that of other IF proteins, which, among others, involves protecting cells from mechanical stress (Parry et al. 2007). Syncoilin’s association with the DAPC and other IF proteins (in particular, the finding that it binds to desmin) adds weight to it having a structural role in muscle. Our studies indicate that at least in an acute setting there are sufficient mechanisms in existence in myocytes to cope with a loss of syncoilin. One such potential compensatory mechanism may exist in skeletal muscle in the form of the type IV IF protein synemin, although it is worth noting that there are significant size and structural differences between these two IF proteins.

Synemin is a large IF protein expressed in all types of human muscle cells as two splice variants, α-synemin and β-synemin (Sun et al. 2008). Synemin has been shown to act as a binding partner for a range of cytoskeletal and structural proteins and contains a binding site for α-actinin (Bellin et al. 1999, 2001). Recent studies have indicated that the two variants may play different roles in muscle, with the ability to interact with vinculin and metavinculin limited to α-synemin, which contains an additional C-terminal 312-residue insert not found in β-synemin (Sun et al. 2008). Importantly for our investigation of syncoilin,
synemin has also been shown to act as a binding partner for desmin and α-dystrobrevin, and the spatial organization of synemin exhibits a number of parallels to that of syncoilin (Bhosle et al. 2006). As such, it has the potential to act in a compensatory role in a

Our Western blot analysis for both H and M variants (analogous to α and β variants in humans) of synemin showed no difference in net protein levels between sync+/+ and sync−/− mice, indicating that a lack of overt phenotype in the sync−/− mouse is not due to compensation by synemin. It is possible, however, that the presence of synemin in what appears to be a similar role in muscle offsets or masks the deleterious effects of a loss of syncoilin, such that under normal laboratory cage or mild exercise conditions, syncoilin exhibits a degree of functional redundancy.

In summary, our investigations show that we have generated a functional example of a syncoilin knockout mouse model. Syncoilin-null mice do not display an overt phenotype. An absence of syncoilin does not (in the proteins investigated in this study) induce a compensatory protein upregulation in skeletal muscle or the disruption of protein localization in isolated fibers. No difference in muscle regeneration following cardiotoxin-induced necrosis was observed, and an absence of syncoilin did not affect voluntary or enforced exercise capacity.

The mechanical analysis of isolated muscle revealed a significant alteration in contractile properties, indicating that syncoilin may play a role in the lateral transmission of force across contractile tissues. Most interestingly, we have shown that under extended exercise conditions, an absence of syncoilin predisposes muscle to increased damage.

Given the findings in this and other studies, it is possible that the role of syncoilin is analogous to that of “a single spoke in a wheel,” in that its structural role is not appreciated (nor its absence noted) until significant levels of mechanical stress are applied to the syncoilin-deficient system.

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