Dystrophin is a microtubule-associated protein

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Introduction

The plakins are a class of giant cytolinker proteins that can link transmembrane protein complexes to the actin, intermediate filament, and microtubule (MT) cytoskeletons in various combinations (Fuchs and Karakesisoglou, 2001). Plakins can bind actin filaments via tandem calponin homology (CH) domains, intermediate filaments via plakin repeat domains, and MTs through either a Gas2-related domain or a glycine-serine/arginine domain (for review see Leung et al., 2002). The ability to cross-link multiple components of the cellular cytoskeleton allows cytolinkers to stabilize cells from mechanically induced damage. Mouse knockout studies exemplify the stabilizing effects of cytolinkers, as loss of the cytolinker plectin resulted in skin blistering and a form of muscular dystrophy (Andra et al., 1997), and ablation of BPAG2 in mice caused skin blistering upon mechanical stimulation (Guo et al., 1995).

Dystrophin, the protein absent in patients with Duchenne muscular dystrophy (Hoffman et al., 1987), shows structural and functional similarities to cytolinkers, which suggests the hypothesis that dystrophin performs a cytolinker role in muscle. Dystrophin’s large molecular mass of 427 kD, spectrinlike repeats, and ability to bind actin filaments via tandem CH domain (Way et al., 1992) highlight three similarities with cytolinkers. Although dystrophin lacks a plakin repeat domain, dystrophin–intermediate filament interactions have been documented (Stone et al., 2005; Bhosle et al., 2006). Thus, the ability to link the actin and intermediate filament cytoskeletons to the transmembrane dystroglycan complex (Suzuki et al., 1992) illustrates how dystrophin functions similarly to other cytolinkers. Finally, the muscle membrane fragility associated with the loss of dystrophin (Petofy et al., 1993) parallels the structural deficiencies observed in other cytolinker-deficient tissues, further demonstrating a close relationship between dystrophin and other cytolinkers. Collectively, these data support the hypothesis that dystrophin may function as a cytolinker in skeletal muscle.

Although dystrophin exhibits many characteristics of a cytolinker, a direct dystrophin–MT interaction has not been documented. Dystrophin lacks either a Gas2-related or a glycine-serine/arginine domain, but recent studies indicated that dystrophin at least indirectly influences MT organization or stability (Percival et al., 2007; Ayalon et al., 2008). For instance, the dystrophin-deficient mdx mouse exhibited MT disorganization in skeletal muscle with the costameric MTs most severely affected (Percival et al., 2007). Dystrophin’s enrichment at costameres...
and the restoration of costameric MT organization through virally mediated expression of a microdystrophin (Percival et al., 2007) indicates that dystrophin is necessary for proper costameric MT organization in skeletal muscle. Moreover, acute knockdown of ankyrin-B, a protein necessary for delivery of dystrophin to the sarcolemma and neuromuscular junction, caused the loss of costameric MTs and aberrant MT organization in a subset of MTs underlying the neuromuscular junction (Ayalon et al., 2008).

In this study, we investigated the hypothesis that dystrophin directly interacts with costameric MTs. We confirmed that costameric MTs were disrupted in dystrophin-deficient skeletal muscle and showed endogenous dystrophin cosedimented with MTs in tissue homogenates. Using purified proteins, we found that the carboxyl-terminal two thirds of dystrophin bound MTs with a Kd of 0.66 µM and stabilized MTs from cold-induced depolymerization. Finally, we documented a 2.5-fold increased expression of α- and β-tubulin without alteration in the tubulin--MT equilibrium in mdx skeletal muscle. These results demonstrate that dystrophin is a MT-associated protein (MAP) that stabilizes costameric MTs and functions as a costameric cytolinker in skeletal muscle.

Results and discussion

To determine whether dystrophin and MTs localize to similar structures in skeletal muscle, we conducted immunofluorescence analysis on teased extensor digitorum longus muscle fibers co-labeled with antidystrophin and anti–α-tubulin antibodies (Fig. 1, A and B). Dystrophin forms a subsarcolemmal network with transverse components along the I bands and the M line and with longitudinal components (Williams and Bloch, 1999), whereas MTs form a subsarcolemmal lattice, which in fast fibers, has transverse and longitudinal components plus an accumulation of MTs around myonuclei (Ralston et al., 1999). We found that the transverse MTs (Fig. 1 A, arrowheads) weave their course along the I band dystrophin staining for long distances. MTs were also transverse MTs (Fig. 1 A, arrowheads) weave their course along the I band dystrophin staining for long distances both transversely (arrowheads) and longitudinally (arrows). These data identify domains of the subsarcolemmal cytoskeleton where dystrophin and MTs may interact either directly or indirectly. Next, we examined MT organization in mouse models lacking dystrophin (mdx), dystrophin’s autosomal homologue utrophin (utrn<sup>−/−</sup>), or both dystrophin and utrophin (mdx/utrn<sup>−/−</sup>). Consistent with previous results (Percival et al., 2007), loss of dystrophin resulted in MT disorganization with the costameric MTs appearing to be most severely affected (Fig. 1 C) when compared with wild type (wt; Fig. 1 C). Ablation of utrophin had no effect on MT organization (Fig. 1 C), which is likely a result of its very low expression (Rybakov et al., 2002) and restriction to the neuromuscular junction (Ohtsuka et al., 1991). Finally, mdx/utrn<sup>−/−</sup> skeletal muscle exhibited MT disorganization comparable with that of mdx (Fig. 1 C). MT organization in 24-d-old prenecrotic mdx skeletal muscle fibers was also disorganized, whereas age-matched wt mice displayed a MT lattice nearly identical to mature wt mice (Fig. 1 D). Collectively, these results confirm a role for dystrophin in the stabilization and proper organization of costameric MTs independent of muscle necrosis and regeneration.

Figure 1. Dystrophin guides MTs at the surface of the muscle fibers and is necessary for proper MT organization. (A) Isolated muscle fibers from the extensor digitorum longus of 7-wk-old wt mice were costained for dystrophin (left) and α-tubulin (middle). The right panel shows that MTs (red) follow dystrophin (green) bands for long distances both transversely (arrowheads) and longitudinally (arrows). (B) At a higher magnification, dystrophin staining is granular; MTs are studded with dystrophin “dots.” Arrows indicate longitudinal MTs that follow dystrophin. (C) Muscle fibers from the extensor digitorum longus of 7-wk-old mdx, utrn<sup>−/−</sup>, and mdx/utrn<sup>−/−</sup> mice were stained with DM1A anti–α-tubulin and Hoechst dye. Both wt and utrn<sup>−/−</sup> fibers show the lattice of transverse and longitudinal MTs characteristic of fast fibers (arrowheads). In mdx and mdx/utrn<sup>−/−</sup> fibers, the regularity of the lattice is lost, and mostly oblique MTs originate from cytoplasmic nucleation points (arrows). (D) Peripherally nucleated prenecrotic muscle fibers from 24-d-old mdx mice also displayed MT disorganization, indicating that MT derangement occurred before muscle cell necrosis and regeneration. Bars: (A and B) 10 µm; (C and D) 20 µm.
rod domain absent from Dp260. A small amount of Dp260 pelleted in the absence of MTs, but substantially more Dp260 shifted to the pellet fraction when MTs were present (Fig. 3 A). After subtracting self-pelleting Dp260, Dp260 displayed a concentration-dependent and saturable cosedimentation with a Dp260/α-tubulin heterodimer stoichiometry of 1:1.4 and a Kd of 0.66 µM (Fig. 3 C). As predicted, DysNTerm-R10 did not cosediment with MTs up to concentrations approaching 10 µM (Fig. 3, B and C).

Next, we assessed how the presence of 1 µM Dp260 affected the tubulin–MT equilibrium in vitro. Dp260 had no significant effect on the fraction of tubulin in the MT fraction when incubated at room temperature (67.3 ± 0.72% vs. 68.6 ± 1.3%). However, the presence of Dp260 significantly increased the fraction of tubulin retained in the MT pellet (33.6 ± 2.9% vs. 42.2 ± 2.0%) when MTs were induced to depolymerize by incubating at 4°C. (Fig. 3, D and E). Collectively, these results demonstrate that dystrophin directly binds and stabilizes MTs from cold-induced depolymerization.

Because misregulation of other MAPs can alter tubulin expression and MT stability (Harada et al., 1994; Takahashi et al., 2003), we investigated how the loss of dystrophin affects the regulation of tubulin expression and the tubulin–MT equilibrium in skeletal muscle fibers. Tubulin levels in wt and mdx skeletal muscle extracts were examined by quantitative Western blot analysis. With mAb B512, we observed no difference...
in α-tubulin expression in wt and mdx skeletal muscle extracts (Fig. 4, A and B), which was consistent with what we (Prins et al., 2008) and others (Barton et al., 2002) reported previously. However, mAb DM1A showed an ∼2.5-fold increase in α-tubulin expression in mdx skeletal muscle (Fig. 4, A and B). Because levels of α- and β-tubulin are coregulated (Gonzalez-Garay and Cabral, 1995), we investigated β-tubulin levels to determine whether α-tubulin is up-regulated in mdx skeletal muscle. β-Tubulin expression was elevated 2.5-fold in mdx skeletal muscle (Fig. 4, A and B), suggesting that expression of both α- and β-tubulin is increased in mdx skeletal muscle. Thus, we conclude that mAb DM1A is able to recognize a population of α-tubulin not detected by mAb B512. To examine MT stability in mdx skeletal muscle, we analyzed levels of tyrosinated α-tubulin, a marker of dynamic MTs (Gundersen et al., 1984, 1987), and acetylated α-tubulin, a marker of long-lived MTs (Bulinski and Gundersen, 1991). The levels of tyrosinated α-tubulin were increased ∼2.5-fold in mdx extracts (Fig. 4, A and B), whereas the levels of acetylated α-tubulin were not (Fig. 4, A and B). The loss of dystrophin’s MT-stabilizing ability may explain why acetylated α-tubulin was not more abundant in mdx skeletal muscle extracts, but alterations in the tubulin–MT equilibrium could also explain the lack of more stable MTs. Therefore, we examined the tubulin–MT equilibrium in wt and mdx skeletal muscles and found that the loss of dystrophin did not affect the equilibrium (Fig. 4, C and D). Collectively, these results show that tubulins are misregulated in dystrophin-deficient skeletal muscle without affecting the tubulin–MT equilibrium. The loss of dystrophin’s MT-stabilizing ability likely explains why there are not more stabilized MTs even in the presence of more tubulin dimer in dystrophin-deficient skeletal muscle.

An indirect link between dystrophin and MTs mediated by ankyrin-B was recently shown to be important for proper trafficking of dystrophin and β-dystroglycan to the sarcolemma (Ayalon et al., 2008). However, costameric MTs are disorganized in mdx skeletal muscle even in the presence of properly localized ankyrin-B (Ayalon et al., 2008). Because the MT- and ankyrin-B-binding domains of dystrophin do not overlap (Fig. 5), our results and previous results suggest that dystrophin interacts with MTs in vivo through two distinct mechanisms. We propose that ankyrin-B delivers dystrophin to the sarcolemma dependent on MTs and that dystrophin and ankyrin-B collaborate to stabilize and organize MTs in skeletal muscle.

As with other cytolinkers, the ability to bind multiple components of the filamentous cytoskeleton likely allows dystrophin to protect the sarcolemma from mechanically induced damage. One highly truncated microdystrophin construct (∆R4-23) is very effective in restoring function in the dystrophin-deficient mdx mouse (Harper et al., 2002). Interestingly, the ∆R4-23 microdystrophin contains all sequences required for interaction with the three cytoskeletal filament systems: the amino-terminal tandem CH domain, which binds actin (Way et al., 1992) and cytokeratin filaments (Stone et al., 2005), the spectrinlike repeat 3 and the cysteine-rich regions, which are necessary for synemin intermediate filament binding (Bhosle et al., 2006), and the MT-binding domain. In contrast, Dp260 lacks the cytokeratin filament–binding domain and portions of the synemin- and actin-binding domains, which likely alters the binding affinities to both actin and synemin filaments and may explain why transgenic overexpression of Dp260 only partially alleviates the mdx
pathophysiology. For example, disorganized MTs are also associated with Golgi mislocalization (Percival et al., 2007), which in combination, would likely lead to impaired trafficking of membrane-bound proteins and may explain the decreased levels of β-dystroglycan and the sarcoglycans at the sarcolema of mdx skeletal muscle (Ohlendieck and Campbell, 1991). Because no MT knockout mouse has been generated, the exact function of MTs in skeletal muscle remains unknown. However, the importance of MTs in skeletal muscle biology is illustrated by the muscle weakness and increased levels of serum creatine kinase associated with colchicine toxicity in human patients (Boomershine, 2002; Caglar et al., 2003; Wilbur and Makowsky, 2004; Altman et al., 2007). Therefore, it is possible that derangement of the MT cytoskeleton contributes to some of the phenotypes associated with dystrophin deficiency.

**Materials and methods**

**Mice**

Control C57BL/6 and mdx mice were initially obtained from The Jackson Laboratory. The utrn⁻/⁻ and mdx/utrn⁻/⁻ mice were provided by D. Lowe (University of Minnesota, Minneapolis, MN). Mdx mice transgenically expressing Dp260 and ΔR4-R23 were provided by J. Chamberlain (University of Washington, Seattle, WA), and the Dp71 line was provided by J. Rafael-Fontney (Ohio State University, Columbus, OH). All animals were housed and treated following guidelines set by the University of Minnesota Institutional Animal Care and Use Committee.

**Antibodies**

The mAbs to α-tubulin (B512), β-tubulin (DM1A), and acetylated tubulin (6-11B-1) were purchased from Sigma-Aldrich. The mAb to α-tubulin (DM1A) was purchased from Abcam. The mAb to dystrophin (Dys2) was purchased from Novacastra. The polyclonal antibody to dystrophin (Rb2) was described previously (Rybakova et al., 1996). Infrared dye-conjugated anti-mouse secondary antibodies were purchased from LI-COR Biosciences.
Immuno- and fluorescence analysis
To analyze the MT lattice in dystrophic animal models, hindlegs of wt (3 and 8 wk), mdx (3, 5, and 8 wk), utm-/- (8–10 wk), and mdx:utm-/- (3, 5, and 8 wk) mice were skinned, cut as close as possible to the body, and fixed at room temperature for 2 h with 4% para-formaldehyde in phos- phate buffer. They were stored in phosphate buffer until the extensor digitorum longus muscle was dissected and separated with fine forceps into mostly single fibers. These were transferred to a 24-well tissue culture plate and incubated with mouse on mouse (MOM)–blocking buffer (Vector Labo- ratories) for 2 h at room temperature. Blocking buffer and every subse- quent buffer for incubation or washing contained 0.04% saponin for permeabilization and 0.05% sodium azide. Fibers were incubated over- night with mouse antitubulin (DM1A, 1:500; or B512, 1:4,000) in MOM diluent, washed three times for 20 min, and stained with 1:500 dilution of Alexa Fluor 488 anti-mouse and Alexa Fluor 568 anti–rabbit secondary antibodies (Invitrogen) in MOM diluent for 2 h at room temperature. After three 20-min washes, one of which contained the nuclear stain Hoechst 33342 (Sigma-Aldrich) at 2 µg/ml, fibers were mounted onto a glass slide in a drop of Vectashield (Vector Laboratories). Confocal images were captured with a 63× NA 1.4 oil immersion lens on a TCS SPS confocal micro- scope (Leica) in the Light Imaging Section of the National Institute of Arthritis and Musculoskeletal and Skin Diseases. Gain and laser power set- tings were adjusted to avoid saturation and use the whole linear range of fluorescence intensity. Unless specified, the parameters were adjusted for each new fiber imaged. The raw TIF images were transferred to a com- puter (Macintosh G5; Apple), opened in Photoshop (CS2; Adobe), assem- bled into montages, and adjusted for brightness when needed. The final illustrations give a faithful representation of the collected images.

Tissue MT cosedimentation assay
Tissue-based cosedimentation was performed as described previously (Hughes et al., 2008) with the following exception. The starting material was 200 µg of frozen skeletal muscle that was pulverized in a mortar and pestle cooled with liquid nitrogen then added to MT buffer (1% Triton X-100, 50 mM Hepes, 50 mM KCl, 1 mM MgCl2, 1 mM EGTA, 0.75 mM benz- amidine, 0.1 mM PMSF, 0.6 µg/ml pepstatin A, 0.5 µg/ml aprolin, 0.5 µg/ml leupeptin, iodoacetamide, and E64). The extracts were incu- bated for 1 h at 4°C and centrifuged at 100,000 g for 30 min at 25°C. A 1:1 ratio of both GTP and DTT was added to the soluble fraction of the extract and incubated at 37°C for 5 min. The extract was split into two fractions, one that was incubated on ice for 15 min, and 20 µM taxol was added to the other and incubated at 37°C for 15 min. 300 µl of each fraction was layered onto a cushion buffer (MT buffer plus 40% sucrose) and centrifuged at 100,000 g for 30 min at 25°C. The supernatant was removed, and the pellet fraction was resuspended in a Laemmli sample buffer.

Protein purification
A cDNA encoding Flag-tagged Dp260 (Warner et al., 2002) provided by J. Chamberlain was cloned into pFastBac1 to generate a recombinant baculovirus expression vector using previously described methods (Rybakova et al., 2002). Dp260 and dystrophin Nterm-R10 were expressed and purified with the baculovirus expression system and anti-Flag M2 affinity chroma- tography, respectively, as previously described (Rybakova et al., 2002), and 25 µg of skeletal muscle extract was subjected to SDS-PAGE and transferred to nitrocellulose membranes, which were washed/ blocked in a 5% milk solution in PBS for 1 h. The membranes were incu- bated overnight with primary antibody at room temperature. The primary antibodies and dilutions used were mAb Dys2 (1:50), mAb B512 (1:250), mAb D1MA (1:250; Sigma-Aldrich), mAb D66 (1:100), and mAb 6-11B-1 (1:100). Membranes were washed two times for 10 min in 5% milk solu- tion at room temperature, incubated with infrared dye–conjugated second- ary antibody (1:10,000) for 30 min at room temperature, and the membranes were washed in a 0.5% Tween solution in PBS two times for 10 min. Western blots were imaged and quantified with an infrared imag- ing system (Odyssey); LI-COR Biosciences). The Coomasie blue-stained posttransfer gel was analyzed densitometrically using UVP software and served as the loading control.

In vivo tubulin-MT equilibrium assay
The tibialis anterior was dissected, immediately placed in 1 ml of MT stabi- lization buffer (1% Triton X-100, 50% glycerol, 5% DMSO, 10 mM Na-HPO4, 0.5 mM EGTA, and 0.5 mM MgSO4), and homogenized with 10 strokes in a homogenizer. The resulting homogenate was centrifuged at 100,000 g for 30 min at 25°C. The soluble portion (tubulin containing) was saved for analysis, whereas the pellet portion (MT fraction) was re- suspended in 1 ml of 1% SDS buffer then boiled for 10 min. The pellet fraction was centrifuged at 13,000 g for 10 min, and the soluble portion was saved for analysis. 25 µl of the tubulin and MT fraction was analyzed and quantified via Western blot using mAb D1MA on the infrared imaging system (Odyssey).

Statistical analysis
All data are presented as mean ± SEM. Comparison between groups was performed using a t test with significance defined as P < 0.05.

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