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## Overexpression of SERCA1a in the *mdx* Diaphragm Reduces Susceptibility to Contraction-Induced Damage

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### Abstract

Although the precise pathophysiological mechanism of muscle damage in dystrophin-deficient muscle remains disputed, calcium appears to be a critical mediator of the dystrophic process. Duchenne muscular dystrophy patients and mouse models of dystrophin deficiency exhibit extensive abnormalities of calcium homeostasis, which we hypothesized would be mitigated by increased expression of the sarcoplasmic reticulum calcium pump. Neonatal adeno-associated virus gene transfer of sarcoplasmic reticulum ATPase 1a to the *mdx* diaphragm decreased centrally located nuclei and resulted in reduced susceptibility to eccentric contraction-induced damage at 6 months of age. As the diaphragm is the mouse muscle most representative of human disease, these results provide impetus for further investigation of therapeutic strategies aimed at enhanced cytosolic calcium removal.

### Disciplines

Comparative and Laboratory Animal Medicine | Medicine and Health Sciences | Musculoskeletal Diseases | Veterinary Medicine

## Overexpression of SERCA1a in the *mdx* Diaphragm Reduces Susceptibility to Contraction-Induced Damage

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### Abstract

Although the precise pathophysiological mechanism of muscle damage in dystrophin-deficient muscle remains disputed, calcium appears to be a critical mediator of the dystrophic process. Duchenne muscular dystrophy patients and mouse models of dystrophin deficiency exhibit extensive abnormalities of calcium homeostasis, which we hypothesized would be mitigated by increased expression of the sarcoplasmic reticulum calcium pump. Neonatal adeno-associated virus gene transfer of sarcoplasmic reticulum ATPase 1a to the *mdx* diaphragm decreased centrally located nuclei and resulted in reduced susceptibility to eccentric contraction-induced damage at 6 months of age. As the diaphragm is the mouse muscle most representative of human disease, these results provide impetus for further investigation of therapeutic strategies aimed at enhanced cytosolic calcium removal.

### Introduction

**D**UCHENNE MUSCULAR DYSTROPHY is due to the absence of dystrophin and characterized by progressive muscle degeneration, weakness, and death due to cardiopulmonary failure. Dystrophin serves as a mechanical link between the extracellular matrix and cytoskeleton and functions to stabilize the sarcolemma during contraction. The absence of dystrophin at the sarcolemmal membrane leads to a secondary loss of the dystrophin-associated glycoprotein complex, sarcolemmal instability, and increased susceptibility to muscle degradation following contraction (Petrof *et al.*, 1993). Although the precise mechanism of muscle damage has not been delineated, one hypothesis is that dystrophin deficiency promotes transient excess calcium influx following contraction and activates proteolytic breakdown of the cytoskeleton primarily by calpains (Deconinck and Dan, 2007; Clafin and Brooks, 2008; Zhang *et al.*, 2008). The source of the elevated calcium found in dystrophic muscle is controversial. Possible sources of excess calcium influx are tears in the plasma membrane, stretch-activated calcium channels, and/or leak channels (Bodensteiner and Engel, 1978; Fong *et al.*, 1990; Franco and Lansman, 1990; Millay *et al.*, 2009).

Extensive evidence suggests calcium homeostasis is dysfunctional in dystrophic muscle. Sarcoplasmic reticulum calcium ATPase (SERCA) activity is most likely not impaired

in dystrophic muscle, although disparate results have been reported depending on the muscle group examined and method used (Kargacin and Kargacin, 1996; Khammari *et al.*, 1998; Culligan *et al.*, 2002; Divet and Huchet-Cadiou, 2002; Plant and Lynch, 2003). Numerous abnormalities of SERCA and other calcium-handling proteins have been reported in dystrophin-deficient skeletal muscle. In the *mdx* mouse, levels of SERCA1a are elevated in the spared intrinsic laryngeal and toe muscles, unchanged in the mildly affected tibialis anterior, and reduced in the moderately affected extensor digitorum longus (EDL) muscle (Culligan *et al.*, 2002; Dowling *et al.*, 2002, 2003; Ferretti *et al.*, 2009). Similarly, calsequestrin, a high-capacity intraluminal calcium-binding protein, is elevated in intrinsic laryngeal muscle and reduced in the EDL (Ferretti *et al.*, 2009). Other reports describe decreased levels of calsequestrin-like protein, whose function is unknown but may contribute to reduced calcium-binding capacity in the dystrophic sarcoplasmic reticulum (Leberer *et al.*, 1988; Culligan *et al.*, 2002; Doran *et al.*, 2004).

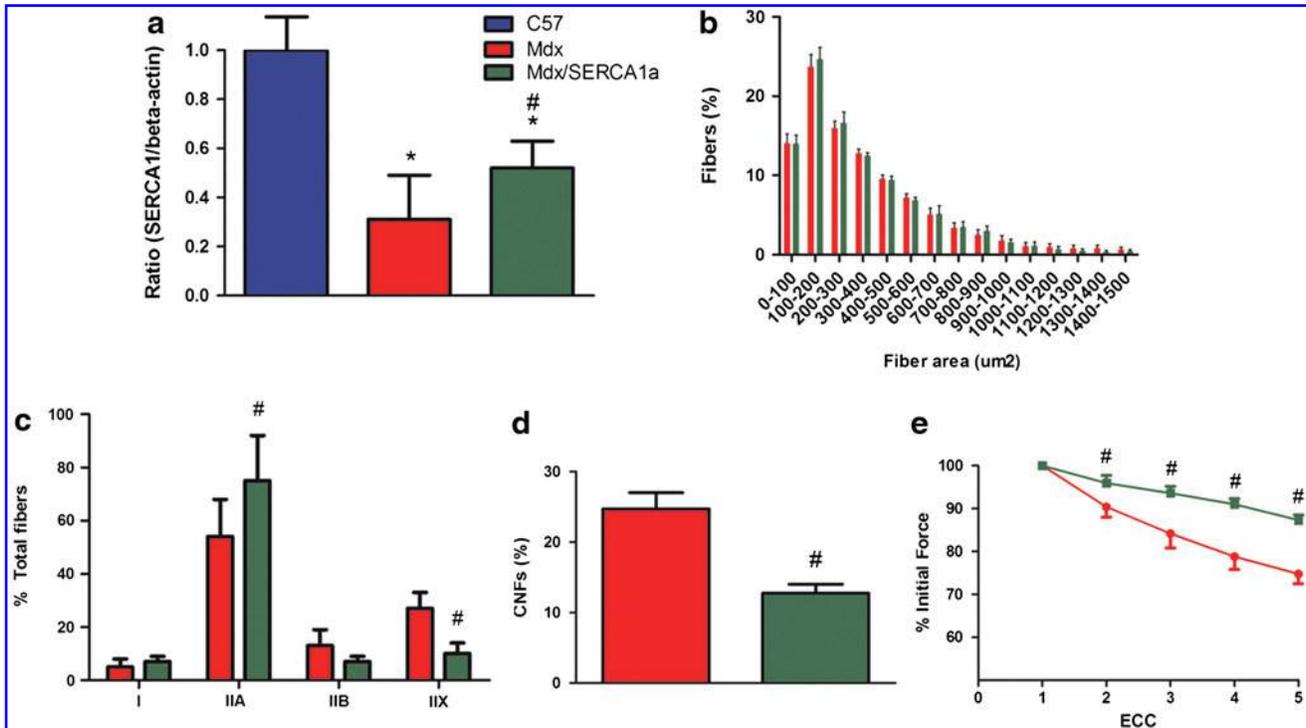
These observations led us to hypothesize that increased expression of sarcoplasmic reticulum ATPase 1a would reduce muscle damage following eccentric contraction. We used neonatal gene transfer of adeno-associated virus (AAV) pseudotype 2/6 to overexpress the fast skeletal muscle isoform of SERCA in the diaphragm of the dystrophin-deficient *mdx* mouse model and analyzed muscle function and

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**FIG. 1.** Overexpression of SERCA1a in the *mdx* diaphragm protects from eccentric contraction-induced loss of force production. (a) Immunoblotting of diaphragm homogenates demonstrated 69% less SERCA1a in the *mdx* diaphragm (red column) in comparison with C57 Bl/6 controls (navy column). AAV-mediated gene transfer increased SERCA1a content in *mdx*/SERCA1a diaphragm (green column) to 52% of control. (b) The muscle fiber size distribution was unaffected by SERCA1a overexpression. (c) An increase in the proportion of type IIA fibers and a decrease in the proportion of type IIX fibers were observed in the *mdx*/SERCA1a group. (d) The percentage of centrally nucleated fibers (CNFs) was reduced from 25% in the *mdx* diaphragm to 13% in the *mdx*/SERCA1a diaphragm. (e) Force production was measured following a series of eccentric contractions (ECC). Beginning at the second contraction, there was a significant attenuation of the force loss in *mdx*/SERCA1a diaphragm in comparison with the untreated *mdx* diaphragm. Data represent means  $\pm$  SD. \* $P < 0.05$ , *mdx* vs. C57/Bl6. # $P < 0.05$ , *mdx* vs. *mdx*/SERCA1a. Color images available online at [www.liebertonline.com/hum](http://www.liebertonline.com/hum).

morphology at 6 months of age. Increased SERCA1a protein levels in the diaphragm increased the proportion of type IIA fibers, reduced the percentage of centrally nucleated fibers, and attenuated the loss of force production following eccentric contractions.

## Materials and Methods

The use of mice in these experiments was approved by the University of Pennsylvania Animal Care and Use Committee. The viral construct used in this study consisted of the constitutive chicken  $\beta$ -actin promoter and the murine fast skeletal muscle isoform of SERCA (SERCA1a). AAV pseudotype 2/6 was produced by the University of Pennsylvania Vector Core, and 1E12 genome copies of virus in a total volume of 50  $\mu$ l was injected into neonatal mice (Bish *et al.*, 2008). At 6 months of age, muscle function was measured, and tissues were harvested for further analysis. Immunoblotting, muscle histology, and functional assessments were performed as described previously (Morine *et al.*, 2010), with the following modification for the assessment of eccentric contraction-induced force deficit. Following measurement of maximum isometric tetanus, the diaphragm was subjected to a series of five eccentric contractions with a 5-min rest between contractions. Muscles were stimulated

for a total of 700 msec and stretched 10% of optimum length in the final 200 msec of stimulation. The two-tailed unpaired Student's *t* test was used to compare means between groups; data are reported as means  $\pm$  SD.

## Results and Discussion

Neonatal *mdx* mice ( $n=9$  control *mdx*,  $n=9$  *mdx*/SERCA1a) were injected via a subxyphoid approach with 1E12 genome copies of AAV2/6 CB.SERCA1a to target transduction of the diaphragm (Bish *et al.*, 2008). *Mdx* diaphragm was found to contain approximately one third of the SERCA1a protein content of C57 Bl/6 muscle, whereas gene transfer restored SERCA1a protein levels to 52% of normal (Fig. 1a). Analysis of muscle morphology revealed that increased expression of SERCA1a did not alter fiber size, increased the proportion of type IIA fibers from 54% to 75%, and decreased the proportion of type IIX fibers from 27% to 10% (Fig. 1b and c). Quantification of centrally nucleated fibers, a marker of fibers that had previously undergone degeneration and regeneration, demonstrated a reduced percentage of centrally nucleated fibers in the *mdx*/SERCA1a diaphragm (Fig. 1d). Dystrophin-deficient muscle fibers are profoundly sensitive to eccentric muscle contractions and exhibit a rapid loss of force production following repetitive

stimulation (Petrof *et al.*, 1993). Following a series of five eccentric contractions, the *mdx*/SERCA1a diaphragm was partially protected from force loss (Fig. 1e). Specific force, or absolute force production normalized to cross-sectional area, was unchanged ( $9.4 \pm 0.7 \text{ N/cm}^2$  in control *mdx* vs.  $9.9 \pm 1 \text{ N/cm}^2$  in *mdx*/SERCA1a).

Here we show that SERCA1a overexpression was sufficient to ameliorate eccentric contraction-induced damage and improve some aspects of disease pathology in the *mdx* mouse model. The progression of dystrophic pathology is largely accounted for by the loss of muscle fibers following muscle contraction (Lynch, 2004). In dystrophin-deficient muscle, sarcolemmal stabilization achieved by dystrophin replacement or utrophin up-regulation prevents abnormal calcium influx and muscle damage due to contraction and slows or prevents disease progression (Harper *et al.*, 2002; Krag *et al.*, 2004; Liu *et al.*, 2005; Gregorevic *et al.*, 2006, 2008; Odom *et al.*, 2008). Modulation of sarcoplasmic reticulum calcium uptake activity thus represents a novel therapeutic target for muscular dystrophy.

The prevention of abnormal calcium influx and the activation of downstream effectors in muscular dystrophy have been explored in prior preclinical and clinical studies. In the 1980s, trials of the calcium-channel blockers nifedipine and diltiazem did not attenuate Duchenne muscular dystrophy in humans (Moxley *et al.*, 1987; Bertorini *et al.*, 1988). Although there is *in vitro* and *in vivo* evidence from *mdx* mice that the calcium-channel blockers diltiazem and verapamil can prevent calcium influx (Iwata *et al.*, 2005; Matsumura *et al.*, 2009), these agents predominantly act on L-type voltage-dependent calcium channels, and it has not been demonstrated that they act on other pathways of calcium influx implicated in muscular dystrophy, such as stretch-operated and store-operated channels. At doses sufficient to effect clinically meaningful blockade of calcium influx in dystrophic muscle, the untoward cardiovascular side effects of these agents may limit use in humans. Alternatively, selective inhibition of the degradative pathways activated by calcium, such as the calpain and ubiquitin-proteasome system, have shown some promise in preclinical studies (Spencer and Mellgren, 2002; Bonuccelli *et al.*, 2003; Burdi *et al.*, 2006; Briguet *et al.*, 2008).

An important limitation of our study is that transgene overexpression of SERCA1a was lost over time secondary to muscle-fiber turnover. Although we demonstrate that partial restoration of SERCA1a protein content to normal levels provides ~50% protection from contraction-induced damage, the diaphragm was not fully protected from cumulative muscle damage as specific force was not improved.

Prior evidence indicates that chronic calcium influx can activate the slow/oxidative gene program through the calcineurin/NFAT (nuclear factor of activated T cells) pathway (Chakkalakal *et al.*, 2003; McCullagh *et al.*, 2004; Calabria *et al.*, 2009). Hence, it would be expected that correction of the chronic calcium overload present in dystrophic muscle would prevent the fast to slow transition of myosin heavy-chain expression present in the *mdx* diaphragm. However, we found that SERCA1a overexpression led to an abundance of type IIA fibers in lieu of type IIX fibers. Type I and IIA fibers express significantly more utrophin than type IIB and IIX fibers (Chakkalakal *et al.*, 2003). In addition to enhanced calcium uptake afforded by SERCA1a overexpression, the

fiber-type switch and concomitant increase in utrophin expression may underlie the partial protection from eccentric contraction-induced damage observed in our study. Although increasing extrasynaptic utrophin content is unequivocally beneficial in dystrophin-deficient muscle, it is unclear if inducing the slow gene program in general would be a desirable aim in Duchenne muscular dystrophy patients. Studies of activated calcineurin in the mildly affected *mdx* mice and the severely affected  $\delta$ -sarcoglycan-deficient mouse have yielded contrasting results (Chakkalakal *et al.*, 2004; Parsons *et al.*, 2007). Differences in responsiveness to calcineurin inhibition or deletion in these models may reflect underlying differences in pathophysiology or disease severity. Future studies should examine the efficacy of SERCA overexpression in more severe dystrophic models and determine if inducing a fast to slow fiber-type transition is beneficial independent of utrophin up-regulation.

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### Author Disclosure Statement

No competing financial interests exist.

### References

- Bertorini, T.E., Palmieri, G.M., Griffin, J.W., Igarashi, M., McGee, J., Brown, R., Nutting, D.F., Hinton, A.B., and Karas, J.G. (1988). Effect of chronic treatment with the calcium antagonist diltiazem in Duchenne muscular dystrophy. *Neurology* 38, 609–613.
- Bish, L.T., Morine, K., Sleeper, M.M., Sanmiguel, J., Wu, D., Gao, G., Wilson, J.M., and Sweeney, H.L. (2008). Adeno-associated virus (AAV) serotype 9 provides global cardiac gene transfer superior to AAV1, AAV6, AAV7, and AAV8 in the mouse and rat. *Hum. Gene Ther.* 19, 1359–1368.
- Bodensteiner, J.B., and Engel, A.G. (1978). Intracellular calcium accumulation in Duchenne dystrophy and other myopathies: a study of 567,000 muscle fibers in 114 biopsies. *Neurology* 28, 439–446.
- Bonuccelli, G., Sotgia, F., Schubert, W., Park, D.S., Frank, P.G., Woodman, S.E., Insabato, L., Cammer, M., Minetti, C., and Lisanti, M.P. (2003). Proteasome inhibitor (MG-132) treatment of *mdx* mice rescues the expression and membrane localization of dystrophin and dystrophin-associated proteins. *Am. J. Pathol.* 163, 1663–1675.
- Briguet, A., Erb, M., Courdier-Fruh, I., Barzaghi, P., Santos, G., Herzner, H., Lescop, C., Siendt, H., Henneboehle, M., Weyermann, P., Magyar, J.P., Dubach-Powell, J., Metz, G., and Meier, T. (2008). Effect of calpain and proteasome inhibition on  $\text{Ca}^{2+}$ -dependent proteolysis and muscle histopathology in the *mdx* mouse. *FASEB J.* 22, 4190–4200.
- Burdi, R., Didonna, M.P., Pignol, B., Nico, B., Mangieri, D., Rolland, J.F., Camerino, C., Zallone, A., Ferro, P., Andreatta, F., Confalonieri, P., and De Luca, A. (2006). First evaluation of the potential effectiveness in muscular dystrophy of a novel chimeric compound, BN 82270, acting as calpain-inhibitor and anti-oxidant. *Neuromuscul. Disord.* 16, 237–248.
- Calabria, E., Ciciliot, S., Moretti, I., Garcia, M., Picard, A., Dyar, K.A., Pallafacchina, G., Tothova, J., Schiaffino, S., and Murgia, M. (2009). NFAT isoforms control activity-dependent muscle

- fiber type specification. *Proc. Natl. Acad. Sci. U.S.A.* 106, 13335–13340.
- Chakkalakal, J.V., Stocksley, M.A., Harrison, M.A., Angus, L.M., Deschenes-Furry, J., St-Pierre, S., Megoney, L.A., Chin, E.R., Michel, R.N., and Jasmin, B.J. (2003). Expression of utrophin A mRNA correlates with the oxidative capacity of skeletal muscle fiber types and is regulated by calcineurin/NFAT signaling. *Proc. Natl. Acad. Sci. U.S.A.* 100, 7791–7796.
- Chakkalakal, J.V., Harrison, M.A., Carbonetto, S., Chin, E., Michel, R.N., and Jasmin, B.J. (2004). Stimulation of calcineurin signaling attenuates the dystrophic pathology in mdx mice. *Hum. Mol. Genet.* 13, 379–388.
- Clafin, D.R., and Brooks, S.V. (2008). Direct observation of failing fibers in muscles of dystrophic mice provides mechanistic insight into muscular dystrophy. *Am. J. Physiol. Cell Physiol.* 294, C651–658.
- Culligan, K., Banville, N., Dowling, P., and Ohlendieck, K. (2002). Drastic reduction of calsequestrin-like proteins and impaired calcium binding in dystrophic mdx muscle. *J. Appl. Physiol.* 92, 435–445.
- Deconinck, N., and Dan, B. (2007). Pathophysiology of Duchenne muscular dystrophy: current hypotheses. *Pediatr. Neurol.* 36, 1–7.
- Divet, A., and Huchet-Cadiou, C. (2002). Sarcoplasmic reticulum function in slow- and fast-twitch skeletal muscles from mdx mice. *Pflugers Arch.* 444, 634–643.
- Doran, P., Dowling, P., Lohan, J., McDonnell, K., Poetsch, S., and Ohlendieck, K. (2004). Subproteomics analysis of Ca<sup>2+</sup>-binding proteins demonstrates decreased calsequestrin expression in dystrophic mouse skeletal muscle. *Eur. J. Biochem.* 271, 3943–3952.
- Dowling, P., Culligan, K., and Ohlendieck, K. (2002). Distal mdx muscle groups exhibiting up-regulation of utrophin and rescue of dystrophin-associated glycoproteins exemplify a protected phenotype in muscular dystrophy. *Naturwissenschaften* 89, 75–78.
- Dowling, P., Lohan, J., and Ohlendieck, K. (2003). Comparative analysis of Dp427-deficient mdx tissues shows that the milder dystrophic phenotype of extraocular and toe muscle fibres is associated with a persistent expression of beta-dystroglycan. *Eur. J. Cell Biol.* 82, 222–230.
- Ferretti, R., Marques, M.J., Pertille, A., and Santo Neto, H. (2009). Sarcoplasmic-endoplasmic-reticulum Ca<sup>2+</sup>-ATPase and calsequestrin are overexpressed in spared intrinsic laryngeal muscles of dystrophin-deficient mdx mice. *Muscle Nerve* 39, 609–615.
- Fong, P.Y., Turner, P.R., Denetclaw, W.F., and Steinhart, R.A. (1990). Increased activity of calcium leak channels in myotubes of Duchenne human and mdx mouse origin. *Science* 250, 673–676.
- Franco, A., Jr., and Lansman, J.B. (1990). Calcium entry through stretch-inactivated ion channels in mdx myotubes. *Nature* 344, 670–673.
- Gregorevic, P., Allen, J.M., Minami, E., Blankinship, M.J., Haraguchi, M., Meuse, L., Finn, E., Adams, M.E., Froehner, S.C., Murry, C.E., and Chamberlain, J.S. (2006). rAAV6-microdystrophin preserves muscle function and extends lifespan in severely dystrophic mice. *Nat. Med.* 12, 787–789.
- Gregorevic, P., Blankinship, M.J., Allen, J.M., and Chamberlain, J.S. (2008). Systemic microdystrophin gene delivery improves skeletal muscle structure and function in old dystrophic mdx mice. *Mol. Ther.* 16, 657–664.
- Harper, S.Q., Hauser, M.A., DelloRusso, C., Duan, D., Crawford, R.W., Phelps, S.F., Harper, H.A., Robinson, A.S., Engelhardt, J.F., Brooks, S.V., and Chamberlain, J.S. (2002). Modular flexibility of dystrophin: implications for gene therapy of Duchenne muscular dystrophy. *Nat. Med.* 8, 253–261.
- Iwata, Y., Katanosaka, Y., Shijun, Z., Kobayashi, Y., Hanada, H., Shigekawa, M., and Wakabayashi, S. (2005). Protective effects of Ca<sup>2+</sup> handling drugs against abnormal Ca<sup>2+</sup> homeostasis and cell damage in myopathic skeletal muscle cells. *Biochem. Pharmacol.* 70, 740–751.
- Kargacin, M.E., and Kargacin, G.J. (1996). The sarcoplasmic reticulum calcium pump is functionally altered in dystrophic muscle. *Biochim. Biophys. Acta* 1290, 4–8.
- Khamari, A., Pereon, Y., Baudet, S., and Noireaud, J. (1998). In situ study of the sarcoplasmic reticulum function in control and mdx mouse diaphragm muscle. *Can. J. Physiol. Pharmacol.* 76, 1161–1165.
- Krag, T.O., Bogdanovich, S., Jensen, C.J., Fischer, M.D., Hansen-Schwartz, J., Javazon, E.H., Flake, A.W., Edvinsson, L., and Khurana, T.S. (2004). Heregulin ameliorates the dystrophic phenotype in mdx mice. *Proc. Natl. Acad. Sci. U.S.A.* 101, 13856–13860.
- Leberer, E., Hartner, K.T., and Pette, D. (1988). Postnatal development of Ca<sup>2+</sup>-sequestration by the sarcoplasmic reticulum of fast and slow muscles in normal and dystrophic mice. *Eur. J. Biochem.* 174, 247–253.
- Liu, M., Yue, Y., Harper, S.Q., Grange, R.W., Chamberlain, J.S., and Duan, D. (2005). Adeno-associated virus-mediated microdystrophin expression protects young mdx muscle from contraction-induced injury. *Mol. Ther.* 11, 245–256.
- Lynch, G.S. (2004). Role of contraction-induced injury in the mechanisms of muscle damage in muscular dystrophy. *Clin. Exp. Pharmacol. Physiol.* 31, 557–561.
- Matsumura, C.Y., Pertille, A., Albuquerque, T.C., Santo Neto, H., and Marques, M.J. (2009). Diltiazem and verapamil protect dystrophin-deficient muscle fibers of MDX mice from degeneration: a potential role in calcium buffering and sarcolemmal stability. *Muscle Nerve* 39, 167–176.
- McCullagh, K.J., Calabria, E., Pallafacchina, G., Ciciliot, S., Serrano, A.L., Argentini, C., Kahlvovde, J.M., Lomo, T., and Schiaffino, S. (2004). NFAT is a nerve activity sensor in skeletal muscle and controls activity-dependent myosin switching. *Proc. Natl. Acad. Sci. U.S.A.* 101, 10590–10595.
- Millay, D.P., Goonasekera, S.A., Sargent, M.A., Maillat, M., Aronow, B.J., and Molkenin, J.D. (2009). Calcium influx is sufficient to induce muscular dystrophy through a TRPC-dependent mechanism. *Proc. Natl. Acad. Sci. U.S.A.* 106, 19023–19028.
- Morine, K.J., Bish, L.T., Pendrak, K., Sleeper, M.M., Barton, E.R., and Sweeney, H.L. (2010). Systemic myostatin inhibition via liver-targeted gene transfer in normal and dystrophic mice. *PLoS One* 5, e9176.
- Moxley, R.T., 3rd, Brooke, M.H., Fenichel, G.M., Mendell, J.R., Griggs, R.C., Miller, J.P., Province, M.A., and Patterson, V. (1987). Clinical investigation in Duchenne dystrophy. VI. Double-blind controlled trial of nifedipine. *Muscle Nerve* 10, 22–33.
- Odom, G.L., Gregorevic, P., Allen, J.M., Finn, E., and Chamberlain, J.S. (2008). Microdystrophin delivery through rAAV6 increases lifespan and improves muscle function in dystrophic dystrophin/utrophin-deficient mice. *Mol. Ther.* 16, 1539–1545.
- Parsons, S.A., Millay, D.P., Sargent, M.A., Naya, F.J., McNally, E.M., Sweeney, H.L., and Molkenin, J.D. (2007). Genetic disruption of calcineurin improves skeletal muscle pathology and

- cardiac disease in a mouse model of limb-girdle muscular dystrophy. *J. Biol. Chem.* 282, 10068–10078.
- Petrof, B.J., Shrager, J.B., Stedman, H.H., Kelly, A.M., and Sweeney, H.L. (1993). Dystrophin protects the sarcolemma from stresses developed during muscle contraction. *Proc. Natl. Acad. Sci. U.S.A.* 90, 3710–3714.
- Plant, D.R., and Lynch, G.S. (2003). Depolarization-induced contraction and SR function in mechanically skinned muscle fibers from dystrophic *mdx* mice. *Am. J. Physiol. Cell Physiol.* 285, C522–C528.
- Spencer, M.J., and Mellgren, R.L. (2002). Overexpression of a calpastatin transgene in *mdx* muscle reduces dystrophic pathology. *Hum. Mol. Genet.* 11, 2645–2655.
- Zhang, B.T., Yeung, S.S., Allen, D.G., Qin, L., and Yeung, E.W. (2008). Role of the calcium-calpain pathway in cytoskeletal damage after eccentric contractions. *J. Appl. Physiol.* 105, 352–357.

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3. Stefan M. Gehrig, Chris van der Poel, Timothy A. Sayer, Jonathan D. Schertzer, Darren C. Henstridge, Jarrod E. Church, Severine Lamon, Aaron P. Russell, Kay E. Davies, Mark A. Febbraio, Gordon S. Lynch. 2012. Hsp72 preserves muscle function and slows progression of severe muscular dystrophy. *Nature* . [[CrossRef](#)]
4. Sanjeewa A. Goonasekera, Chi K. Lam, Douglas P. Millay, Michelle A. Sargent, Roger J. Hajjar, Evangelia G. Kranias, Jeffery D. Molkenkin. 2011. Mitigation of muscular dystrophy in mice by SERCA overexpression in skeletal muscle. *Journal of Clinical Investigation* **121**:3, 1044-1052. [[CrossRef](#)]
5. Virginie Carmignac, Madeleine Durbeej. 2011. Cell-matrix interactions in muscle disease. *The Journal of Pathology* n/a-n/a. [[CrossRef](#)]
6. Joshua T. Selsby. 2010. Increased catalase expression improves muscle function in mdx mice. *Experimental Physiology* no-no. [[CrossRef](#)]