

EUK-134 ameliorates nNOS μ translocation and skeletal muscle fiber atrophy during short-term mechanical unloading

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AQ:2 Lawler JM, Kunst M, Hord JM, Lee Y, Joshi K, Botchlett RE, Ramirez A, Martinez DA. EUK-134 ameliorates nNOS μ translocation and skeletal muscle fiber atrophy during short-term mechanical unloading. *Am J Physiol Regul Integr Comp Physiol* 306: R000–R000, 2014. First published January 29, 2014; doi:10.1152/ajpregu.00371.2013.—Reduced mechanical loading during bedrest, spaceflight, and casting, causes rapid morphological changes in skeletal muscle: fiber atrophy and reduction of slow-twitch fibers. An emerging signaling event in response to unloading is the translocation of neuronal nitric oxide synthase (nNOS μ) from the sarcolemma to the cytosol. We used EUK-134, a cell-permeable mimetic of superoxide dismutase and catalase, to test the role of redox signaling in nNOS μ translocation and muscle fiber atrophy as a result of short-term (54 h) hindlimb unloading. Fischer-344 rats were divided into ambulatory control, hindlimb-unloaded (HU), and hindlimb-unloaded + EUK-134 (HU-EUK) groups. EUK-134 mitigated the unloading-induced phenotype, including muscle fiber atrophy and muscle fiber-type shift from slow to fast. nNOS μ immunolocalization at the sarcolemma of the soleus was reduced with HU, while nNOS μ protein content in the cytosol increased with unloading. Translocation of nNOS from the sarcolemma to cytosol was virtually abolished by EUK-134. EUK-134 also mitigated dephosphorylation at Thr-32 of FoxO3a during HU. Hindlimb unloading elevated oxidative stress (4-hydroxynonenal) and increased sarcolemmal localization of Nox2 subunits gp91phox (Nox2) and p47phox, effects normalized by EUK-134. Thus, our findings are consistent with the hypothesis that oxidative stress triggers nNOS μ translocation from the sarcolemma and FoxO3a dephosphorylation as an early event during mechanical unloading. Thus, redox signaling may serve as a biological switch for nNOS to initiate morphological changes in skeletal muscle fibers.

atrophy; disuse; skeletal muscle; nNOS; oxidative stress; FoxO3a

SKELETAL MUSCLE IS A HIGHLY specialized and adaptable mesodermic tissue capable of rapid remodeling in response to changes in mechanical loading and stretch (28, 38). Transmittance and detection of loading in skeletal muscle are paramount in regulating differentiation, cell growth, and protein turnover. The ability to sense and relay loading (i.e., mechanotransduction) is, in part, performed by proteins adjacent to and spanning cell membranes (i.e., sarcolemma) in skeletal muscle (47). Mechanical unloading that occurs with limb casting, splinting, bed rest, and spaceflight, elicits a substantial loss of force-generating capacity, linked to a diminishment in muscle fiber

cross-sectional area or atrophy (12, 23). The unloading phenotype also includes a shift of a portion of skeletal muscle fibers from slow-twitch to fast-twitch (12). Muscle atrophy that occurs with unloading is coupled to a net loss of contractile proteins, a function of increased protein degradation combined with a decrease in protein synthesis (12, 23). Recent studies emphasize the importance of proteolytic pathways, including ubiquitin proteasome, calpains, autophagy, and caspase-3 in triggering the early stages of atrophy (4, 10, 11, 26, 34, 35, 39). For example, elevation of ubiquitin ligases with unloading is initiated via dephosphorylation of FoxO3a and activation of nuclear factor- κ B (NF- κ B) (26, 34, 42).

Elegant experiments by Suzuki et al. (42) ushered in a new paradigm, whereby disassembly of skeletal muscle in response to mechanical unloading is accelerated by translocation of neuronal nitric oxide synthase (nNOS) from the sarcolemma to the sarcoplasm. The mu (μ)-splice variant of nNOS is a key signaling protein located in the dystrophin-glycoprotein complex (DGC) of skeletal muscle (29, 42), anchored through binding of a PDX motif at its NH₂ terminus to α and β -syntrophins, and, thus, to dystrophin and dystrobrevin (13, 29). Fourteen days of mechanical unloading of the soleus resulted in movement of nNOS μ to the sarcoplasm, where nNOS μ directly led to dephosphorylation of FoxO3a at Thr-32, and stimulated ubiquitin ligases and atrophy (42). Interestingly, the loss of nNOS μ from the DGC also occurs in Duchenne muscular dystrophy (DMD), where it may exacerbate further disruption of the DGC proteins, thus increasing susceptibility to muscle damage and inflammation (19, 43). Furthermore, knockout of the nNOS μ gene has been recently shown to elicit myopathy and a reduction in contractile force (29). In addition, new data from Ito et al. (15) demonstrate that nNOS activation of the NADPH oxidase isoform Nox4 contributes to hypertrophy when skeletal muscle is overloaded. These data suggest in total that nNOS μ may serve as an effector of dynamic changes in mechanical loading, regardless of whether loading is increased or decreased.

While the mechanisms underlying translocation of nNOS are unknown, increase in reactive oxygen species (ROS) and subsequent oxidative stress is possible. For example, oxidative stress from NADPH oxidase and possibly mitochondrial sources play a role in damage in the *mdx* mouse model of Duchenne muscular dystrophy, where nNOS that is lost from the sarcolemma contributes directly to pathology (17, 45). Elevated skeletal muscle ROS from mitochondrial sources during mechanical unloading has been reported by Powers and colleagues (27). Recent data suggest that the Nox2 isoform of

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NADPH oxidase, a gp91phox containing NADPH oxidase, is important in stretch-induced Ca^{2+} dyshomeostasis and damage (17, 45). NADPH oxidase is a family of membrane-bound oxidoreductases in nonphagocytic and phagocytic cells that produce superoxide anions (O_2^-) in response to mechanical strain, hypoxia, and damage (17). Elevated oxidative stress from mitochondria and possibly Nox2 could result in a loss or translocation of nNOS from the sarcolemma (21, 41).

Therefore, we hypothesized that translocation of nNOS μ during an early stage of mechanical unloading would be linked to changes in oxidative stress and downstream dephosphorylation of FoxO3a at Thr-32. We further postulated that attenuation of oxidative stress would mitigate translocation of nNOS from the sarcolemmal region and significantly ameliorate two predominant phenotypic changes that begin early in mechanical unloading: 1) muscle fiber atrophy and 2) fiber-type shift from slow to fast. Previous antioxidant supplementation studies using nonspecific scavengers have yielded inconsistent findings in reducing morphological responses to unloading (3, 6, 20). In contrast, transfection of the antioxidant enzyme catalase yielded significant protection against atrophy (9). To test the importance of redox signaling, we used a novel cell-permeable, saline-manganese compound (EUK-134), which mimics the catalytic function of superoxide dismutase (SOD) coupled with catalase. In addition, to target redox signaling that is active early in the unloading process, and, therefore, involvement in initiation of atrophy, we chose a 54-h period of hindlimb unloading when oxidative stress is peaking (4).

METHODS

Animals. Young adult (4–6 mo) Fischer-344 (F344) rats were used to model disuse via hindlimb unloading. Animals were housed and cared for in accordance with National Institutes of Health policy (NIH; DHEW publication no. 85–23, revised 1985). Procedures were approved by the Institutional Animal Care and Use Committee at Texas A&M University and meet all federal requirements, as defined in the Animal Welfare Act (AWA) and the Public Health Service Policy and the Humane Care and Use of Laboratory Animals. Rat chow and water were provided ad libitum, and the animals were maintained in a temperature-controlled room ($23 \pm 2^\circ\text{C}$) with a 12:12-h light-dark cycle. Rodents are valuable mammalian models for muscle wasting that occur in human disuse and unloading, with changes in muscle fiber cross-sectional area, weakness, and fiber-type shift from slow to fast, resembling responses in humans.

Hindlimb unloading. An adaptation of the rodent hindlimb unloading model (7) was utilized to induce mechanical unloading. Briefly, the hindlimbs of the hindlimb unloading (HU) group were elevated to a spinal orientation of 55° above horizontal. We secured the tail with flexible, orthopedic tape around the middle one-third of the tail. The tail harness was attached to a linear mobile system with the line and adjusted, so that the hindlimb paws were suspended, while the rats were free to ambulate around the cage with their forelimbs for food and water.

Experimental design. A hindlimb unloading period was chosen to target the early phase of skeletal muscle remodeling during hindlimb unloading. Markers of atrophy and oxidative stress are first seen in the hindlimb unloading model for limb muscles at 2–3 days (4). We used a cell-permeable, catalytic saline-manganese compound developed by Eukarion (EUK-134) as an intervention for elevated oxidative stress. Adult F344 rats were divided into three groups ($n = 8/\text{group}$): loaded controls (CON), 54 h of hindlimb unloading, and hindlimb unloading + 3 $\text{mg}\cdot\text{kg}^{-1}\cdot\text{day}^{-1}$ ip of the SOD/catalase mimetic EUK-134 (HU-

EUK). This dosage used for EUK-134 was similar to that used by Bianca et al. (5). Administration of EUK-134 commenced 24 h prior to hindlimb unloading. The EUK series mimics the antioxidant enzymes superoxide dismutase (SOD) and catalase and, thus, remove superoxide anions (O_2^-) and hydrogen peroxide (H_2O_2). SOD/catalase mimetics have high specificity for O_2^- , H_2O_2 , and low toxicity with potential for translational use (32). EUK-134 had no effect on endogenous antioxidant enzyme (e.g., MnSOD and Cu, ZnSOD) protein levels (data not shown).

Skeletal muscle preparation. The soleus was chosen as a model of skeletal muscle response to mechanical unloading. As a postural muscle with a high percentage of Type I (slow-twitch) fibers, the soleus is susceptible to rapid atrophy and remodeling when exposed to mechanical unloading (7). Following a dose of pentobarbital sodium (100 mg/kg), the soleus muscles were carefully dissected, cleaned, rinsed with Krebs-Ringer solution, laid along the longitudinal axis on a polyurethane splint, and then frozen in isopentane cooled in liquid nitrogen for histological analysis or directly in liquid nitrogen for protein assays. Muscle samples were ground into homogenates with a glass-on-glass pestle at a 26:1 dilution of lysis buffer cooled in melting ice (4°C). The lysis buffer contained 20 mM HEPES free acid, 10 mM HEPES Na salt, 350 mM mannitol, 10% glycerol, 25 mM KCl, and 0.5 mM EDTA. Homogenates were centrifuged 10 min at 3,000 g at 4°C , and *supernatant 1* was removed as the soluble fraction. The pellet was resuspended (9:1 vol/vol) in buffer containing 20 mM HEPES free acid, 20 mM HEPES Na salt, 350 mM NaCl, 10% glycerol, 1 mM MgCl_2 , and 0.5 mM EDTA, and then centrifuged at 12,000 g for 30 min at 4°C . *Supernatant 1* was then centrifuged at 20,000 g for 10 min at 4°C . *Supernatant 2* from this spin served as the cytosolic fraction and was verified using MnSOD. The remaining pellet was the crude membrane fraction, which was resuspended in buffer containing 20 mM HEPES free acid, 20 mM HEPES Na salt, 150 mM NaCl, 300 mM sucrose, 10% glycerol, and 1 mM MgCl_2 , and 0.5 mM EDTA. Protein content was assessed using a Pierce BCA protein assay kit.

Fiber-type analysis and morphology. We adapted a method for fiber-type identification previously described by Kanatous et al. (16). Transverse muscle sections from the soleus of six animals per group were cut (10 μm) in a cryotome at -15°C and then air-dried for 30 min. Sections were placed in acidic ($\text{pH} = 4.25$) or alkaline ($\text{pH} = 10.4$) preincubation medium. The sections were rinsed thrice in 100 mM Tris buffer ($\text{pH} = 7.8$) with 18 mM CaCl_2 for 2 min each. Slides were then placed in an ATP incubation solution ($\text{pH} = 9.4$) for 25 min at room temperature, and rinsed 3 times in 1% CaCl_2 solution. Sections were counterstained in 0.1% toluidine blue solution for 90 s and then briefly washed in dH₂O. Sections were dehydrated in 95% ethanol, twice in 100% ethanol, and then cleared in xylene 2 times (5 min each). Coverslips were then mounted with Vectamount (Vector Laboratories, Burlingame, CA). Type I fibers stain lightly with alkaline preincubation. Type II fibers have differential lability under acidic conditions where Type IIa stain lightest and Type IIb/d fibers stain medium blue. Delp and Duan (8) previously estimated the fiber-type distribution in Fischer-344 rats as 84% Type I, 7% Type IIa, and 9% Type IIb.

Muscle cross-sectional areas were measured, recorded, and calibrated against a stage micrometer. Images were captured on a Zeiss Axioplot Vision-series microscope, and quantified using the NIH ImageJ program. Six soleus muscles per group were analyzed with each fiber cross-sectional area in square micrometer units calculated using ImageJ against the stage micrometer standard. There were ~400–600 fibers/muscle quantified, with three groups of six samples, for a total of nearly 9,000 fibers analyzed for morphology and fiber type.

Histochemistry and immunohistochemistry. For histochemical and immunohistochemistry (IHC) assays, muscle samples were mounted and embedded in optimum cutting temperature (OCT) medium. Cross sections were cut at a temperature of -15°C in a Shandon Cryotome

FSE (Thermo-Fisher, Waltham, MA) and dried for 30 min. For visualization via hematoxylin-and-eosin (H&E) staining, each section had a brief gentle rinse of dH₂O after drying. Briefly, soleus sections were then incubated with alum hematoxylin for 4 min at room temperature and rinsed with tap water to enhance bluing of nuclei. Differentiation was induced to remove background with 75% ethanol and 0.3% glacial acetic acid (acid-alcohol solution) for 15 min in a Coplar jar. Following another brief rinse in tap water, sections were incubated in eosin Y for 4 min at room temperature. Samples were then dehydrated with two changes of 95% ethanol and two short incubations (2 min) in 100% ethanol. Slides were cleared in xylene and sealed under a cover slip in mounting media (Vector Laboratories). Images were captured on an Olympus IX-51 (Olympus Imaging America, Center Valley, PA) inverted epifluorescence microscope.

For immunohistochemistry assays, muscle cross sections were cut at -15°C and fixed in acetone for 60 min. Sections were then blocked in TBS with 0.05% Tween 20 and 10% serum from a source matched with the host of the secondary antibody. The appropriate primary antibody for nNOS (1:50 dilution; Cayman 160870), gp91phox (1:100; SC-5827; specific for Nox2; Santa Cruz Biotechnology, Santa Cruz, CA), and p47phox (1:100; Santa Cruz SC-7660) were applied in blocking buffer and placed on each section for an hour. After washing in PBS with 0.1% Tween 20, biotinylated secondary antibody from Vector Laboratories was applied to the sections (1:200 dilution) and incubated in PBS buffer for 30 min. Cross sections were stained for 30 min with Vectastain Elite ABC reagent and incubated in NovaRed peroxidase substrate solution (Vector Laboratories) for 5 min.

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For nNOS localization, IHCs were also run using a TSA (tyramide signal amplification) kit (Perkin-Elmer, Waltham, MA) to enhance localization of both sarcolemmal and cytosolic locations of nNOS, as recommended by Punkt et al. (31). Within each protein of interest, slides from samples in all three groups were completed during the same day. Images were taken on a Zeiss Axioplot Vision-series microscope (Carl Zeiss Microimaging, Thornwood, NY). Membrane protein localization for p47phox and gp91phox was quantified using an edge filter macrodeveloped for the NIH ImageJ software program. Samples in each group were analyzed under identical conditions.

Immunofluorescence. Soleus sections were cut 10 µm thick in a cryostat at -15°C and placed onto microscope slides. Sections were dried for 30 min and fixed in methanol (-20°C) for 10 min, then acetone (-20°C) for 10 min. After 20 min of air-drying, samples were washed for 5 min 3 times in PBS. Sections were then blocked in TBS with 0.05% Tween 20, and 10% goat serum for 15 min. After air drying for 10 min, primary antibodies (1:100 dilution) for 4-hydroxynonenal (Calbiochem; no. 393206) and laminin (Sigma; no. L8271) were applied and incubated for 60 min in blocking buffer at 25°C. After three 10-min washes in PBS, sections were incubated in the appropriate secondary antibody (1:200 dilution) with a fluorochrome attached (e.g., goat anti-rabbit Alexa Fluor 488, goat anti-mouse Alexa Fluor 594) for 30 min at room temperature. Sections are washed thrice in PBS, and air-dried for 20 min. Slides were then mounted with Prolong Gold anti-fade medium (Invitrogen/Life Technologies, Grand Island, NY). Images were captured on a Zeiss Axioplot Vision series microscope. The antibody used for 4-hydroxynonenal is specific for any protein adducted or attached to 4-HNE, a stable breakdown product of lipid peroxidation. Therefore, visual 4-HNE adducts were used as a marker of damage or alteration of muscle proteins by oxidative stress. 4-HNE and laminin localization staining for all three groups were completed in a single day's run under the same conditions. We also verified sarcolemmal localization of Nox2 subunits (gp91phox, p47phox) using β-sarcoglycan, a membrane protein integrated into the dystrophin-glycoprotein complex.

Membrane protein localization for 4-hydroxynonenol (4-HNE) was quantified using an edge filter macro developed for the NIH ImageJ software program. Cell membranes were traced using the membrane-binding protein laminin as a standard. Each sample among

groups was analyzed using the same procedure. The edge filter width was set at 5 µm.

Western immunoblot. Protein abundance was determined by Western immunoblot analysis in the cytosolic and crude membrane fractions. Twenty micrograms of protein were loaded on 10% polyacrylamide gels, and electrophoresed using a Bio-Rad Protein III gel-box. Briefly, separating gel (375 mM Tris-HCl; pH = 8.8; 0.4% SDS; 10% acrylamide) and stacking gel (125 mM Tris-HCl; pH = 6.8; 0.4% SDS; 10% acrylamide monomer) solutions were made, and polymerization was initiated by TEMED and ammonium persulfate. Separating and stacking gels were quickly poured into a Bio-Rad Protein III gel-box (Bio-Rad, Hercules, CA). Soleus samples in sample buffer (Tris pH = 6.8 with 2% SDS, 30 mM DTT, 25% glycerol) were then loaded into the wells of the gels, and electrophoresed at 150 V. The gels were transferred at 30 V overnight onto a nitrocellulose membrane (Bio-Rad). Membranes were blocked in 5% nonfat milk in PBS with 0.1% Tween-20 for 6 h and then incubated in the appropriate 1° Ab. Antibodies for nNOS (BD: 610309) and FoxO3a phosphorylation at Thr-32 (Cell Signaling: 9464) (1:1,000) were incubated for 12 h at room temperature. Following three washings in PBS with 0.1% Tween-20, membranes were incubated in a horseradish peroxidase-conjugated secondary antibody for 90 min. Then an enhanced chemiluminescence detection system (Amersham, Piscataway, NJ) was used for visualization. Densitometry and quantification were performed using the NIH ImageJ software program. The product of blot intensity and area was used to identify abundance. To ensure equal loading of protein, Ponceau-S-staining was performed for each membrane to verify loading, and the lane background reading was subtracted from each protein blot density reading. Membranes were stripped and reprobed for GAPDH or dystrophin when normalizing for abundance vs. a control protein. Both GAPDH and β-actin were tested as loading controls. Neither GAPDH nor β-actin exhibited any mean differences in protein abundance (Fig. 10).

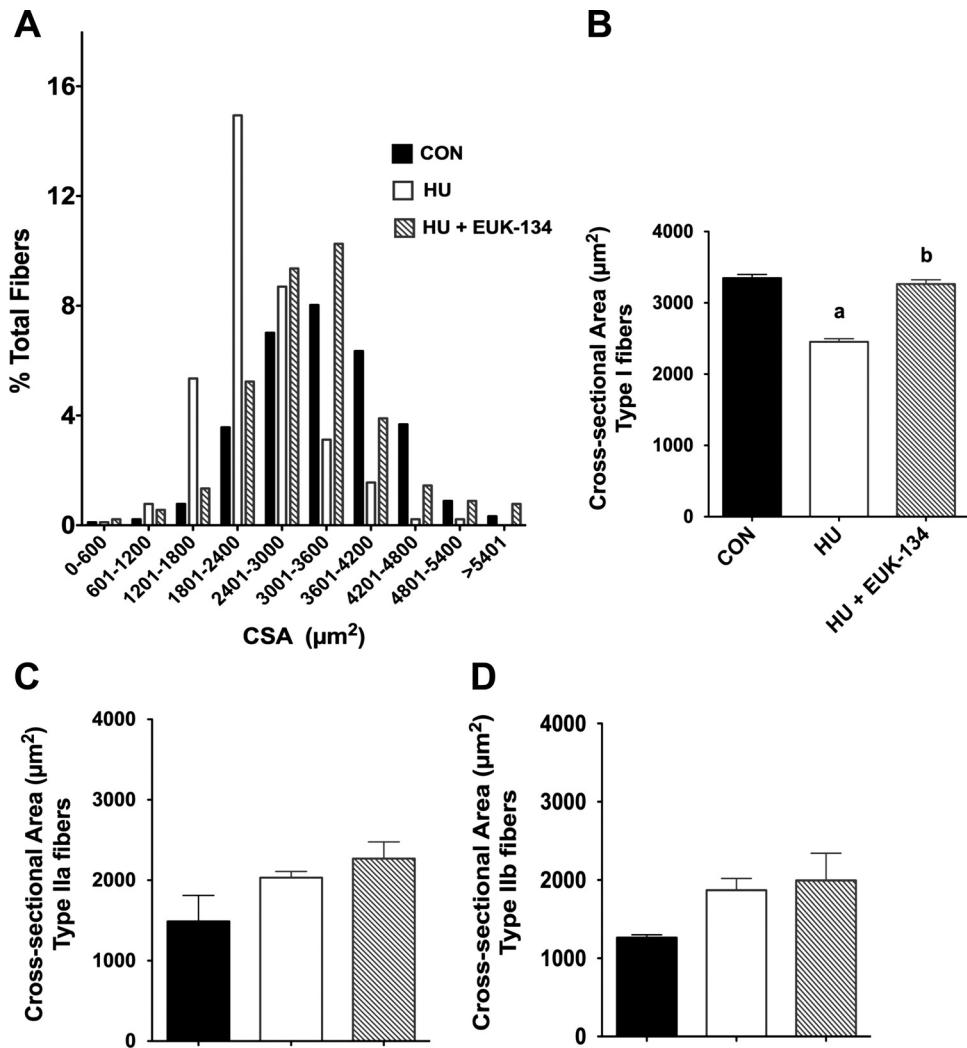
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Statistics. A one-way ANOVA was conducted for protein content, mean muscle cross-sectional area (CSA), and fiber-type with Fisher-LSD run post hoc to determine group mean differences. Analyses were verified for normal distribution with variance homogenously distributed among groups. A Chi-square test was used for determining group differences among muscle fiber CSA profiles. Statistical significance was set at $P < 0.05$.

RESULTS

Alterations in soleus muscle morphology with 54 h of mechanical unloading are regulated by oxidative stress. In the first experiment, we tested the hypothesis that reactive oxygen species were a causal factor in the unloading-induced phenotype: reduction of muscle fiber cross-sectional and partial shift of fiber type from slow-to-fast twitch. Previously, nitric oxide synthase inhibition was shown to regulate muscle hypertrophy and fiber-type shift in response to overloading (15, 38). In addition, inhibition or genetic ablation of nNOS substantially reduced proteolysis and muscle fiber atrophy during mechanical unloading (42). nNOSμ may, indeed, be responsible for guiding skeletal muscle remodeling in response to a dynamic change in mechanical loading (42).

Body masses (means ± SD) were not statistically different among ambulatory control (435.1 ± 17.6 g), hindlimb unloaded (437.4 ± 22.4 g), and hindlimb unloaded + EUK-134 (415.8 ± 22.5 g). While the EUK-134 trended lower, their mass also trended lower prior to the commencement of the study. Soleus-to-body mass ratio trended lower for HU (158.6 ± 11.4 mg/kg) than CON (165.5 ± 15.6 mg/kg) or HU+EUK (163.8 ± 9.27 mg/kg) but did not reach statistical significance. Total protein concentration was significantly lower in HU



($1,160 \pm 126 \mu\text{g/g}$) than CON ($1,649 \pm 90 \mu\text{g/g}$), suggestive of greater water and/or reduced protein levels during the early phase of mechanical unloading. However, HU + EUK total protein content ($1,302 \pm 175 \mu\text{g/g}$) was not significantly lower than that of controls.

We quantified muscle CSA and observed that the fiber cross-sectional area profile (Fig. 1A) for the soleus exposed to hindlimb unloading shifted to the left, illustrating a marked reduction in muscle fiber cross-sectional area (Fig. 1B). The greatest number of soleus fibers was found in the $1,800\text{--}2,400 \mu\text{m}^2$ range after unloading, in contrast with the $3,000\text{--}3,600 \mu\text{m}^2$ range in the ambulatory control group. Remarkably, muscle fiber atrophy was virtually completely abolished by

EUK-134 (Fig. 1A). Interestingly, we found that EUK-134 reduced muscle fiber atrophy primarily in Type I fibers (Fig. 1, A and B) but offered no protection in Type IIa and Type IIb/d fibers (Fig. 1, C and D).

To determine whether EUK-134 affected muscle mass and fiber cross-sectional area, independent of unloading, we conducted a set of follow-up control experiments. We tested 10 additional rats under loaded, ambulatory conditions, in which rats received three saline or EUK-134 injections over a 78-h period. This strategy covered the 54-h unloading period plus the injection, given 24 h prior to HU. Body weights, soleus mass (Fig. 2A), and the ratio of muscle mass to body mass (Fig. 2B) were not different between the saline and EUK-134 groups. In

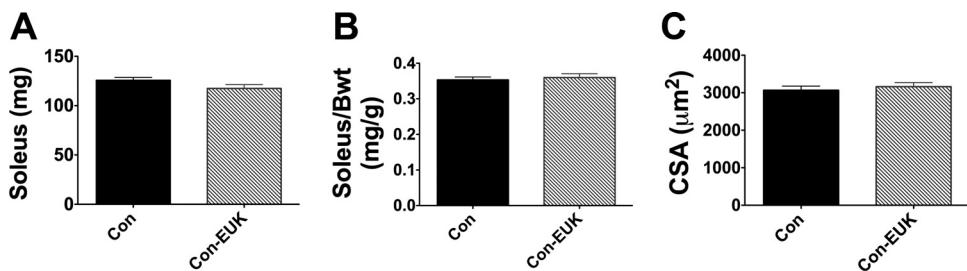


Fig. 2. EUK-134 has no effect on soleus mass or fiber cross-sectional area in ambulatory rats. Seventy-eight hours EUK-134 or saline injection had no differential effect on soleus muscle mass (A), soleus mass/body mass ratio (B), or soleus muscle cross-sectional area (C).

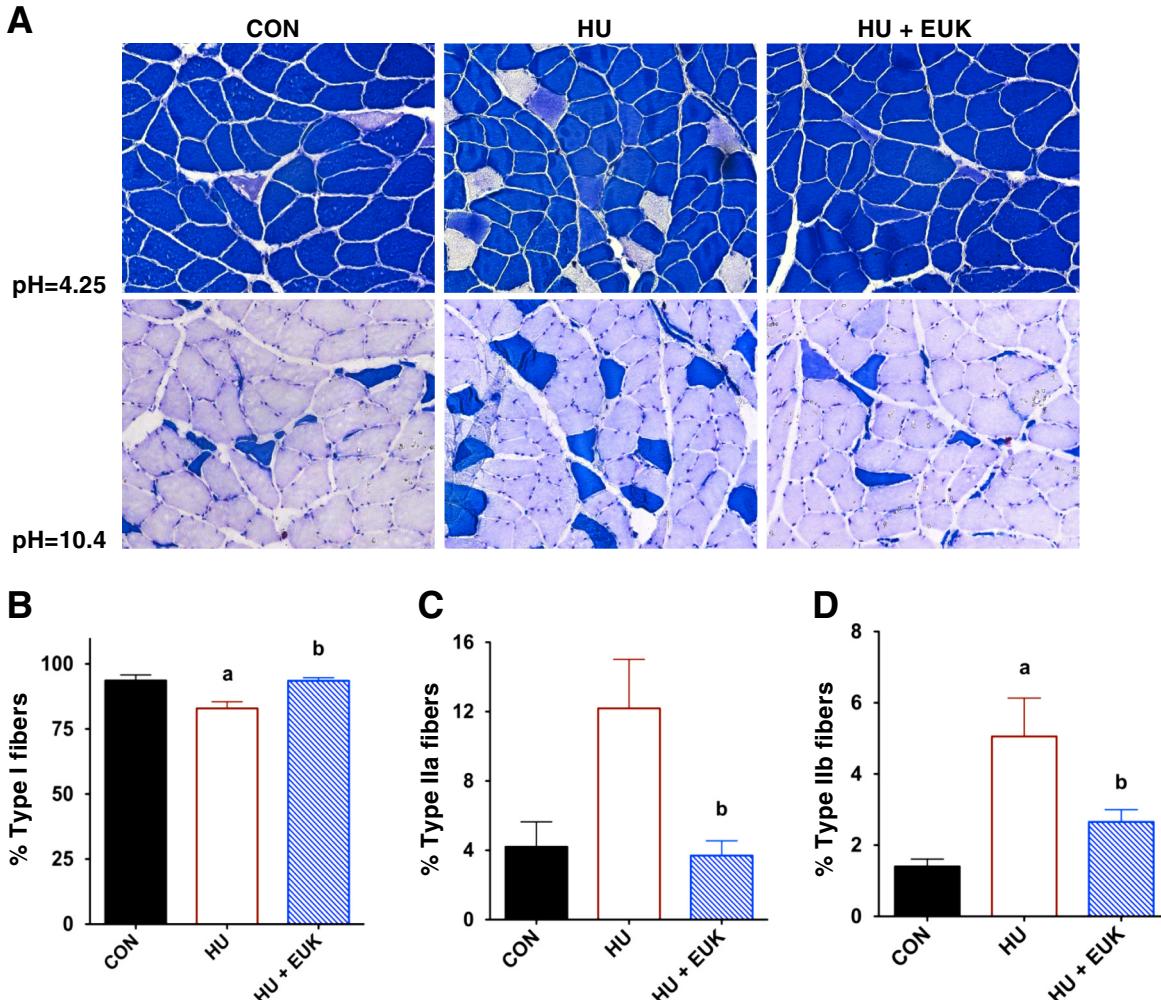


Fig. 3. Unloading-induced fiber-type shift is redox regulated in CON, HU, and HU-EUK rats ($n = 8/\text{group}$). A modified ATPase-staining technique revealed that significant shift in muscle soleus muscle fiber-type from fast to slow in the soleus as a result of HU. *A*: representative sections for both alkaline (pH = 10.4) and acidic (pH = 4.25) preincubations are presented. A reduction of slow-twitch fibers (Type I) was noted with HU (*B*) ($P < 0.05$), accompanied by an elevation in Type IIa (*C*), and Type IIb/d fibers (*D*). EUK-134 abrogated the unloading-induced fiber-type shift (*A–D*). ^{a,b}Letters that differ are significantly different from each other ($P < 0.05$).

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addition, means for skeletal muscle fiber cross-sectional area (Fig. 2C) did not significantly differ between saline and EUK-134-injected rats when loaded with normal ambulation. These data are consistent with the notion that EUK-134's effects on skeletal muscle mass were specific to the unloading protocol.

We tested the hypothesis that unloading-induced shift of skeletal muscle fiber-type from slow to fast was dependent on oxidative stress. Representative ATPase stains with acid and alkaline preincubation that are labile for Type II and Type I fibers, respectively, are displayed in Fig. 3A. Mechanical unloading caused a significant reduction in the percentage of Type I fibers, as expected (Fig. 3B). In addition, hindlimb unloading also caused an elevation in the percentage of Type IIa- (Fig. 3C) and Type IIb/d (Fig. 3D)-positive fibers. Remarkably, EUK-134 virtually halted the unloading-induced reduction in slow-twitch or Type I fibers within the rat soleus (Fig. 3, *A* and *B*). Consistent with the effect on Type I fibers, EUK-134 also prevented elevation of fast-twitch Type IIa (Fig. 3C) and Type IIb/d fibers observed with unloading (Fig. 3D). To our knowledge, these are the first data to indicate that redox

signaling plays a significant role in the fiber-type shift that occurs in response to mechanical unloading.

EUK-134 attenuates nNOS translocation with hindlimb unloading. In the next set of experiments, we tested the hypothesis that perturbations in the location of nNOS μ would be reduced by EUK-134. Hematoxylin-and-eosin stains are presented in Fig. 4A. Using immunohistochemistry, we initially observed substantial delocalization of nNOS from the sarcolemma as a result of short-term hindlimb unloading (Fig. 4, *B* and *C*). Suzuki et al. (42) had previously reported that nNOS leaves the sarcolemma and is dispersed through the sarcoplasm after 14 days of hindlimb unloading. Figure 4B features IHCs for nNOS without tyramide signaling amplification (TSA), and Fig. 1C with TSA. Utilizing both methods, we found that localization of nNOS at the sarcolemma was dramatically reduced in the soleus by 54 h of hindlimb unloading. However, cytosolic nNOS localization (29) was more readily observed with TSA with the greatest positive staining in the sarcoplasm within the HU + saline group. Remarkably, 3 days of injections with EUK-134 virtually prevented the loss

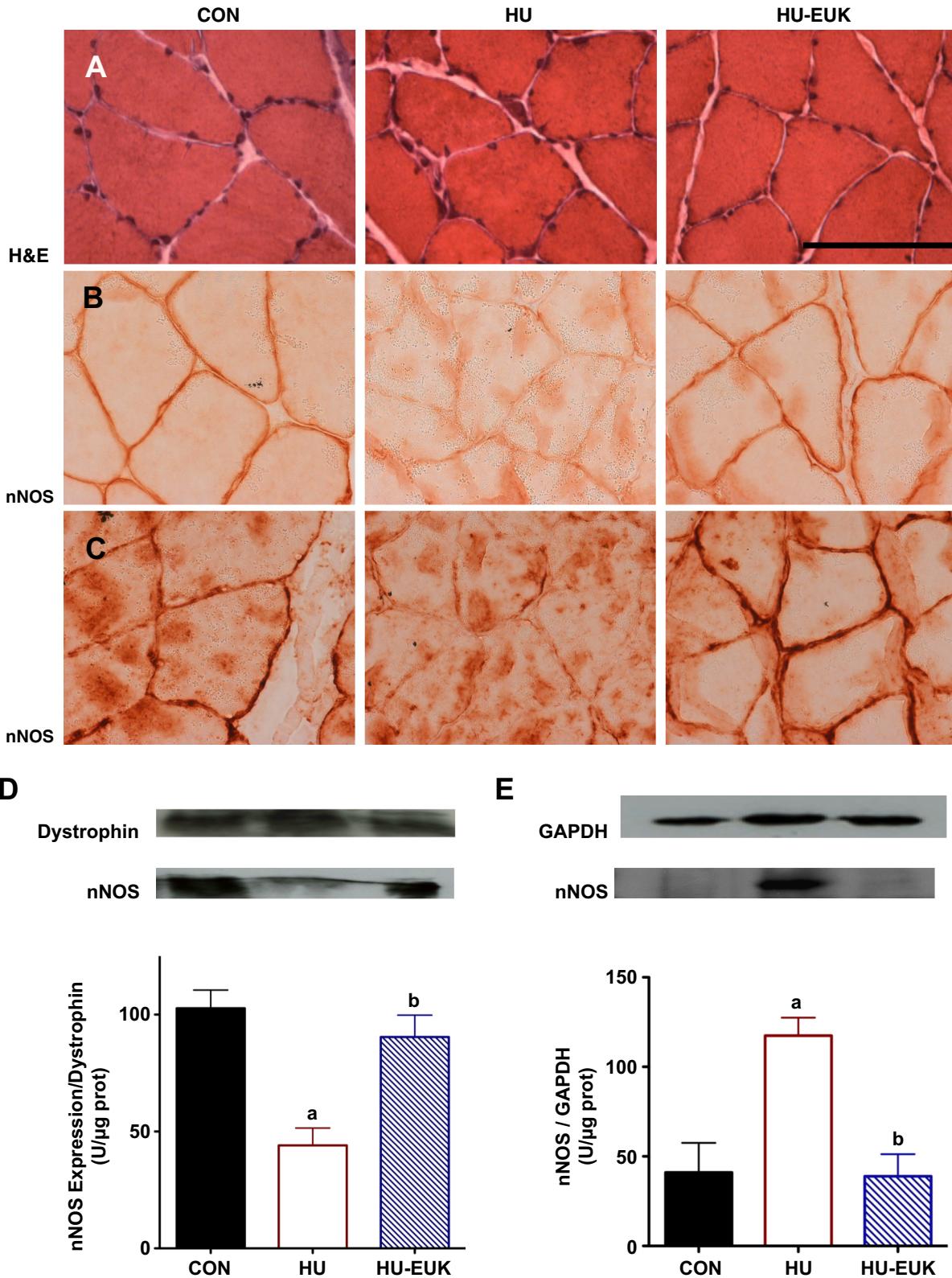


Fig. 4. EUK-134 reduces dislocation of nNOS μ from the sarcolemma of the soleus from HU Fischer-344 rats. Hematoxylin-and-eosin (H&E)-stained muscle cross sections are visualized in Fig. 4 (left) for control (CON), HU (middle), and HU + EUK-134 (HU-EUK) ($n = 6$ /group; right). Immunolocalization of neuronal nitric oxide synthase (nNOS) in the sarcolemma was lost in the rat soleus muscle as a result of HU, an effect abrogated by the catalytic SOD/catalase mimetic EUK-134 as visualized without tyramide signaling amplification (TSA) (B) and with TSA (C). Protein immunoblotting for nNOS was analyzed from membrane (D) and cytosolic (E) fractions. nNOS protein abundance per expression of dystrophin decreased with HU and was protected by EUK-134. EUK-134 attenuated HU-induced elevation in the cytosolic fraction. ^aSignificantly different ($P < 0.01$) than the CON group. ^bSignificantly different ($P < 0.01$) than the HU group. Scale bar = 100 μ m.

of nNOS-positive staining from the sarcolemmal region in Fischer-344 rats (Fig. 4, *B* and *C*). Indeed, there was robust positive staining of nNOS on the soleus muscle fiber periphery in the EUK-treated group, similar to ambulatory control rats injected with saline. Thus, we provide the first evidence that dislocation of nNOS μ may occur in less than 3 days of mechanical unloading in the rat soleus muscle.

nNOS protein abundance measured via immunoblot paralleled our IHC findings. nNOS levels in the membrane fraction levels, when equated for dystrophin, significantly decreased in the hindlimb unloading group when compared with ambulatory controls (Fig. 4, *D* and *E*). Treatment with EUK-134 prevented the reduction in membrane nNOS levels observed with HU, consistent with localization results visualized in Fig. 4, *B* and *C*. In contrast, nNOS protein abundance in soluble, cytosolic fraction, equated per GAPDH levels, increased significantly in the HU group (Fig. 4*B*). Remarkably, EUK-134 prevented the unloading-induced increase in cytosolic nNOS protein abundance. Our immunohistological and protein localization data, therefore, clearly demonstrate that nNOS μ is, indeed, translocated away from the sarcolemma within the first 54 h of mechanical unloading. Furthermore, translocation of nNOS μ with mechanical unloading appeared to be redox-signaling dependent.

EUK-134 protects against unloading-related reduction in FoxO3a phosphorylation (Thr-32). Important regulators of muscle atrophy with mechanical unloading include FoxO3a via dephosphorylation and NF- κ B via phosphorylation of its inhibitor regulator I κ B (I κ B) (35). Suzuki et al. (42) demonstrated that dephosphorylation and activation of FoxO3a was dependent upon nNOS translocation from the sarcolemma to the cytosol. Given that EUK-134 attenuated nNOS translocation with 54 h of HU, we tested the hypothesis that EUK-134 would also attenuate loss of phosphorylation at Thr-32 for FoxO3a. Levels of FoxO3a phosphorylation (Thr-32) significantly decreased with the short-term unloading protocol (Fig. 5), as expected. EUK-134 treatment resulted in an increase in phosphorylation levels at Thr-32 for FoxO3a during HU that were significantly greater than the soleus of unloaded rats injected with saline. In addition, FoxO3a phosphorylation was not significantly different in the HU group injected with EUK-134.

Unloading-induced elevation of oxidative stress and Nox2 is reduced by EUK-134. To determine whether EUK-134, indeed, ameliorated unloading-induced oxidative stress in the soleus muscle, we tracked 4-HNE-positive imaging. Previously, elevation of oxidative stress in response to mechanical unloading has been reported in both hindlimb unloading and mechanical ventilation models (23, 24, 26). In the current study, we found that 4-HNE protein adducts increased in the soleus following 54 h of hindlimb unloading (Fig. 6*A*). Interestingly, there was increased localization of 4-HNE around the sarcolemma with short-term unloading, using laminin immunofluorescence in parallel images to confirm localization (Fig. 6*B*). Furthermore, elevation in soleus 4-HNE fluorescence caused by hindlimb unloading was partially abrogated by EUK-134 treatment (Fig. 6*A*), confirming the inhibitory effect of EUK-134 on oxidative stress.

We also measured NF- κ B using an ELISA technique previously outlined (22). However, we saw no effects of HU or

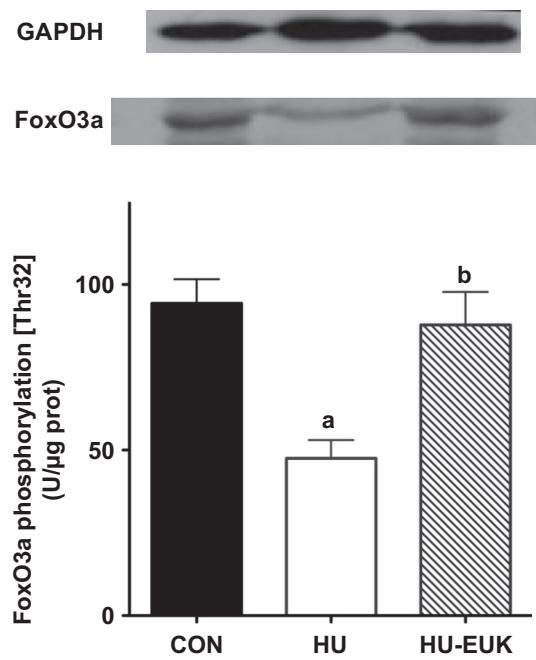


Fig. 5. Effect of EUK-134 on FoxO3a phosphorylation. Effect of EUK-134 on FoxO3a phosphorylation at Thr-32 in the soleus of rats that were hindlimb-unloaded for 54 h in CON, HU, and HU-EUK rats ($n = 6$ /group). ^aSignificantly different than the CON group ($P < 0.01$). ^bSignificantly different than the HU group ($P < 0.05$).

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EUK-134 (data not shown), suggesting the importance of NF- κ B likely increases as unloading duration increases.

Given the potential for high oxidative stress in the sarcolemmal microenvironment (14) during mechanical unloading, 4-HNE oxidation, and upregulation of Nox2 noted in DMD (46), we also determined immunolocalization of two Nox2 subunits: gp91phox (Nox2) and p47phox. Short-term hindlimb unloading, indeed, elevated immunostaining intensity at the sarcolemma of the p47phox-positive in the soleus of F344 rats (Fig. 7, *B* and *D*). The EUK-134 group displayed lower gp91phox-positive sarcolemmal localization, while p47phox sarcolemmal intensity levels in the EUK-134 group were not significantly different than controls (Fig. 7, *A–C*). Membrane-positive staining for gp91phox was demonstrated using immunofluorescence against β -sarcoglycan as our membrane proteins. gp91phox, β -sarcoglycan, merged images, and negative control are pictured in Fig. 8. We also determined p47phox protein abundance using Western immunoblot analysis (Fig. 9). p47phox levels were significantly upregulated in both HU groups. Combined with our DHE data, this is, to our knowledge, the first evidence that the Nox2 isoform of NADPH oxidase in skeletal muscle is elevated in the first 3 days of mechanical unloading.

Both GAPDH and β -actin were tested as loading controls. Neither GAPDH nor β -actin exhibited any mean differences in protein abundance (Fig. 10).

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DISCUSSION

In this study, EUK-134, a mimetic of superoxide dismutase and catalase, attenuated the mechanical unloading or “space-flight phenotype” in skeletal muscle in the hindlimb-unloading ground model: muscle fiber atrophy and slow-twitch to fast-

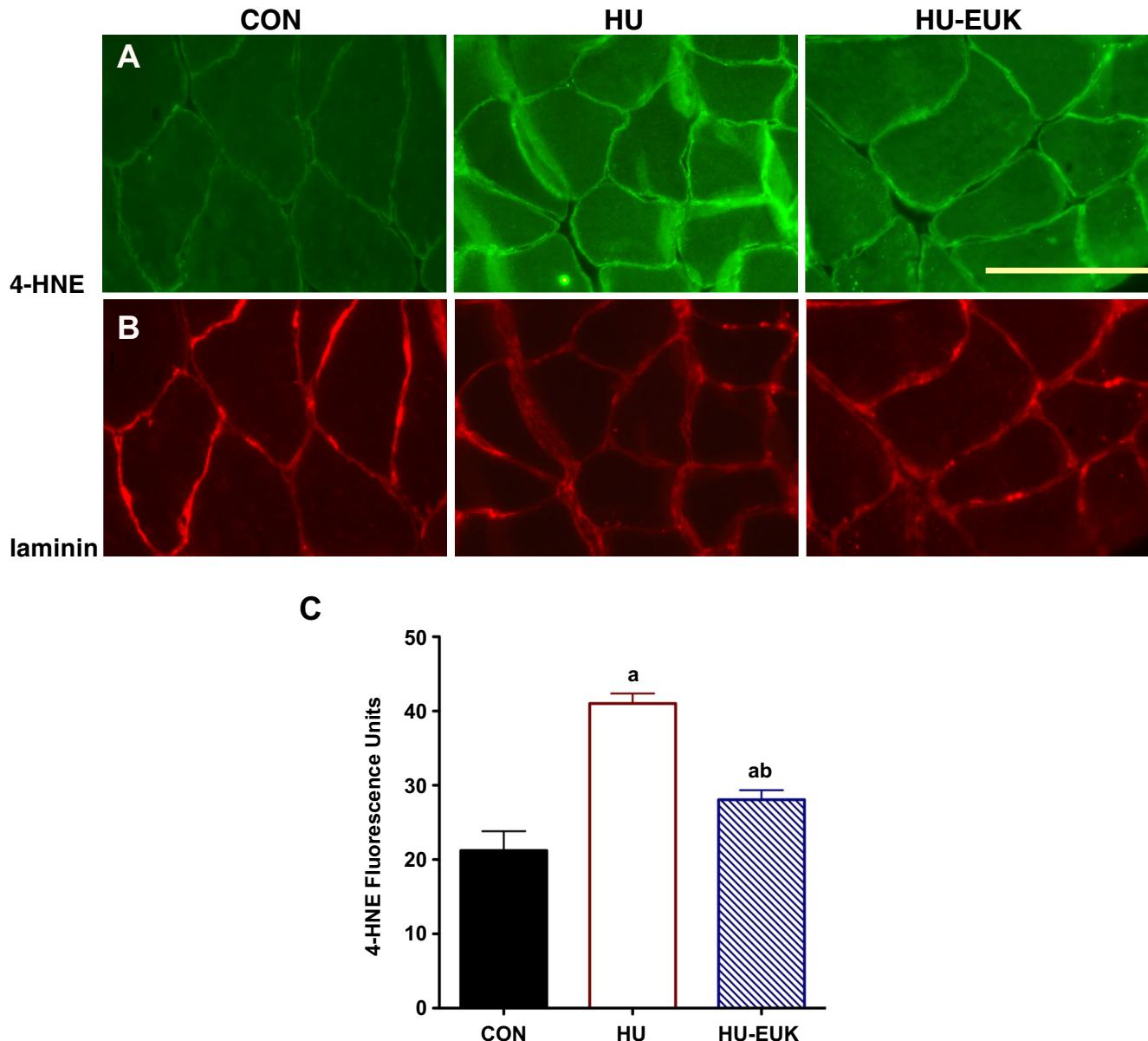


Fig. 6. EUK-134 reduces 4-hydroxynonenal (4-HNE) in the HU soleus. We demonstrated that oxidative stress (A) was increased by hindlimb unloading using 4-hydroxynonenal as a marker. Highest positive staining occurred near the sarcolemma as localized via immunofluorescence with laminin (B) used as a control for identifying muscle cells. Quantification of 4-HNE fluorescence intensity can be found in Fig. 6C. ^aSignificantly different than the CON group ($P < 0.001$). ^bSignificantly different than the HU group ($P < 0.01$).

twitch fiber-type shift. EUK-134 protection of unloading-induced muscle atrophy was linked to abrogation of nNOS translocation away from the sarcolemma. Furthermore, EUK-134 also prevented dephosphorylation of FoxO3a at Thr-32, previously shown to be regulated by nNOS translocation. EUK-134 reduced oxidative stress and upregulation of Nox2. A discussion of the principle finding follows.

EUK-134 attenuation of nNOS translocation and morphological changes. Our data indicate that unloading-induced translocation of nNOS μ away from the sarcolemma is part of an early adaptation to alterations in mechanotransduction during the unloading process. Moreover, given that EUK-134 mitigated unloading-related upregulation of oxidative stress, we propose nNOS μ translocation in response to mechanical

unloading in skeletal muscle is a redox-dependent phenomenon. nNOS μ translocation has previously been associated with increased FoxO3a activation and MuRF-1 activation and muscle atrophy during ground hindlimb unloading (42) and during 91 days of spaceflight (33). In contrast, a new publication indicates that nNOS is necessary in triggering an anabolic signaling pathway involving Nox4 and mTOR, which responds to overloading and regulates muscle hypertrophy (15). Furthermore, nitric oxide and NF- κ B were also found to be essential for stretch-induced proliferation of myoblasts (40). Indeed, inhibition of nitric oxide synthase has been shown to attenuate muscle remodeling and fiber-type shift in response to altered mechanical loading, whether the magnitude be increased or decreased (15, 38, 42). Therefore, redox signaling may couple

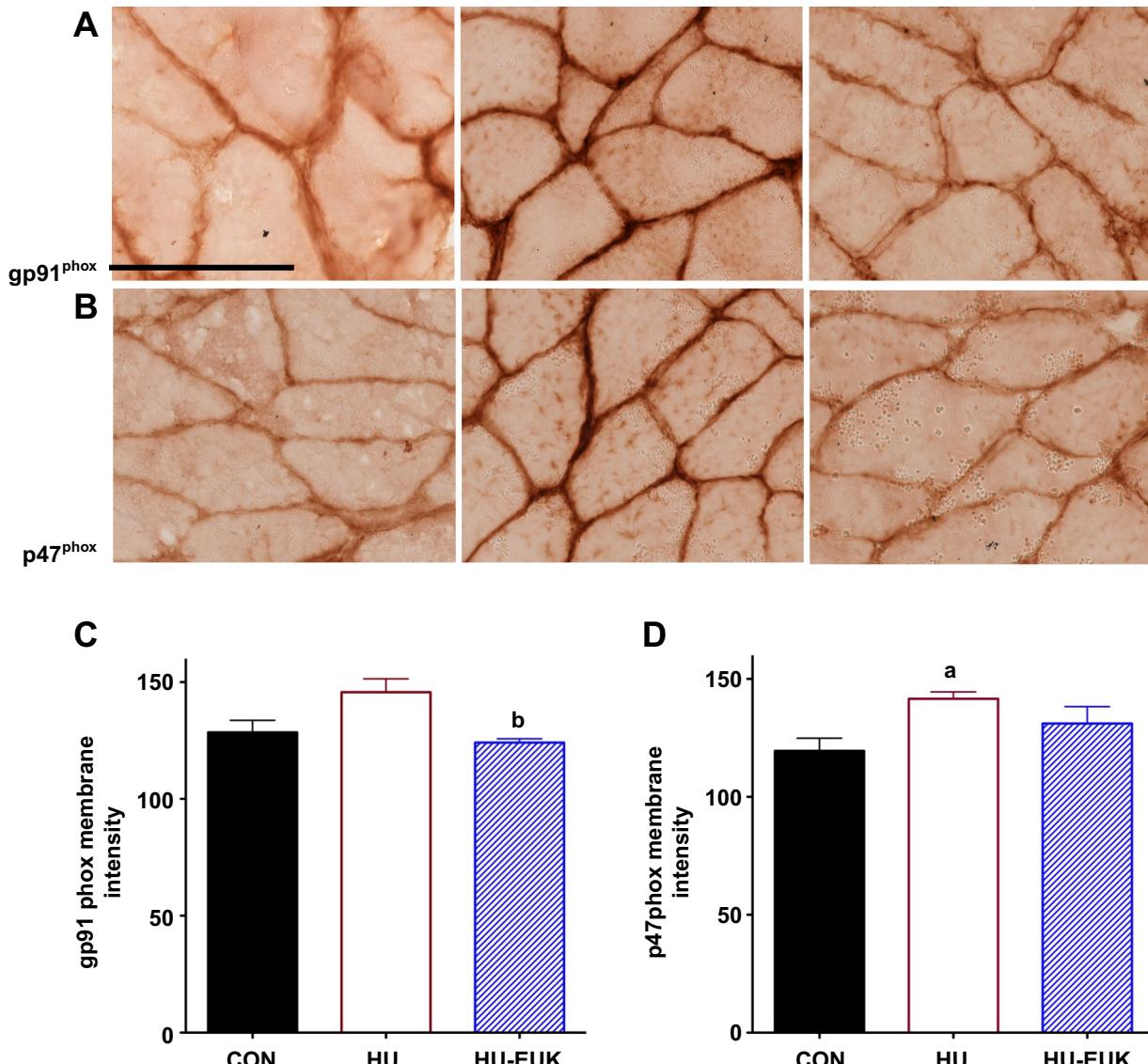


Fig. 7. Effect of EUK-134 on Nox2 subunit localization at the sarcolemma in for CON, HU, and HU-EUK groups ($n = 6$ /group). Localization of NAD(P)H oxidase subunits gp91^{phox} (Nox2) (A) and p47^{phox} (B) increased in response to mechanical unloading. The HU-EUK-134 displayed significantly lower ($P < 0.05$) levels of gp91^{phox} (A, C) than the HU group. p47^{phox}-positive immunostaining (B and D) was significantly greater ($P < 0.05$) for HU vs. CON, but not for rats treated with EUK-134. Scale bar = 100 μ m.

with nNOS to regulate remodeling of skeletal muscle morphology in response to dynamic changes in loading. In contrast, nitric oxide appears not to be requisite for basal proliferation of myoblasts and development (1, 39).

Intervention with EUK-134 not only mitigated unloading-induced atrophy, but also appeared to arrest a shift away from the slow-twitch phenotype in the rat soleus. This was an unexpected finding, but potentially significant, in understanding regulation of slow- and fast-twitch fibers in response to use and disuse. For example, Smith et al. (38) reported that inhibition of nitric oxide synthase abrogated a fiber-type shift from fast to slow in the plantaris with increased loading as a result of ablation of the soleus and gastrocnemius muscles, as they are all plantar flexor muscles. Therefore, we postulate that nNOS translocation may contribute to a shift in fiber type from slow to fast. We did note that EUK-134 protection against muscle fiber atrophy was confined to Type I fibers. These

findings could be consistent with preferential response of Type I fibers by mechanical unloading or an unloading-induced shift in muscle fiber type from slow-twitch fibers, which are paradoxically larger in the soleus (Fig. 1). It is possible that the time course of the study (54 h) may be too short to elicit morphological alterations in Type II fibers. In addition, the shift from slow- to fast-twitch fibers may seek to preserve cross-sectional area. Thus, additional study is warranted.

EUK-134 protection of FoxO3a phosphorylation. EUK-134 ameliorated the reduction in phosphorylation of FoxO3a at Thr-32 caused by hindlimb unloading (Fig. 5), thus reducing a trigger of proteolysis and fiber atrophy (9). Previously, Suzuki et al. (42) demonstrated that translocation of nNOS and cytosolic nitric oxide release by nNOS were required to dephosphorylate FoxO3a in the rat soleus during 14 days of mechanical unloading. FoxO3a activation stimulates upregulation of the ubiquitin ligases MuRF-1 and atrogin-1 (42). Therefore,

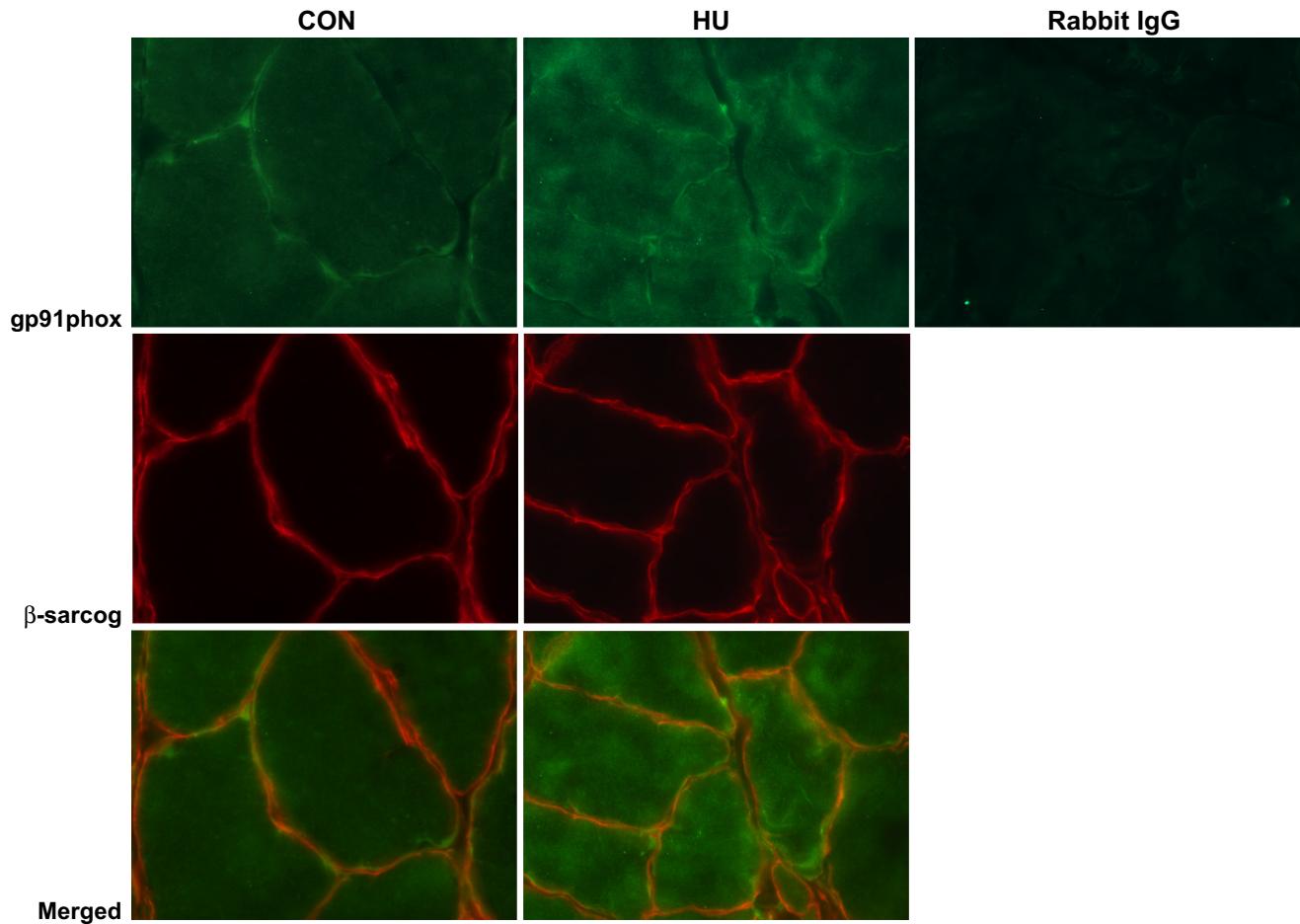


Fig. 8. Immunofluorescence of gp91phox in soleus muscles from control (CON) and hindlimb unloaded (HU) mice. β -sarcoglycan is used as a sarcolemmal marker. Merged images are also displayed. Rabbit IgG is used as a negative control for the primary antibody.

protection against FoxO3a dephosphorylation at Thr-32 by EUK-134 is consistent with a pathway, whereby oxidative stress participates in translocation of nNOS with mechanical unloading, directly leading to FoxO3a activation, as well as increased proteolysis through ubiquitin ligases (e.g., MuRF-1), leading to fiber atrophy (42).

Oxidative stress and mechanical unloading. Our results are consistent with the hypotheses that oxidative stress 1) is elevated early during mechanical unloading in skeletal muscle and 2) contributes significantly to the atrophy as a consequence of unloading. Protective effects of EUK-134 were specific to changes caused during the unloading protocol, as EUK-134 injections had no effect on normally loaded skeletal muscle mass or morphology (Fig. 2). Furthermore, elevated oxidative stress, as detected with 4-hydroxyneonal, was attenuated with EUK-134, which corresponded with sparing of muscle fiber atrophy.

Elevation of oxidative stress after 54 h of mechanical unloading of the rat soleus is consistent with previous studies that indicate oxidative stress is elevated during the first few days of mechanical unloading. Localization of 4-HNE adducts near the sarcolemma (Fig. 6) suggests a source nearby (e.g., Nox2, subsarcolemmal mitochondria). Previously, Abrogast et al. (4) reported that oxidative stress was elevated in unloaded skeletal muscle after 1–3 days. Bowman-Birk inhibitor complex (BBIC),

which exhibits antiproteasome and antioxidant properties, significantly reduced unloading-induced fiber atrophy and reduction of force-generating capacity (4). In addition, Powers and colleagues (48) also demonstrated that elevation in oxidative stress occurs in just 12–24 h of unloading in the diaphragm muscle during mechanical ventilation. In a series of studies, oxidative stress was found to be causal in stimulating apoptosis and muscle fiber atrophy in the unloaded diaphragm muscle (25, 26, 34, 48). Recently, transfection of catalase in the soleus has been shown to reduce soleus muscle atrophy during leg immobilization, while lowering oxidative stress (9).

It should be noted that evidence supportive of a contribution of oxidative stress in eliciting muscle atrophy with unloading has not been consistent across studies. Indeed, while some antioxidant strategies have been successful in reducing skeletal muscle atrophy with mechanical unloading (4, 25, 30), other interventions have not (6, 20). For example, administration of BBIC significantly attenuated muscle atrophy in limb muscles (4), while Trolox was less successful (6). Vitamin E reduced unloading-induced atrophy when preinjected for 21 days (3), while a broad, nonspecific cocktail of antioxidants was ineffective (20).

However, more recent, mechanistic studies indicate that ROS play an important role in unloading-induced atrophy. For example, catalase transfection suppressed activation of

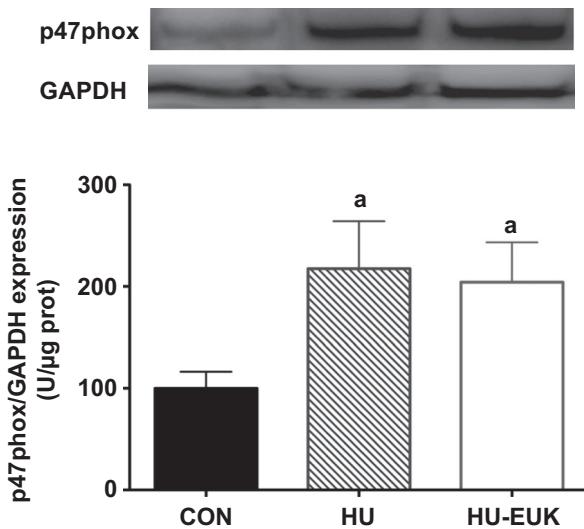


Fig. 9. Protein abundance for p47phox, expressed as a ratio relative to GAPDH (top) and β -actin expression (bottom). Representative blots and summary statistics are presented in HU and HU + EUK-134 groups exhibited significantly greater p47phox than controls (CON).

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FoxO3a and NF- κ B, while attenuating one-half of the soleus mass loss during 7 days of unloading (9). Inhibition of mitochondrial oxidative stress using SS-31 also significantly ameliorated muscle atrophy in a casting model of mechanical unloading (27). The reasons underlying differential responses to antioxidant strategies have not been resolved, but length of time, localized effects, specificity, and catalytic vs. scavenger approaches could all play a role. It is also possible that antioxidant therapeutics may be more effective in the early phase of mechanical unloading (1–7 days) than in later phases (14+ days).

While redox regulation of subsarcolemmal proteins, including nNOS in skeletal muscle remains poorly understood, there is growing acceptance that ROS sources participate in subcellular regulation of microenvironments, including Nox2 signaling in caveolae and lipid rafts (18, 44). We found increased Nox2 subunit localization at the membrane (Fig. 4) and elevated protein abundance of p47phox (Fig. 9); however, we cannot conclude a causal relationship. In addition, it is possible that Nox2 could serve to amplify or augment mitochondrial ROS (27), recently established as a causal factor in disuse-related atrophy.

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Study limitations. Recent studies have suggested that the cytokine TNF- α may play a significant role in apoptosis and ubiquitin ligase activation in skeletal muscle during mechanical unloading (2). It is possible that EUK-134 might play a protective role in unloading-induced atrophy by suppressing TNF- α levels, possibly via NF- κ B. However, we found no change in NF- κ B activity with unloading or EUK-134 at this stage of mechanical unloading.

It is possible that under high levels (mM) of H_2O_2 EUK-134 may oxidize to oxo-EUK and quench nitric oxide ($\cdot NO$) (37). While H_2O_2 levels in skeletal muscle are believed to exist in the high-nM to low- μ M range, future experiments could specifically inhibit iNOS as a source of RNS. In addition, high levels of EUK-134 at the tissue level may also remove or prevent the production of peroxynitrite ($OONO^-$). Thus, it is

possible that nNOS may contribute to its own dislocation. It is also possible that mechanical unloading affects nNOS via transcriptional regulation rather than posttranslational mechanisms or interaction with other proteins (e.g., caveolin-3). These provocative questions should be foci of future investigations.

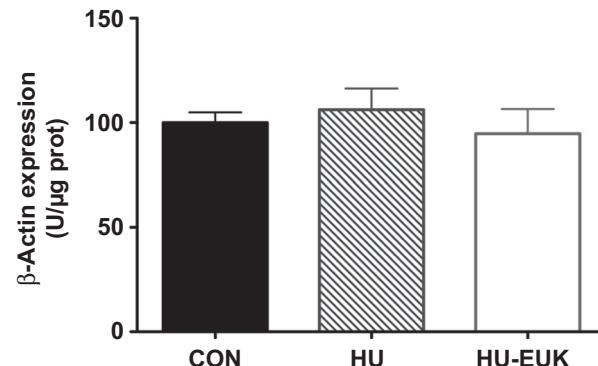
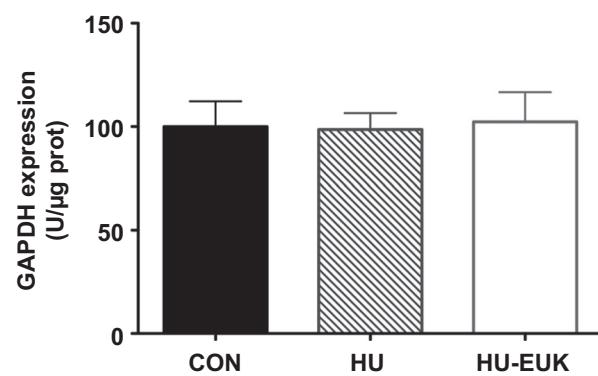
In summary, translocation of nNOS μ during short-term mechanical unloading in the rat soleus was reduced by the SOD/catalase mimetic EUK-134 and linked to protection against muscle fiber atrophy and fiber-type shift from slow to fast. In addition, prevention of nNOS translocation by EUK-134 was directly associated with protection against dephosphorylation of FoxO3a. Finally, EUK-134 reduced oxidative stress and normalized Nox2 levels. Our results suggest that loss of nNOS μ from the sarcolemma is an early, redox-dependent event in the response to mechanical unloading, linked to morphological changes in skeletal muscle phenotype.

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COLOR

Fig. 10. No significant differences in GAPDH nor β -actin among controls (CON), hindlimb unloaded (HU), and hindlimb unloaded + EUK-134 in the rat soleus muscle.

DISCLOSURES

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No conflicts of interest, financial or otherwise, are declared by the authors.

AUTHOR CONTRIBUTIONS

Author contributions: J.M.L. conception and design of research; J.M.L., M.K., J.M.H., Y.L., K.J., R.E.B., A.R., and D.A.M. performed experiments; J.M.L., M.K., J.M.H., Y.L., K.J., R.E.B., A.R., and D.A.M. analyzed data; J.M.L. and D.A.M. interpreted results of experiments; J.M.L. and Y.L. prepared figures; J.M.L. drafted manuscript; J.M.L., J.M.H., Y.L., R.E.B., and D.A.M. edited and revised manuscript; J.M.L. approved final version of manuscript.

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