



Medicine & Science in Sports & Exercise

Issue: Volume 41(5) Supplement 1, May 2009, p 131

Copyright: ©2009The American College of Sports Medicine

Publication Type: [F-11 Free Communication/Slide - Aging Effects on Muscle Structure and Function: MAY 29, 2009 1:00 PM - 3:00 PM ROOM: 4C4]

DOI: 10.1249/01.mss.0000353691.33400.08

ISSN: 0195-9131

Accession: 00005768-200905001-01157

[Hide Cover](#)

[F-11 Free Communication/Slide - Aging Effects on Muscle Structure and Function: MAY 29, 2009 1:00 PM - 3:00 PM ROOM: 4C4]

Myosin Immunohistochemistry And Fiber Adaptation Following Stretch-shortening Contraction Loading: Influence Of Glutathione And Age: 957: May 29 2:45 PM - 3:00 PM

Baker, Brent A.¹; Hollander, Melinda S.²; Kashon, Michael L.¹; Cutlip, Robert G.¹

Author Information

¹NIOSH, Morgantown, WV. ²West Virginia University, Morgantown, WV.

(Sponsor: Stephen E. Alway, Ph.D., FACSM, FACSM)

Email: bwb3@cdc.gov

(No relationships reported)

The extent and contribution of myosin heavy chain plasticity in the adaptive response following glutathione modulation and repetitive mechanical loading is not known.

PURPOSE: The purpose of this research was to identify the influence dietary supplementation with a glutathione antagonist (L-Buthionine Sulfoximine (BSO)) has on the adaptability of skeletal muscle during chronic high-intensity mechanical loading via stretch-shortening contractions (SSCs) in young and old rats.

METHODS: Left dorsiflexor muscles of young (12 weeks, N= 22) and old (30 months, N = 22), vehicle- (VEH) and BSO-treated rats were exposed 3 times per week for 4.5-weeks to a protocol of 80 maximal SSCs per exposure *in vivo*, while cage, age-matched rats served as controls (CON). Myofiber plasticity was characterized by myosin heavy chain (MHC_{slow}⁺, MHC_{fast}⁺, or MHC_{dev}⁺) response by immunohistochemical analysis, as well as by changes in muscle cross-sectional area distribution following the exposure period.

RESULTS: MHC_{slow}⁺ content was increased in the old rats, irrespective of treatment or exposure. Conversely, MHC_{fast}⁺ content was increased in the young rats, regardless of treatment of exposure. MHC_{dev}⁺ content was not changed following chronic SSC loading. Shifts to larger fiber cross-sectional areas were apparent for all SSC-exposed groups.

CONCLUSIONS: MHC_{dev}⁺ is not expressed following chronic SSC exposure, regardless of age, treatment, or exposure. Aging skeletal muscle does retain some capacity for adaptation; however aging apparently diminishes the adaptive response following chronic SSC exposure. Evidently, a population of small myofibers does exist that do not undergo regeneration in aged skeletal muscle, perhaps accounting for the disparity observed between morphological adaptation and functional performance.

Copyright (c) 2000-2014 Ovid Technologies, Inc.

[Terms of Use](#) | [Support & Training](#) | [About Us](#) | [Contact Us](#)

Version: OvidSP_UI03.12.00.116, SourceID 60384