Accepted Manuscript

Lipolysis and lipophagy in lipid storage myopathies

Corrado Angelini, Anna Chiara Nascimbeni, Giovanna Cenacchi, Elisabetta Tasca

PII: S0925-4439(16)30079-5

DOI: doi: 10.1016/j.bbadis.2016.04.008

Reference: BBADIS 64452

To appear in: BBA - Molecular Basis of Disease

Received date: 5 January 2016 Revised date: 31 March 2016 Accepted date: 11 April 2016



Please cite this article as: Corrado Angelini, Anna Chiara Nascimbeni, Giovanna Cenacchi, Elisabetta Tasca, Lipolysis and lipophagy in lipid storage myopathies, *BBA - Molecular Basis of Disease* (2016), doi: 10.1016/j.bbadis.2016.04.008

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

LIPOLYSIS AND LIPOPHAGY IN LIPID STORAGE MYOPATHIES

Corrado Angelini ¹, Anna Chiara Nascimbeni ², Giovanna Cenacchi ³, Elisabetta Tasca ¹

¹Fondazione San Camillo Hospital IRCCS, via Alberoni 70, 30126 Lido Venice, Italy; ²Department of Neurosciences, University of Padova, via Giustiniani 5, 35128 Padova, Italy; ³Department of Radiological and Histopathological Sciences, Alma Mater University of Bologna, via Massarenti 9, 40138 Bologna, Italy

Corresponding Author: Corrado Angelini, MD. Neuromuscular Department, Fondazione San Camillo Hospital IRCCS, via Alberoni 70, 30126 Lido Venice, Italy. Tel.: +39.49.8216155. Fax: +39.49.8216163. Email: corrado.angelini@unipd.it

ABBREVIATIONS

ATGL: adipose triglyceride lipase; ATP-ase: adenosine tri-phosphatase; CD: carnitine deficiency; CK: creatine kinase; COX: cytochrome oxidase; CPT: carnitine-palmitoyl-transferase; DAPI: 4',6-diamidin-2-phenylindole; EMG: electromyography; ETF: electron transfer flavoprotein; FOXO: fork head box protein; GAPDH: glyceraldehyde 3-phosphate dehydrogenase; HSL: hormone-sensitive lipase; LC3: microtubule-associated proteins light chain-3; LSM: lipid storage myopathy; MADD: multiple acyl-CoA dehydrogenase deficiency; MCT: medium chain triglyceride; NADH-TR: nicotinamide adenine dinucleotide dehydrogenase tetrazolium reductase; NLSD-M: neutral lipid storage disease with myopathy; OCTN2: carnitine organic cation transporter-2; p62-SQSTM1: p62-sequestosome-1; PAS: perjodic acid Schiff; PPARα: peroxisome proliferator-activated receptor-γ-coactivator-1α; SDH: succinate dehydrogenase; TFEB: transcription factor-EB.

Download English Version:

https://daneshyari.com/en/article/8259242

Download Persian Version:

https://daneshyari.com/article/8259242

<u>Daneshyari.com</u>