## Maria Lina Massimino

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Alteration in Calcium Handling at the Subcellular Level inmdx Myotubes. Journal of Biological Chemistry, 2001, 276, 4647-4651.	1.6	136
2	Involvement of caveolae and caveolae-like domains in signalling, cell survival and angiogenesis. Cellular Signalling, 2002, 14, 93-98.	1.7	72
3	The Metabolism and Imaging in Live Cells of the Bovine Prion Protein in Its Native Form or Carrying Single Amino Acid Substitutions. Molecular and Cellular Neurosciences, 2001, 17, 521-538.	1.0	62
4	Cellular Prion Protein Promotes Regeneration of Adult Muscle Tissue. Molecular and Cellular Biology, 2010, 30, 4864-4876.	1.1	58
5	Cellular prion protein is implicated in the regulation of local Ca <sup>2+</sup> movements in cerebellar granule neurons. Journal of Neurochemistry, 2011, 116, 881-890.	2.1	41
6	Cell surface nucleolin interacts with and internalizes Bothrops asper Lys49 phospholipase A2 and mediates its toxic activity. Scientific Reports, 2018, 8, 10619.	1.6	36
7	The cellular prion protein counteracts cardiac oxidative stress. Cardiovascular Research, 2014, 104, 93-102.	1.8	29
8	Absolute quantification of myosin heavy chain isoforms by selected reaction monitoring can underscore skeletal muscle changes in a mouse model of amyotrophic lateral sclerosis. Analytical and Bioanalytical Chemistry, 2017, 409, 2143-2153.	1.9	26
9	SOD1 in ALS: Taking Stock in Pathogenic Mechanisms and the Role of Glial and Muscle Cells. Antioxidants, 2022, 11, 614.	2.2	26
10	The prion protein constitutively controls neuronal store-operated Ca2+ entry through Fyn kinase. Frontiers in Cellular Neuroscience, 2015, 9, 416.	1.8	24
11	Human Doppel and prion protein share common membrane microdomains and internalization pathways. International Journal of Biochemistry and Cell Biology, 2004, 36, 2016-2031.	1.2	22
12	Heterogeneous PrPCmetabolism in skeletal muscle cells. FEBS Letters, 2006, 580, 878-884.	1.3	15
13	The Link of the Prion Protein with Ca2+ Metabolism and ROS Production, and the Possible Implication in Al² Toxicity. International Journal of Molecular Sciences, 2019, 20, 4640.	1.8	12
14	Nucleolin Rescues TDP-43 Toxicity in Yeast and Human Cell Models. Frontiers in Cellular Neuroscience, 2021, 15, 625665.	1.8	12
15	Altered behavioral aspects of aged mice lacking the cellular prion protein. Physiology and Behavior, 2013, 119, 86-91.	1.0	11
16	The prion protein regulates glutamate-mediated Ca2+ entry and mitochondrial Ca2+ accumulation in neurons. Journal of Cell Science, 2017, 130, 2736-2746.	1.2	11
17	Microglia in Prion Diseases: Angels or Demons?. International Journal of Molecular Sciences, 2020, 21, 7765.	1.8	11
18	Age-dependent neuromuscular impairment in prion protein knockout mice. Muscle and Nerve, 2016, 53, 269-279	1.0	10

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19	ALS-Associated SOD1(G93A) Decreases SERCA Pump Levels and Increases Store-Operated Ca2+ Entry in Primary Spinal Cord Astrocytes from a Transgenic Mouse Model. International Journal of Molecular Sciences, 2019, 20, 5151.	1.8	10
20	Generation and validation of novel adeno-associated viral vectors for the analysis of Ca2+ homeostasis in motor neurons. Scientific Reports, 2017, 7, 6521.	1.6	9
21	The Prion Protein Regulates Synaptic Transmission by Controlling the Expression of Proteins Key to Synaptic Vesicle Recycling and Exocytosis. Molecular Neurobiology, 2019, 56, 3420-3436.	1.9	9
22	Perturbations of the Proteome and of Secreted Metabolites in Primary Astrocytes from the hSOD1(G93A) ALS Mouse Model. International Journal of Molecular Sciences, 2021, 22, 7028.	1.8	9
23	Regulation of Endoplasmic Reticulum–Mitochondria Tethering and Ca2+ Fluxes by TDP-43 via GSK3β. International Journal of Molecular Sciences, 2021, 22, 11853.	1.8	9
24	The Cellular Prion Protein Is Expressed in Olfactory Sensory Neurons of Adult Mice but Does Not Affect the Early Events of the Olfactory Transduction Pathway. Chemical Senses, 2011, 36, 791-797.	1.1	7
25	Prion and TNFα: TAC(E)it agreement between the prion protein and cell signaling. Cell Cycle, 2010, 9, 4616-4621.	1.3	6