Paolo Broda

List of Publications by Year in descending order

14 papers	1,286 citations	933264 10 h-index	1058333 14 g-index
14 all docs	14 docs citations	14 times ranked	1745 citing authors

#	Article	IF	Citations
1	Mutations in the caveolin-3 gene cause autosomal dominant limb-girdle muscular dystrophy. Nature Genetics, 1998, 18, 365-368.	9.4	555
2	Hyperactivation of oxidative mitochondrial metabolism in epithelial cancer cells in situ. Cell Cycle, 2011, 10, 4047-4064.	1.3	256
3	Impairment of Caveolae Formation and T-System Disorganization in Human Muscular Dystrophy with Caveolin-3 Deficiency. American Journal of Pathology, 2002, 160, 265-270.	1.9	117
4	Increased Number of Caveolae and Caveolin-3 Overexpression in Duchenne Muscular Dystrophy. Biochemical and Biophysical Research Communications, 1999, 261, 547-550.	1.0	93
5	POMT2 gene mutation in limb-girdle muscular dystrophy with inflammatory changes. Biochemical and Biophysical Research Communications, 2007, 363, 1033-1037.	1.0	91
6	POMGnT1 Mutations in Congenital Muscular Dystrophy. Archives of Neurology, 2006, 63, 1491.	4.9	38
7	Genetic and Early Clinical Manifestations of Females Heterozygous for Duchenne/Becker Muscular Dystrophy. Pediatric Neurology, 2016, 55, 58-63.	1.0	37
8	Phenotypic Heterogeneity in Two Unrelated Danon Patients Associated with the SameLAMPâ€2Gene Mutation. Neuropediatrics, 2005, 36, 309-313.	0.3	32
9	Aquaporin-4 expression is severely reduced in human sarcoglycanopathies and dysferlinopathies. Cell Cycle, 2008, 7, 2199-2207.	1.3	20
10	Novel mutation in sarcotubular myopathy. Acta Myologica, 2019, 38, 8-12.	1.5	13
11	Expanding the histopathological spectrum of CFL2â€related myopathies. Clinical Genetics, 2018, 93, 1234-1239.	1.0	11
12	Spinal motor neuron involvement in a patient with homozygous PRUNE mutation. European Journal of Paediatric Neurology, 2018, 22, 541-543.	0.7	10
13	Clinical and molecular consequences of exon 78 deletion in DMD gene. Journal of Human Genetics, 2018, 63, 761-764.	1.1	7
14	Distal motor neuropathy associated with novel EMILIN1 mutation. Neurobiology of Disease, 2020, 137, 104757.	2.1	6