Lucia Artuso

List of Publications by Year
in descending order

9 papers	285 citations	1162889 8 h-index	1474057 9 g-index
9 all docs	9 docs citations	9 times ranked	474 citing authors

#	Article	IF	Citations
1	Impact of mutational status on outcomes in myelofibrosis patients treated with ruxolitinib in the COMFORT-II study. Blood, 2014, 123, 2157-2160.	0.6	115
2	Mitochondrial genome aberrations in skeletal muscle of patients with motor neuron disease. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2013, 14, 261-266.	1.1	14
3	Deep sequencing unearths Nuclear mitochondrial Sequences under Leber's hereditary optic neuropathy-associated false heteroplasmic mitochondrial DNA variants. Human Molecular Genetics, 2012, 21, 3753-3764.	1.4	15
4	Mitochondrial DNA metabolism in early development of zebrafish (Danio rerio). Biochimica Et Biophysica Acta - Bioenergetics, 2012, 1817, 1002-1011.	0.5	78
5	Mitochondrial genome large rearrangements in the skeletal muscle of a patient with <scp>PMA</scp>. European Journal of Neurology, 2012, 19, e63-4.	1.7	9
6	Bilateral progressive visual loss in an epileptic, mentally retarded boy. Middle East African Journal of Ophthalmology, 2011, 18, 67.	0.5	5
7	Late-onset Leber hereditary optic neuropathy mimicking Susacâ€™s syndrome. Journal of Neurology, 2010, 257, 1999-2003.	1.8	14
8	A new locus on 3p23â€“p25 for an autosomal-dominant limb-girdle muscular dystrophy, LGMD1H. European Journal of Human Genetics, 2010, 18, 636-641.	1.4	27
9	Molecular analysis in a family presenting with a mild form of late-onset autosomal dominant chronic progressive external ophthalmoplegia. Neuromuscular Disorders, 2009, 19, 423-426.	0.3	8