

# Lucia Artuso

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3860316/publications.pdf>

Version: 2024-02-01

9  
papers

285  
citations

1162367  
8  
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1473754  
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g-index

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9  
docs citations

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