

Manuela Spagnolo

List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	Homozygous mutations in <i>C1QBP</i> as cause of progressive external ophthalmoplegia (PEO) and mitochondrial myopathy with multiple mtDNA deletions. <i>Human Mutation</i> , 2020, 41, 1745-1750.	2.5	15
2	Biallelic pathogenic variants in <i>NDUFC2</i> cause early-onset Leigh syndrome and stalled biogenesis of complex I. <i>EMBO Molecular Medicine</i> , 2020, 12, e12619.	6.9	17
3	Paraoxonase, arylesterase and lactonase activities of paraoxonase-1 (PON1) in obese and severely obese women. <i>Scandinavian Journal of Clinical and Laboratory Investigation</i> , 2018, 78, 18-24.	1.2	25