Paulien Smits

List of Publications by Year
in descending order

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6 papers	659 citations	1478505 6 h-index	1872680 6 g-index
7 all docs	7 docs citations	7 times ranked	1163 citing authors

#	Article	IF	Citations
1	Mutation in subdomain G' of mitochondrial elongation factor G1 is associated with combined OXPHOS deficiency in fibroblasts but not in muscle. European Journal of Human Genetics, 2011, 19, 275-279.	2.8	42
2	Mutation in mitochondrial ribosomal protein MRPS22 leads to Cornelia de Lange-like phenotype, brain abnormalities and hypertrophic cardiomyopathy. European Journal of Human Genetics, 2011, 19, 394-399.	2.8	90
3	Functional consequences of mitochondrial tRNATrp and tRNAArg mutations causing combined OXPHOS defects. European Journal of Human Genetics, 2010, 18, 324-329.	2.8	32
4	Mitochondrial Translation and Beyond: Processes Implicated in Combined Oxidative Phosphorylation Deficiencies. Journal of Biomedicine and Biotechnology, 2010, 2010, 1-24.	3.0	158
5	Reconstructing the evolution of the mitochondrial ribosomal proteome. Nucleic Acids Research, 2007, 35, 4686-4703.	14.5	168
6	Distinct Clinical Phenotypes Associated with a Mutation in the Mitochondrial Translation Elongation Factor EFTs. American Journal of Human Genetics, 2006, 79, 869-877.	6.2	169