Takuya Fushimi

List of Publications by Year in descending order

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1684188 1720034 7 396 5 7 citations g-index h-index papers 7 7 7 860 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Neonatal-onset mitochondrial disease: clinical features, molecular diagnosis and prognosis. Archives of Disease in Childhood: Fetal and Neonatal Edition, 2022, 107, 329-334.	2.8	9
2	Development of Leigh syndrome with a high probability of cardiac manifestations in infantile-onset patients with m.14453GÂ>ÂA. Mitochondrion, 2022, 63, 1-8.	3.4	2
3	Genome sequencing and RNAâ€seq analyses of mitochondrial complex I deficiency revealed <i>Alu</i> insertionâ€mediated deletion in <i>NDUFV2</i> . Human Mutation, 2021, 42, 1422-1428.	2.5	4
4	Mortality of Japanese patients with Leigh syndrome: Effects of age at onset and genetic diagnosis. Journal of Inherited Metabolic Disease, 2020, 43, 819-826.	3.6	32
5	Cardiomyopathy in children with mitochondrial disease: Prognosis and genetic background. International Journal of Cardiology, 2019, 279, 115-121.	1.7	35
6	Clinical validity of biochemical and molecular analysis in diagnosing Leigh syndrome: a study of 106 Japanese patients. Journal of Inherited Metabolic Disease, 2017, 40, 685-693.	3.6	78
7	A Comprehensive Genomic Analysis Reveals the Genetic Landscape of Mitochondrial Respiratory Chain Complex Deficiencies. PLoS Genetics, 2016, 12, e1005679.	3 . 5	236