

Takuya Fushimi

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9621736/publications.pdf>

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7
papers

396
citations

1684188
5
h-index

1720034
7
g-index

7
all docs

7
docs citations

7
times ranked

860
citing authors

#	ARTICLE	IF	CITATIONS
1	A Comprehensive Genomic Analysis Reveals the Genetic Landscape of Mitochondrial Respiratory Chain Complex Deficiencies. PLoS Genetics, 2016, 12, e1005679.	3.5	236
2	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
3	Cardiomyopathy in children with mitochondrial disease: Prognosis and genetic background. International Journal of Cardiology, 2019, 279, 115-121.	1.7	35
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	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
6	Genome sequencing and RNA-seq analyses of mitochondrial complex I deficiency revealed <i>Alu</i> insertion-mediated deletion in <i>NDUFV2</i> . Human Mutation, 2021, 42, 1422-1428.	2.5	4
7	Development of Leigh syndrome with a high probability of cardiac manifestations in infantile-onset patients with m.14453G>A. Mitochondrion, 2022, 63, 1-8.	3.4	2