

Laurence Walsh

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/2551486/publications.pdf>

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6
papers

258
citations

1478505

6
h-index

1872680

6
g-index

6
all docs

6
docs citations

6
times ranked

893
citing authors

#	ARTICLE	IF	CITATIONS
1	Enhanced MAPK1 Function Causes a Neurodevelopmental Disorder within the RASopathy Clinical Spectrum. <i>American Journal of Human Genetics</i> , 2020, 107, 499-513.	6.2	48
2	Variants in TCF20 in neurodevelopmental disability: description of 27 new patients and review of literature. <i>Genetics in Medicine</i> , 2019, 21, 2036-2042.	2.4	23
3	De Novo and Inherited Loss-of-Function Variants in TLK2: Clinical and Genotype-Phenotype Evaluation of a Distinct Neurodevelopmental Disorder. <i>American Journal of Human Genetics</i> , 2018, 102, 1195-1203.	6.2	37
4	Heterozygous De Novo UBTF Gain-of-Function Variant Is Associated with Neurodegeneration in Childhood. <i>American Journal of Human Genetics</i> , 2017, 101, 267-273.	6.2	41
5	WDR26 Haploinsufficiency Causes a Recognizable Syndrome of Intellectual Disability, Seizures, Abnormal Gait, and Distinctive Facial Features. <i>American Journal of Human Genetics</i> , 2017, 101, 139-148.	6.2	45
6	The expanding clinical phenotype of Bosch-Boonstra-Schaaf optic atrophy syndrome: 20 new cases and possible genotype-phenotype correlations. <i>Genetics in Medicine</i> , 2016, 18, 1143-1150.	2.4	64