

Jennifer A Lee

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/8565275/publications.pdf>

Version: 2024-02-01

10
papers

1,274
citations

1163117

8
h-index

1474206

9
g-index

10
all docs

10
docs citations

10
times ranked

2312
citing authors

#	ARTICLE	IF	CITATIONS
1	A DNA Replication Mechanism for Generating Nonrecurrent Rearrangements Associated with Genomic Disorders. <i>Cell</i> , 2007, 131, 1235-1247.	28.9	756
2	Genomic Rearrangements and Gene Copy-Number Alterations as a Cause of Nervous System Disorders. <i>Neuron</i> , 2006, 52, 103-121.	8.1	284
3	Spastic paraplegia type 2 associated with axonal neuropathy and apparent <i>PLP1</i> position effect. <i>Annals of Neurology</i> , 2006, 59, 398-403.	5.3	83
4	Role of genomic architecture in PLP1 duplication causing Pelizaeus-Merzbacher disease. <i>Human Molecular Genetics</i> , 2006, 15, 2250-2265.	2.9	73
5	Phenotype and mutation expansion of the PTPN23 associated disorder characterized by neurodevelopmental delay and structural brain abnormalities. <i>European Journal of Human Genetics</i> , 2020, 28, 76-87.	2.8	21
6	Prenatal diagnosis of PLP1 copy number by array comparative genomic hybridization. <i>Prenatal Diagnosis</i> , 2005, 25, 1188-1191.	2.3	17
7	Three additional patients with EED-associated overgrowth: potential mutation hotspots identified?. <i>Journal of Human Genetics</i> , 2019, 64, 561-572.	2.3	16
8	Xq22 deletions and correlation with distinct neurological disease traits in females: Further evidence for a contiguous gene syndrome. <i>Human Mutation</i> , 2020, 41, 150-168.	2.5	15
9	Steric Clash in the SET Domain of Histone Methyltransferase NSD1 as a Cause of Sotos Syndrome and Its Genetic Heterogeneity in a Brazilian Cohort. <i>Genes</i> , 2016, 7, 96.	2.4	9
10	Inside Back Cover, Volume 41, Issue 1. <i>Human Mutation</i> , 2020, 41, ii.	2.5	0