

Jehan Suleiman

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

22
papers

579
citations

11
h-index

24
g-index

26
ext. papers

695
ext. citations

3.4
avg, IF

3.81
L-index

#	Paper	IF	Citations
22	VPS26C homozygous nonsense variant in two cousins with neurodevelopmental deficits, growth failure, skeletal abnormalities, and distinctive facial features. <i>Clinical Genetics</i> , 2020 , 97, 644-648	4	1
21	Homozygous loss-of-function variants of TASP1, a gene encoding an activator of the histone methyltransferases KMT2A and KMT2D, cause a syndrome of developmental delay, happy demeanor, distinctive facial features, and congenital anomalies. <i>Human Mutation</i> , 2019 , 40, 1985-1992	4.7	6
20	ATP13A2-related juvenile-onset Parkinson disease. <i>Brain and Development</i> , 2019 , 41, 223	2.2	1
19	Pediatric Multiple Sclerosis in the United Arab Emirates: Characteristics From a Multicenter Study and Global Comparison. <i>Journal of Child Neurology</i> , 2018 , 33, 422-427	2.5	6
18	TASP1 is deleted in an infant with developmental delay, microcephaly, distinctive facial features, and multiple congenital anomalies. <i>Clinical Genetics</i> , 2018 , 94, 170-173	4	7
17	Compound heterozygous variants in the multiple PDZ domain protein (MPDZ) cause a case of mild non-progressive communicating hydrocephalus. <i>BMC Medical Genetics</i> , 2018 , 19, 34	2.1	6
16	WDR45B-related intellectual disability, spastic quadriplegia, epilepsy, and cerebral hypoplasia: A consistent neurodevelopmental syndrome. <i>Clinical Genetics</i> , 2018 , 93, 360-364	4	23
15	ATP13A2 novel mutations causing a rare form of juvenile-onset Parkinson disease. <i>Brain and Development</i> , 2018 , 40, 824-826	2.2	10
14	West syndrome, developmental and epileptic encephalopathy, and severe CNS disorder associated with WWOX mutations. <i>Epileptic Disorders</i> , 2018 , 20, 401-412	1.9	11
13	Mitochondrial dynamics: Biological roles, molecular machinery, and related diseases. <i>Molecular Genetics and Metabolism</i> , 2018 , 125, 315-321	3.7	62
12	PPP1R21 homozygous null variants associated with developmental delay, muscle weakness, distinctive facial features, and brain abnormalities. <i>Clinical Genetics</i> , 2018 , 94, 351-355	4	2
11	B3GALNT2-Related Dystroglycanopathy: Expansion of the Phenotype with Novel Mutation Associated with Muscle-Eye-Brain Disease, Walker-Warburg Syndrome, Epileptic Encephalopathy-West Syndrome, and Sensorineural Hearing Loss. <i>Neuropediatrics</i> , 2018 , 49, 289-295	1.6	5
10	17q23.2q23.3 de novo duplication in association with speech and language disorder, learning difficulties, incoordination, motor skill impairment, and behavioral disturbances: a case report. <i>BMC Medical Genetics</i> , 2017 , 18, 119	2.1	7
9	Expanding the genetic heterogeneity of intellectual disability. <i>Human Genetics</i> , 2017 , 136, 1419-1429	6.3	74
8	Clinical and molecular delineation of dysequilibrium syndrome type 2 and profound sensorineural hearing loss in an inbred Arab family. <i>American Journal of Medical Genetics, Part A</i> , 2016 , 170A, 540-543	2.5	15
7	A novel de novo mutation in DYNC1H1 gene underlying malformation of cortical development and cataract. <i>Meta Gene</i> , 2016 , 9, 124-7	0.7	17
6	The recognition and treatment of autoimmune epilepsy in children. <i>Developmental Medicine and Child Neurology</i> , 2015 , 57, 431-40	3.3	57

5	Autoimmune epilepsy in children: case series and proposed guidelines for identification. <i>Epilepsia</i> , 2013 , 54, 1036-45	6.4	58
4	Autoantibodies to neuronal antigens in children with new-onset seizures classified according to the revised ILAE organization of seizures and epilepsies. <i>Epilepsia</i> , 2013 , 54, 2091-100	6.4	45
3	Immune-mediated steroid-responsive epileptic spasms and epileptic encephalopathy associated with VGKC-complex antibodies. <i>Developmental Medicine and Child Neurology</i> , 2011 , 53, 1058-60	3.3	36
2	VGKC antibodies in pediatric encephalitis presenting with status epilepticus. <i>Neurology</i> , 2011 , 76, 1252-56.5	6.5	90
1	Severe encephalopathy with swine origin influenza A H1N1 infection in childhood: case reports. <i>Neurology</i> , 2010 , 74, 1077-8	6.5	37