Jehan Suleiman

List of Publications by Citations

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	VGKC antibodies in pediatric encephalitis presenting with status epilepticus. *Neurology*, **2011**, 76, 1252-5	6.5	90
21	Expanding the genetic heterogeneity of intellectual disability. *Human Genetics*, **2017**, 136, 1419-1429	6.3	74
20	Mitochondrial dynamics: Biological roles, molecular machinery, and related diseases. *Molecular Genetics and Metabolism*, **2018**, 125, 315-321	3.7	62
19	Autoimmune epilepsy in children: case series and proposed guidelines for identification. *Epilepsia*, **2013**, 54, 1036-45	6.4	58
18	The recognition and treatment of autoimmune epilepsy in children. *Developmental Medicine and Child Neurology*, **2015**, 57, 431-40	3.3	57
17	Autoantibodies to neuronal antigens in children with new-onset seizures classified according to the revised ILAE organization of seizures and epilepsies. *Epilepsia*, **2013**, 54, 2091-100	6.4	45
16	Severe encephalopathy with swine origin influenza A H1N1 infection in childhood: case reports. *Neurology*, **2010**, 74, 1077-8	6.5	37
15	Immune-mediated steroid-responsive epileptic spasms and epileptic encephalopathy associated with VGKC-complex antibodies. *Developmental Medicine and Child Neurology*, **2011**, 53, 1058-60	3.3	36
14	WDR45B-related intellectual disability, spastic quadriplegia, epilepsy, and cerebral hypoplasia: A consistent neurodevelopmental syndrome. *Clinical Genetics*, **2018**, 93, 360-364	4	23
13	A novel de novo mutation in DYNC1H1 gene underlying malformation of cortical development and cataract. *Meta Gene*, **2016**, 9, 124-7	0.7	17
12	Clinical and molecular delineation of dysequilibrium syndrome type 2 and profound sensorineural hearing loss in an inbred Arab family. *American Journal of Medical Genetics, Part A*, **2016**, 170A, 540-543	2.5	15
11	West syndrome, developmental and epileptic encephalopathy, and severe CNS disorder associated with WWOX mutations. *Epileptic Disorders*, **2018**, 20, 401-412	1.9	11
10	ATP13A2 novel mutations causing a rare form of juvenile-onset Parkinson disease. *Brain and Development*, **2018**, 40, 824-826	2.2	10
9	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. *BMC Medical Genetics*, **2017**, 18, 119	2.1	7
8	TASP1 is deleted in an infant with developmental delay, microcephaly, distinctive facial features, and multiple congenital anomalies. *Clinical Genetics*, **2018**, 94, 170-173	4	7
7	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. *Human Mutation*, **2019**, 40, 1985-1992	4.7	6
6	Pediatric Multiple Sclerosis in the United Arab Emirates: Characteristics From a Multicenter Study and Global Comparison. *Journal of Child Neurology*, **2018**, 33, 422-427	2.5	6

5	Compound heterozygous variants in the multiple PDZ domain protein (MPDZ) cause a case of mild non-progressive communicating hydrocephalus. *BMC Medical Genetics*, **2018**, 19, 34	2.1	6
4	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. *Neuropediatrics*, **2018**, 49, 289-295	1.6	5
3	PPP1R21 homozygous null variants associated with developmental delay, muscle weakness, distinctive facial features, and brain abnormalities. *Clinical Genetics*, **2018**, 94, 351-355	4	2
2	ATP13A2-related juvenile-onset Parkinson disease. *Brain and Development*, **2019**, 41, 223	2.2	1
1	VPS26C homozygous nonsense variant in two cousins with neurodevelopmental deficits, growth failure, skeletal abnormalities, and distinctive facial features. *Clinical Genetics*, **2020**, 97, 644-648	4	1