

Deb K Pal

List of Publications by Citations

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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.

59
papers

1,334
citations

22
h-index

35
g-index

65
ext. papers

1,673
ext. citations

5.1
avg, IF

4.29
L-index

#	Paper	IF	Citations
59	BRD2 (RING3) is a probable major susceptibility gene for common juvenile myoclonic epilepsy. <i>American Journal of Human Genetics</i> , 2003 , 73, 261-70	11	146
58	Attention impairment in rolandic epilepsy: systematic review. <i>Epilepsia</i> , 2008 , 49, 1570-80	6.4	110
57	High risk of reading disability and speech sound disorder in rolandic epilepsy families: case-control study. <i>Epilepsia</i> , 2007 , 48, 2258-65	6.4	85
56	Gene Panel Testing in Epileptic Encephalopathies and Familial Epilepsies. <i>Molecular Syndromology</i> , 2016 , 7, 210-219	1.5	65
55	A neurocognitive endophenotype associated with rolandic epilepsy. <i>Epilepsia</i> , 2012 , 53, 705-11	6.4	56
54	Defining the phenotypic spectrum of SLC6A1 mutations. <i>Epilepsia</i> , 2018 , 59, 389-402	6.4	54
53	Genetic evaluation and counseling for epilepsy. <i>Nature Reviews Neurology</i> , 2010 , 6, 445-53	15	54
52	Autosomal dominant inheritance of centrotemporal sharp waves in rolandic epilepsy families. <i>Epilepsia</i> , 2007 , 48, 2266-72	6.4	51
51	The role of mTOR signalling in neurogenesis, insights from tuberous sclerosis complex. <i>Seminars in Cell and Developmental Biology</i> , 2016 , 52, 12-20	7.5	46
50	Idiopathic focal epilepsies: the "lost tribe". <i>Epileptic Disorders</i> , 2016 , 18, 252-88	1.9	42
49	The spectrum of intermediate SCN8A-related epilepsy. <i>Epilepsia</i> , 2019 , 60, 830-844	6.4	38
48	A meta-analysis of literacy and language in children with rolandic epilepsy. <i>Developmental Medicine and Child Neurology</i> , 2015 , 57, 1019-26	3.3	34
47	Is Rolandic epilepsy associated with abnormal findings on cranial MRI?. <i>Epilepsy Research</i> , 2007 , 75, 180-5		34
46	Complex inheritance and parent-of-origin effect in juvenile myoclonic epilepsy. <i>Brain and Development</i> , 2006 , 28, 92-8	2.2	33
45	Incorporating epilepsy genetics into clinical practice: a 360° evaluation. <i>Npj Genomic Medicine</i> , 2018 , 3, 13	6.2	32
44	Microdeletions of ELP4 Are Associated with Language Impairment, Autism Spectrum Disorder, and Mental Retardation. <i>Human Mutation</i> , 2015 , 36, 842-50	4.7	31
43	Epilepsy and neurodevelopmental disorders of language. <i>Current Opinion in Neurology</i> , 2011 , 24, 126-31	7.1	27

42	Clinical spectrum of -related epileptic disorders. <i>Neurology</i> , 2019 , 92, e1238-e1249	6.5	25
41	A microRNA-328 binding site in PAX6 is associated with centrotemporal spikes of rolandic epilepsy. <i>Annals of Clinical and Translational Neurology</i> , 2016 , 3, 512-22	5.3	23
40	Antiepileptic drug treatment of rolandic epilepsy and Panayiotopoulos syndrome: clinical practice survey and clinical trial feasibility. <i>Archives of Disease in Childhood</i> , 2015 , 100, 62-7	2.2	22
39	Dissecting the genetic basis of myoclonic-astatic epilepsy. <i>Epilepsia</i> , 2012 , 53, 1303-13	6.4	22
38	Analysis of rare copy number variation in absence epilepsies. <i>Neurology: Genetics</i> , 2016 , 2, e56	3.8	22
37	Association analysis using next-generation sequence data from publicly available control groups: the robust variance score statistic. <i>Bioinformatics</i> , 2014 , 30, 2179-88	7.2	21
36	PIGT-CDG, a disorder of the glycosylphosphatidylinositol anchor: description of 13 novel patients and expansion of the clinical characteristics. <i>Genetics in Medicine</i> , 2019 , 21, 2216-2223	8.1	18
35	Phenotypic and genetic spectrum of epilepsy with myoclonic atonic seizures. <i>Epilepsia</i> , 2020 , 61, 995-1007	6.4	18
34	Evaluating genetic heterogeneity in complex disorders. <i>Human Heredity</i> , 2002 , 53, 216-26	1.1	18
33	Psychosocial complications in juvenile myoclonic epilepsy. <i>Epilepsy and Behavior</i> , 2019 , 90, 122-128	3.2	16
32	Parental mosaicism in epilepsies due to alleged de novo variants. <i>Epilepsia</i> , 2019 , 60, e63-e66	6.4	16
31	Core Health Outcomes in Childhood Epilepsy (CHOICE): Development of a core outcome set using systematic review methods and a Delphi survey consensus. <i>Epilepsia</i> , 2019 , 60, 857-871	6.4	16
30	Phenobarbital for childhood epilepsy: systematic review. <i>Paediatric and Perinatal Drug Therapy</i> , 2006 , 7, 31-42		16
29	Risk factors for reading disability in families with rolandic epilepsy. <i>Epilepsy and Behavior</i> , 2015 , 53, 174-9	3.2	15
28	Identification of new risk factors for rolandic epilepsy: CNV at Xp22.31 and alterations at cholinergic synapses. <i>Journal of Medical Genetics</i> , 2018 , 55, 607-616	5.8	15
27	The genetics of reading disability in an often excluded sample: novel loci suggested for reading disability in Rolandic epilepsy. <i>PLoS ONE</i> , 2012 , 7, e40696	3.7	14
26	Decreased functional connectivity within a language subnetwork in benign epilepsy with centrotemporal spikes. <i>Epilepsia Open</i> , 2017 , 2, 214-225	4	11
25	Phenotypic features of familial febrile seizures: case-control study. <i>Neurology</i> , 2003 , 60, 410-4	6.5	10

24	Epilepsy-specific patient-reported outcome measures of children's health-related quality of life: A systematic review of measurement properties. <i>Epilepsia</i> , 2020 , 61, 230-248	6.4	9
23	Core Health Outcomes In Childhood Epilepsy (CHOICE): protocol for the selection of a core outcome set. <i>Trials</i> , 2017 , 18, 572	2.8	9
22	Trait impulsivity in Juvenile Myoclonic Epilepsy. <i>Annals of Clinical and Translational Neurology</i> , 2021 , 8, 138-152	5.3	9
21	Meta-analysis of response inhibition in juvenile myoclonic epilepsy. <i>Epilepsy and Behavior</i> , 2020 , 106, 107038	3.2	5
20	The genetics of common epilepsies: common or distinct?. <i>Lancet Neurology</i> , 2014 , 13, 859-60	24.1	5
19	SCN8A heterozygous variants are associated with anoxic-epileptic seizures. <i>American Journal of Medical Genetics, Part A</i> , 2020 , 182, 1209-1216	2.5	4
18	Practical aspects of childhood epilepsy. <i>BMJ</i> , 2019 , 367, l6096	5.9	4
17	Novel mechanisms, treatments, and outcome measures in childhood sleep. <i>Frontiers in Psychology</i> , 2015 , 6, 602	3.4	4
16	Remote and Long-Term Self-Monitoring of Electroencephalographic and Noninvasive Measurable Variables at Home in Patients With Epilepsy (EEG@HOME): Protocol for an Observational Study. <i>JMIR Research Protocols</i> , 2021 , 10, e25309	2	4
15	A specific deficit of auditory processing in children with Rolandic Epilepsy and their relatives. <i>Epilepsy and Behavior</i> , 2017 , 72, 135-139	3.2	3
14	Mapping epilepsy-specific patient-reported outcome measures for children to a proposed core outcome set for childhood epilepsy. <i>Epilepsy and Behavior</i> , 2020 , 112, 107372	3.2	3
13	Trait impulsivity correlates with active myoclonic seizures in genetic generalized epilepsy. <i>Epilepsy and Behavior</i> , 2020 , 112, 107260	3.2	3
12	Sex-specific disease modifiers in juvenile myoclonic epilepsy.. <i>Scientific Reports</i> , 2022 , 12, 2785	4.9	3
11	The Effects of Sleep on Emotional Target Detection Performance: A Novel iPad-Based Pediatric Game. <i>Frontiers in Psychology</i> , 2018 , 9, 241	3.4	2
10	Complementing the phenotypical spectrum of TUBA1A tubulinopathy and its role in early-onset epilepsies.. <i>European Journal of Human Genetics</i> , 2022 ,	5.3	2
9	Reply to: Is a microRNA-328 binding site in PAX6 associated with Rolandic epilepsy?. <i>Annals of Clinical and Translational Neurology</i> , 2017 , 4, 278-280	5.3	1
8	ELP4 in rolandic epilepsy and BRD2 in juvenile myoclonic epilepsy. <i>Epilepsia</i> , 2010 , 51, 73-73	6.4	1
7	Neurodevelopmental origins of self-limiting rolandic epilepsy: Systematic review of MR imaging studies. <i>Epilepsia Open</i> , 2021 , 6, 310-322	4	1

6	A Qualitative Investigation Into What Parents Want From an Online Behavioural Sleep Intervention for Children With Epilepsy. <i>Frontiers in Psychology</i> , 2021 , 12, 628605	3.4	1
5	Narcolepsy Following Yellow Fever Vaccination: A Case Report. <i>Frontiers in Neurology</i> , 2016 , 7, 130	4.1	1
4	Development and Evaluation of the CASTLE Trial Online Sleep Intervention for Parents of Children with Epilepsy. <i>Frontiers in Psychology</i> , 2021 , 12, 679804	3.4	1
3	Response to pyridoxine in CACNA1A epilepsy-ataxia does not imply a causal effect. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2021 , 91, 196-197	3.2	1
2	ZMYND11 variants are a novel cause of centrotemporal and generalised epilepsies with neurodevelopmental disorder. <i>Clinical Genetics</i> , 2021 , 100, 412-429	4	0
1	Fashions come and go. <i>Epilepsia</i> , 2011 , 52, 191-2; discussion 193-6	6.4	