

# Petra M C Callenbach

## List of Publications by Year in descending order

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Version: 2024-02-01

34  
papers

1,585  
citations

394286

19  
h-index

377752

34  
g-index

35  
all docs

35  
docs citations

35  
times ranked

2324  
citing authors

#	ARTICLE	IF	CITATIONS
1	Mutations in DEPDC5 cause familial focal epilepsy with variable foci. <i>Nature Genetics</i> , 2013, 45, 546-551.	9.4	301
2	Familial cortical myoclonic tremor with epilepsy: A single syndromic classification for a group of pedigrees bearing common features. <i>Movement Disorders</i> , 2005, 20, 665-673.	2.2	161
3	<i>GRIN2A</i>-related disorders: genotype and functional consequence predict phenotype. <i>Brain</i> , 2019, 142, 80-92.	3.7	143
4	Mortality Risks in New-Onset Childhood Epilepsy. <i>Pediatrics</i> , 2013, 132, 124-131.	1.0	141
5	Familial Occurrence of Epilepsy in Children with Newly Diagnosed Multiple Seizures: Dutch Study of Epilepsy in Childhood. <i>Epilepsia</i> , 1998, 39, 331-336.	2.6	84
6	Long term outcome of benign childhood epilepsy with centrotemporal spikes: Dutch Study of Epilepsy in Childhood. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2010, 19, 501-506.	0.9	72
7	Long-term outcome of childhood absence epilepsy: Dutch Study of Epilepsy in Childhood. <i>Epilepsy Research</i> , 2009, 83, 249-256.	0.8	71
8	Haploinsufficiency of the STX1B gene is associated with myoclonic astatic epilepsy. <i>European Journal of Paediatric Neurology</i> , 2016, 20, 489-492.	0.7	52
9	A Dutch family with 'familial cortical tremor with epilepsy'. <i>Journal of Neurology</i> , 2002, 249, 829-834.	1.8	46
10	Linkage and mutational analysis of CLCN2 in childhood absence epilepsy. <i>Epilepsy Research</i> , 2007, 75, 145-153.	0.8	46
11	Familial Partial Epilepsy with Variable Foci in a Dutch Family: Clinical Characteristics and Confirmation of Linkage to Chromosome 22q. <i>Epilepsia</i> , 2003, 44, 1298-1305.	2.6	43
12	Add-on levetiracetam in children and adolescents with refractory epilepsy: Results of an open-label multi-centre study. <i>European Journal of Paediatric Neurology</i> , 2008, 12, 321-327.	0.7	43
13	Levetiracetam Monotherapy in Children with Epilepsy: A Systematic Review. <i>CNS Drugs</i> , 2015, 29, 371-382.	2.7	41
14	Linkage and association analysis of CACNG3 in childhood absence epilepsy. <i>European Journal of Human Genetics</i> , 2007, 15, 463-472.	1.4	39
15	Benign familial infantile convulsions: a clinical study of seven Dutch families. <i>European Journal of Paediatric Neurology</i> , 2002, 6, 269-283.	0.7	34
16	Evaluation of CACNA1H in European patients with childhood absence epilepsy. <i>Epilepsy Research</i> , 2006, 69, 177-181.	0.8	32
17	Prevalence of sleep disturbances in people with epilepsy and the impact on quality of life: a survey in secondary care. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2019, 69, 298-303.	0.9	31
18	Genome wide high density SNP-based linkage analysis of childhood absence epilepsy identifies a susceptibility locus on chromosome 3p23-p14. <i>Epilepsy Research</i> , 2009, 87, 247-255.	0.8	29

#	ARTICLE	IF	CITATIONS
19	<i>PRRT2</i> mutation causes benign familial infantile convulsions. <i>Neurology</i> , 2012, 79, 2154-2155.	1.5	22
20	Î-Catenin ( <i>CTNND2</i> ) missense mutation in familial cortical myoclonic tremor and epilepsy. <i>Neurology</i> , 2017, 89, 2341-2350.	1.5	22
21	PRRT2-related phenotypes in patients with a 16p11.2 deletion. <i>European Journal of Medical Genetics</i> , 2019, 62, 265-269.	0.7	20
22	Sumatriptan nasal spray in the acute treatment of migraine in adolescents and children. <i>European Journal of Paediatric Neurology</i> , 2007, 11, 325-330.	0.7	18
23	Clinical and genetic aspects of idiopathic epilepsies in childhood. <i>European Journal of Paediatric Neurology</i> , 2005, 9, 91-103.	0.7	14
24	RCTs with new antiepileptic drugs in children: A systematic review of monotherapy studies and their methodology. <i>Epilepsy Research</i> , 2010, 91, 1-9.	0.8	13
25	Copy number variation in a hospital-based cohort of children with epilepsy. <i>Epilepsia Open</i> , 2017, 2, 244-254.	1.3	13
26	Ketogenic Diet in Refractory Childhood Epilepsy. <i>Child Neurology Open</i> , 2018, 5, 2329048X1877949.	0.5	10
27	Contextual Structured Reporting in Radiology: Implementation and Long-Term Evaluation in Improving the Communication of Critical Findings. <i>Journal of Medical Systems</i> , 2020, 44, 148.	2.2	10
28	Antiepileptic drug prescription in Dutch children from 2006â€“2014 using pharmacy-dispensing data. <i>Epilepsy Research</i> , 2018, 146, 21-27.	0.8	9
29	Hereditary epilepsy syndromes. <i>Clinical Neurology and Neurosurgery</i> , 1997, 99, 159-171.	0.6	7
30	CMAP variation over a length of nerve in diabetic neuropathy. <i>Muscle and Nerve</i> , 1995, 18, 907-909.	1.0	4
31	The diagnostic value of duplex ultrasound in detecting the presence and location of a stenosis in an autologous arteriovenous fistula. <i>Journal of Vascular Access</i> , 2020, 21, 217-222.	0.5	4
32	Changes in empowerment and anxiety of patients and parents during genetic counselling for epilepsy. <i>European Journal of Paediatric Neurology</i> , 2021, 32, 128-135.	0.7	4
33	Treatment of prolonged convulsive seizures in children; a single centre, retrospective, observational study. <i>European Journal of Paediatric Neurology</i> , 2014, 18, 663-669.	0.7	3
34	Investigator-initiated randomized controlled trials in children with epilepsy: Mission impossible?. <i>Epilepsia Open</i> , 2017, 2, 32-38.	1.3	3