Petra M C Callenbach

List of Publications by Year
in descending order

34	1,585	394286 19	377752 34
papers	citations	h-index	g-index

35	35	35	2324
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Changes in empowerment and anxiety of patients and parents during genetic counselling for epilepsy. European Journal of Paediatric Neurology, 2021, 32, 128-135.	0.7	4
2	The diagnostic value of duplex ultrasound in detecting the presence and location of a stenosis in an autologous arteriovenous fistula. Journal of Vascular Access, 2020, 21, 217-222.	0.5	4
3	Contextual Structured Reporting in Radiology: Implementation and Long-Term Evaluation in Improving the Communication of Critical Findings. Journal of Medical Systems, 2020, 44, 148.	2.2	10
4	PRRT2-related phenotypes in patients with a 16p11.2 deletion. European Journal of Medical Genetics, 2019, 62, 265-269.	0.7	20
5	Prevalence of sleep disturbances in people with epilepsy and the impact on quality of life: a survey in secondary care. Seizure: the Journal of the British Epilepsy Association, 2019, 69, 298-303.	0.9	31
6	GRIN2A-related disorders: genotype and functional consequence predict phenotype. Brain, 2019, 142, 80-92.	3.7	143
7	Antiepileptic drug prescription in Dutch children from 2006â€“2014 using pharmacy-dispensing data. Epilepsy Research, 2018, 146, 21-27.	0.8	9
8	Ketogenic Diet in Refractory Childhood Epilepsy. Child Neurology Open, 2018, 5, 2329048X1877949.	0.5	10
9	Investigatorâ€initiated randomized controlled trials in children with epilepsy: Mission impossible?. Epilepsia Open, 2017, 2, 32-38.	1.3	3
10	Copy number variation in a hospitalâ€based cohort of children with epilepsy. Epilepsia Open, 2017, 2, 244-254.	1.3	13
11	Î´-Catenin (CTNND2) missense mutation in familial cortical myoclonic tremor and epilepsy. Neurology, 2017, 89, 2341-2350.	1.5	22
12	Haploinsufficiency of the STX1B gene is associated with myoclonic astatic epilepsy. European Journal of Paediatric Neurology, 2016, 20, 489-492.	0.7	52
13	Levetiracetam Monotherapy in Children with Epilepsy: A Systematic Review. CNS Drugs, 2015, 29, 371-382.	2.7	41
14	Treatment of prolonged convulsive seizures in children; a single centre, retrospective, observational study. European Journal of Paediatric Neurology, 2014, 18, 663-669.	0.7	3
15	Mortality Risks in New-Onset Childhood Epilepsy. Pediatrics, 2013, 132, 124-131.	1.0	141
16	Mutations in DEPDC5 cause familial focal epilepsy with variable foci. Nature Genetics, 2013, 45, 546-551.	9.4	301
17	PRRT2 mutation causes benign familial infantile convulsions. Neurology, 2012, 79, 2154-2155.	1.5	22
18	RCTs with new antiepileptic drugs in children: A systematic review of monotherapy studies and their methodology. Epilepsy Research, 2010, 91, 1-9.	0.8	13

#	Article	IF	Citations
19	Long term outcome of benign childhood epilepsy with centrotemporal spikes: Dutch Study of Epilepsy in Childhood. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 501-506.	0.9	72
20	Long-term outcome of childhood absence epilepsy: Dutch Study of Epilepsy in Childhood. Epilepsy Research, 2009, 83, 249-256.	0.8	71
21	Genome wide high density SNP-based linkage analysis of childhood absence epilepsy identifies a susceptibility locus on chromosome 3p23-p14. Epilepsy Research, 2009, 87, 247-255.	0.8	29
22	Add-on levetiracetam in children and adolescents with refractory epilepsy: Results of an open-label multi-centre study. European Journal of Paediatric Neurology, 2008, 12, 321-327.	0.7	43
23	Linkage and mutational analysis of CLCN2 in childhood absence epilepsy. Epilepsy Research, 2007, 75, 145-153.	0.8	46
24	Linkage and association analysis of CACNG3 in childhood absence epilepsy. European Journal of Human Genetics, 2007, 15, 463-472.	1.4	39
25	Sumatriptan nasal spray in the acute treatment of migraine in adolescents and children. European Journal of Paediatric Neurology, 2007, 11, 325-330.	0.7	18
26	Evaluation of CACNA1H in European patients with childhood absence epilepsy. Epilepsy Research, 2006, 69, 177-181.	0.8	32
27	Clinical and genetic aspects of idiopathic epilepsies in childhood. European Journal of Paediatric Neurology, 2005, 9, 91-103.	0.7	14
28	Familial cortical myoclonic tremor with epilepsy: A single syndromic classification for a group of pedigrees bearing common features. Movement Disorders, 2005, 20, 665-673.	2.2	161
29	Familial Partial Epilepsy with Variable Foci in a Dutch Family: Clinical Characteristics and Confirmation of Linkage to Chromosome 22q. Epilepsia, 2003, 44, 1298-1305.	2.6	43
30	Benign familial infantile convulsions: a clinical study of seven Dutch families. European Journal of Paediatric Neurology, 2002, 6, 269-283.	0.7	34
31	A Dutch family with 'familial cortical tremor with epilepsy'. Journal of Neurology, 2002, 249, 829-834.	1.8	46
32	Familial Occurrence of Epilepsy in Children with Newly Diagnosed Multiple Seizures: Dutch Study of Epilepsy in Childhood. Epilepsia, 1998, 39, 331-336.	2.6	84
33	Hereditary epilepsy syndromes. Clinical Neurology and Neurosurgery, 1997, 99, 159-171.	0.6	7
34	CMAP variation over a length of nerve in diabetic neuropathy. Muscle and Nerve, 1995, 18, 907-909.	1.0	4