Lieven Lagae

List of Publications by Year
in descending order

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149 papers	10,909 citations	47 h-index	99 g-index
		47006	33894
151 all docs	151 docs citations	151 times ranked	11798 citing authors

#	Article	IF	Citations
1	Operational classification of seizure types by the International League Against Epilepsy: Position Paper of the ILAE Commission for Classification and Terminology. Epilepsia, 2017, 58, 522-530.	5.1	2,191
2	De Novo Mutations in the Sodium-Channel Gene SCN1A Cause Severe Myoclonic Epilepsy of Infancy. American Journal of Human Genetics, 2001, 68, 1327-1332.	6.2	1,111
3	Instruction manual for the <scp>ILAE</scp> 2017 operational classification of seizure types. Epilepsia, 2017, 58, 531-542.	5.1	699
4	Gain-of-function mutations in IFIH1 cause a spectrum of human disease phenotypes associated with upregulated type I interferon signaling. Nature Genetics, 2014, 46, 503-509.	21.4	490
5	Genetic and phenotypic heterogeneity suggest therapeutic implications in SCN2A-related disorders. Brain, 2017, 140, 1316-1336.	7.6	426
6	Antenatal Maternal Anxiety is Related to HPA-Axis Dysregulation and Self-Reported Depressive Symptoms in Adolescence: A Prospective Study on the Fetal Origins of Depressed Mood. Neuropsychopharmacology, 2008, 33, 536-545.	5.4	387
7	Proposed consensus definitions for newâ€onset refractory status epilepticus (NORSE), febrile infectionâ€related epilepsy syndrome (FIRES), and related conditions. Epilepsia, 2018, 59, 739-744.	5.1	308
8	Fenfluramine hydrochloride for the treatment of seizures in Dravet syndrome: a randomised, double-blind, placebo-controlled trial. Lancet, The, 2019, 394, 2243-2254.	13.7	227
9	High antenatal maternal anxiety is related to impulsivity during performance on cognitive tasks in 14- and 15-year-olds. Neuroscience and Biobehavioral Reviews, 2005, 29, 259-269.	6.1	225
10	Successful use of fenfluramine as an addâ€on treatment for Dravet syndrome. Epilepsia, 2012, 53, 1131-1139.	5.1	175
11	Management of epilepsy associated with tuberous sclerosis complex: Updated clinical recommendations. European Journal of Paediatric Neurology, 2018, 22, 738-748.	1.6	151
12	Altered functional connectivity of the language network in ASD: Role of classical language areas and cerebellum. NeuroImage: Clinical, 2014, 4, 374-382.	2.7	139
13	Prevention of Epilepsy in Infants with Tuberous Sclerosis Complex in the <scp>EPISTOP</scp> Trial. Annals of Neurology, 2021, 89, 304-314.	5.3	137
14	Quality of life and comorbidities associated with Dravet syndrome severity: a multinational cohort survey. Developmental Medicine and Child Neurology, 2018, 60, 63-72.	2.1	125
15	Inborn errors in RNA polymerase III underlie severe varicella zoster virus infections. Journal of Clinical Investigation, 2017, 127, 3543-3556.	8.2	125
16	Long-term cognitive sequelae of antenatal maternal anxiety: involvement of the orbitofrontal cortex. Neuroscience and Biobehavioral Reviews, 2006, 30, 1078-1086.	6.1	124
17	Vagus nerve stimulation for refractory epilepsy: A Belgian multicenter study. European Journal of Paediatric Neurology, 2007, 11, 261-269.	1.6	118
18	ADHD Deficit as Measured in Adolescent Boys with a Continuous Performance Task Is Related to Antenatal Maternal Anxiety. Pediatric Research, 2006, 59, 78-82.	2.3	117

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19	Mutations in SNORD118 cause the cerebral microangiopathy leukoencephalopathy with calcifications and cysts. Nature Genetics, 2016, 48, 1185-1192.	21.4	114
20	Fiveâ€year extended followâ€up status of 10 patients with Dravet syndrome treated with fenfluramine. Epilepsia, 2016, 57, e129-34.	5.1	97
21	Cognitive side effects of anti-epileptic drugs. Seizure: the Journal of the British Epilepsy Association, 2006, 15, 235-241.	2.0	96
22	Pharmacological Analysis of the Anti-epileptic Mechanisms of Fenfluramine in scn1a Mutant Zebrafish. Frontiers in Pharmacology, 2017, 8, 191.	3.5	96
23	Intravenous Immunoglobulins in Refractory Childhood-Onset Epilepsy: Effects on Seizure Frequency, EEG Activity, and Cerebrospinal Fluid Cytokine Profile. Epilepsia, 2007, 48, 1739-1749.	5.1	93
24	Non-EEG seizure detection systems and potential SUDEP prevention: State of the art. Seizure: the Journal of the British Epilepsy Association, 2016, 41, 141-153.	2.0	91
25	Serotonergic Modulation as Effective Treatment for Dravet Syndrome in a Zebrafish Mutant Model. ACS Chemical Neuroscience, 2016, 7, 588-598.	3.5	86
26	Non-EEG seizure-detection systems and potential SUDEP prevention: State of the art. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 345-355.	2.0	85
27	Pharmacological Characterization of an Antisense Knockdown Zebrafish Model of Dravet Syndrome: Inhibition of Epileptic Seizures by the Serotonin Agonist Fenfluramine. PLoS ONE, 2015, 10, e0125898.	2.5	83
28	Aberrant Inclusion of a Poison Exon Causes Dravet Syndrome and Related SCN1A-Associated Genetic Epilepsies. American Journal of Human Genetics, 2018, 103, 1022-1029.	6.2	76
29	Treatment and long term outcome in West syndrome: The clinical reality. A multicentre follow up study. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 159-164.	2.0	75
30	Efficacy and safety of perampanel in adolescent patients with drug-resistant partial seizures inÂthree double-blind, placebo-controlled, phase III randomized clinical studies and a combined extension study. European Journal of Paediatric Neurology, 2015, 19, 435-445.	1.6	73
31	Low-dose fenfluramine in the treatment of neurologic disorders: experience in Dravet syndrome. Therapeutic Advances in Neurological Disorders, 2015, 8, 328-338.	3.5	67
32	Developmental brain alterations in 17 year old boys are related to antenatal maternal anxiety. Clinical Neurophysiology, 2009, 120, 1116-1122.	1.5	64
33	Learning Disabilities: Definitions, Epidemiology, Diagnosis, and Intervention Strategies. Pediatric Clinics of North America, 2008, 55, 1259-1268.	1.8	63
34	Automated Detection of Tonicâ€“Clonic Seizures Using 3-D Accelerometry and Surface Electromyography in Pediatric Patients. IEEE Journal of Biomedical and Health Informatics, 2016, 20, 1333-1341.	6.3	62
35	Diagnostic implications of genetic copy number variation in epilepsy plus. Epilepsia, 2019, 60, 689-706.	5.1	61
36	Current role of carbamazepine and oxcarbazepine in the management of epilepsy. Seizure: the Journal of the British Epilepsy Association, 2020, 83, 251-263.	2.0	59

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37	A pilot, openâ€label study of the effectiveness and tolerability of lowâ€dose <scp>ZX</scp>008 (fenfluramine <scp>HC</scp>l) in Lennoxâ€Gastaut syndrome. Epilepsia, 2018, 59, 1881-1888.	5.1	58
38	Ketogenic diet for the treatment of pediatric epilepsy: review and meta-analysis. Child's Nervous System, 2020, 36, 1099-1109.	1.1	58
39	Severe Myoclonic Epilepsy in Infancy: Toward an Optimal Treatment. Journal of Child Neurology, 2004, 19, 516-521.	1.4	57
40	Adjunctive perampanel in adolescents with inadequately controlled partialâ€onset seizures: A randomized study evaluating behavior, efficacy, and safety. Epilepsia, 2016, 57, 1120-1129.	5.1	57
41	Gain-of-function FHF1 mutation causes early-onset epileptic encephalopathy with cerebellar atrophy. Neurology, 2016, 86, 2162-2170.	1.1	57
42	Dravet syndrome: Treatment options and management of prolonged seizures. Epilepsia, 2019, 60, S39-S48.	5.1	56
43	Long-term home monitoring of hypermotor seizures by patient-worn accelerometers. Epilepsy and Behavior, 2013, 26, 118-125.	1.7	51
44	TSC2 pathogenic variants are predictive of severe clinical manifestations in TSC infants: results of the EPISTOP study. Genetics in Medicine, 2020, 22, 1489-1497.	2.4	51
45	Early onset epileptic encephalopathy or genetically determined encephalopathy with early onset epilepsy? Lessons learned from TSC. European Journal of Paediatric Neurology, 2016, 20, 203-211.	1.6	49
46	Fenfluramine HCl (Fintepla^{Â®}) provides longâ€term clinically meaningful reduction in seizure frequency: Analysis of an ongoing openâ€label extension study. Epilepsia, 2020, 61, 2396-2404.	5.1	49
47	Peri-ictal ECG changes in childhood epilepsy: Implications for detection systems. Epilepsy and Behavior, 2013, 29, 72-76.	1.7	48
48	Coding and small non-coding transcriptional landscape of tuberous sclerosis complex cortical tubers: implications for pathophysiology and treatment. Scientific Reports, 2017, 7, 8089.	3.3	47
49	Epilepsy and cannabidiol: a guide to treatment. , 2020, 22, 1-14.		46
50	ERP correlates of complex human decision making in a gambling paradigm: Detection and resolution of conflict. Psychophysiology, 2008, 45, 714-720.	2.4	45
51	Left fronto-parietal white matter correlates with individual differences in children's ability to solve additions and multiplications: A tractography study. NeuroImage, 2014, 90, 117-127.	4.2	44
52	Efficacy and Safety of Fenfluramine for the Treatment of Seizures Associated With Lennox-Gastaut Syndrome. JAMA Neurology, 2022, 79, 554.	9.0	43
53	Online Automated Seizure Detection in Temporal Lobe Epilepsy Patients Using Single-lead ECG. International Journal of Neural Systems, 2017, 27, 1750022.	5.2	42
54	Early Clinical Predictors of Autism Spectrum Disorder in Infants with Tuberous Sclerosis Complex: Results from the EPISTOP Study. Journal of Clinical Medicine, 2019, 8, 788.	2.4	42

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55	Impact of fenfluramine on the expected SUDEP mortality rates in patients with Dravet syndrome. Seizure: the Journal of the British Epilepsy Association, 2021, 93, 154-159.	2.0	41
56	Cardiovascular safety of low-dose fenfluramine in Dravet syndrome: a review of its benefit-risk profile in a new patient population. Current Medical Research and Opinion, 2017, 33, 1773-1781.	1.9	40
57	Why we urgently need improved seizure and epilepsy therapies for children and neonates. Neuropharmacology, 2020, 170, 107854.	4.1	40
58	The administration of rescue medication to children with prolonged acute convulsive seizures in the community: What happens in practice?. European Journal of Paediatric Neurology, 2013, 17, 14-23.	1.6	39
59	The challenges and innovations for therapy in children with epilepsy. Nature Reviews Neurology, 2014, 10, 249-260.	10.1	38
60	Long-term effects of adjunctive perampanel on cognition in adolescents with partial seizures. Epilepsy and Behavior, 2018, 83, 50-58.	1.7	38
61	Epilepsy and Neurodevelopmental Comorbidities in Tuberous Sclerosis Complex: A Natural History Study. Pediatric Neurology, 2020, 106, 10-16.	2.1	37
62	Caregiver impact and health service use in high and low severity Dravet syndrome: A multinational cohort study. Seizure: the Journal of the British Epilepsy Association, 2019, 65, 72-79.	2.0	35
63	Accurate detection of typical absence seizures in adults and children using a twoâ€channel electroencephalographic wearable behind the ears. Epilepsia, 2021, 62, 2741-2752.	5.1	34
64	Rational treatment options with AEDs and ketogenic diet in Landauâ€Kleffner syndrome: Still waiting after all these years. Epilepsia, 2009, 50, 59-62.	5.1	32
65	Guidance on Dravet syndrome from infant to adult care: Road map for treatment planning in Europe. Epilepsia Open, 2022, 7, 11-26.	2.4	32
66	Drug repurposing for Dravet syndrome in scn1Lab^{â^'/â^'} mutant zebrafish. Epilepsia, 2019, 60, e8-e13.	5.1	31
67	Child development at 6 years after maternal cancer diagnosis and treatment during pregnancy. European Journal of Cancer, 2020, 138, 57-67.	2.8	31
68	mTOR-related neuropathology in mutant tsc2 zebrafish: Phenotypic, transcriptomic and pharmacological analysis. Neurobiology of Disease, 2017, 108, 225-237.	4.4	29
69	Detecting rare events using extreme value statistics applied to epileptic convulsions in children. Artificial Intelligence in Medicine, 2014, 60, 89-96.	6.5	26
70	Vagus Nerve Stimulation in children: A focus on intellectual disability. European Journal of Paediatric Neurology, 2017, 21, 427-440.	1.6	26
71	Dravet syndrome. Current Opinion in Neurology, 2021, 34, 213-218.	3.6	26
72	Can ECG monitoring identify seizures?. Journal of Electrocardiology, 2015, 48, 1069-1074.	0.9	25

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73	Is autism driven by epilepsy in infants with Tuberous Sclerosis Complex?. Annals of Clinical and Translational Neurology, 2020, 7, 1371-1381.	3.7	23
74	Are we failing to provide adequate rescue medication to children at risk of prolonged convulsive seizures in schools?. Archives of Disease in Childhood, 2013, 98, 777-780.	1.9	21
75	The mis-wired language network in children with developmental language disorder: insights from DTI tractography. Brain Imaging and Behavior, 2019, 13, 973-984.	2.1	21
76	Myelin Pathology Beyond White Matter in Tuberous Sclerosis Complex (TSC) Cortical Tubers. Journal of Neuropathology and Experimental Neurology, 2020, 79, 1054-1064.	1.7	21
77	Cortical malformations: a frequent cause of epilepsy in children. European Journal of Pediatrics, 2000, 159, 555-562.	2.7	20
78	Clinical practice. European Journal of Pediatrics, 2011, 170, 413-418.	2.7	20
79	Adaptive nocturnal seizure detection using heart rate and low-complexity novelty detection. Seizure: the Journal of the British Epilepsy Association, 2018, 59, 48-53.	2.0	20
80	Interictal cardiorespiratory variability in temporal lobe and absence epilepsy in childhood. Physiological Measurement, 2015, 36, 845-856.	2.1	19
81	Prediction of Neurodevelopment in Infants With Tuberous Sclerosis Complex Using Early EEG Characteristics. Frontiers in Neurology, 2020, 11, 582891.	2.4	19
82	Recommendations for the treatment of epilepsy in adult and pediatric patients in Belgium: 2020 update. Acta Neurologica Belgica, 2021, 121, 241-257.	1.1	19
83	Early epileptiform EEG activity in infants with tuberous sclerosis complex predicts epilepsy and neurodevelopmental outcomes. Epilepsia, 2021, 62, 1208-1219.	5.1	19
84	Serotonin receptors in epilepsy: Novel treatment targets?. Epilepsia Open, 2022, 7, 231-246.	2.4	19
85	Long-term accelerometry-triggered video monitoring and detection of tonicâ€“clonic and clonic seizures in a home environment: Pilot study. Epilepsy & Behavior Case Reports, 2016, 5, 66-71.	1.5	18
86	Recommendations for the treatment of epilepsy in adult patients in general practice in Belgium: an update. Acta Neurologica Belgica, 2012, 112, 119-131.	1.1	16
87	Overlap between linear scleroderma, progressive facial hemiatrophy and immune-inflammatory encephalitis in a paediatric cohort. European Journal of Pediatrics, 2015, 174, 1247-1254.	2.7	16
88	Effect of rescue medication on seizure duration in non-institutionalized children with epilepsy. European Journal of Paediatric Neurology, 2018, 22, 56-63.	1.6	15
89	2017 International League Against Epilepsy classifications of seizures and epilepsy are steps in the right direction. Epilepsia, 2019, 60, 1040-1044.	5.1	15
90	Automated detection of absence seizures using a wearable electroencephalographic device: a phase 3 validation study and feasibility of automated behavioral testing. Epilepsia, 2023, 64, .	5.1	15

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91	Use of Zebrafish Models to Boost Research in Rare Genetic Diseases. International Journal of Molecular Sciences, 2021, 22, 13356.	4.1	15
92	Whatâ€™s new in: â€œGenetics in childhood epilepsyâ€• European Journal of Pediatrics, 2008, 167, 715-722.	2.7	14
93	Ictal and interictal respiratory changes in temporal lobe and absence epilepsy in childhood. Epilepsy Research, 2013, 106, 410-416.	1.6	14
94	Children with well controlled epilepsy possess different spatio-temporal patterns of causal network connectivity during a visual working memory task. Cognitive Neurodynamics, 2016, 10, 99-111.	4.0	14
95	Renal Replacement Therapy in children with severe developmental disability: guiding questions for decision-making. European Journal of Pediatrics, 2018, 177, 1735-1743.	2.7	14
96	Efficacy of Fenfluramine and Norfenfluramine Enantiomers and Various Antiepileptic Drugs in a Zebrafish Model of Dravet Syndrome. Neurochemical Research, 2021, 46, 2249-2261.	3.3	14
97	Drug Development for Rare Paediatric Epilepsies: Current State and Future Directions. Drugs, 2019, 79, 1917-1935.	10.9	13
98	The importance of assessing behaviour and cognition in antiepileptic drug trials in children and adolescents. Acta Neurologica Belgica, 2017, 117, 425-432.	1.1	12
99	Defining the phenotype of FHF1 developmental and epileptic encephalopathy. Epilepsia, 2020, 61, e71-e78.	5.1	11
100	The Ketogenic Diet Revisited: Beyond Ketones. Frontiers in Neurology, 2021, 12, 720073.	2.4	10
101	Genetic Modulation of Neurocognitive Development in Cancer Patients throughout the Lifespan: a Systematic Review. Neuropsychology Review, 2019, 29, 190-219.	4.9	9
102	Antiepileptic Drug Teratogenicity and De Novo Genetic Variation Load. Annals of Neurology, 2020, 87, 897-906.	5.3	9
103	First line management of prolonged convulsive seizures in children and adults: good practice points. Acta Neurologica Belgica, 2013, 113, 375-380.	1.1	8
104	The administration of rescue medication to children with prolonged acute convulsive seizures in a non-hospital setting: an exploratory survey of healthcare professionalsâ€™ perspectives. European Journal of Pediatrics, 2014, 173, 773-779.	2.7	8
105	Antenatal maternal anxiety modulates the BOLD response in 20-year-old men during endogenous cognitive control. Brain Imaging and Behavior, 2020, 14, 830-846.	2.1	8
106	Association of Early MRI Characteristics With Subsequent Epilepsy and Neurodevelopmental Outcomes in Children With Tuberous Sclerosis Complex. Neurology, 2022, 98, .	1.1	8
107	Treatment of a symptomatic posterior fossa subdural effusion in a child. Child's Nervous System, 1999, 15, 90-93.	1.1	7
108	Health-related quality of life and the burden of prolonged seizures in noninstitutionalized children with epilepsy. Epilepsy and Behavior, 2020, 102, 106340.	1.7	7

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109	Results of quantitative EEG analysis are associated with autism spectrum disorder and development abnormalities in infants with tuberous sclerosis complex. Biomedical Signal Processing and Control, 2021, 68, 102658.	5.7	7
110	The need for broad spectrum and safe anti-epileptic drugs in childhood epilepsy. Acta Neurologica Belgica, 2009, 109, 167-70.	1.1	7
111	Malignant migrating partial seizures in Aicardi syndrome. Developmental Medicine and Child Neurology, 2008, 50, 790-792.	2.1	6
112	The concept of disease modification. European Journal of Paediatric Neurology, 2020, 24, 43-46.	1.6	6
113	Electroencephalography source localization analysis in epileptic children during a visual workingâ€memory task. International Journal for Numerical Methods in Biomedical Engineering, 2020, 36, e3404.	2.1	6
114	Detection of epileptic seizures from single lead ECG by means of phase rectified signal averaging. , 2014, 2014, 3789-90.		5
115	Overview of clinical efficacy and risk data of benzodiazepines for prolonged seizures. Epileptic Disorders, 2014, 16, 44-49.	1.3	5
116	Comparison and combination of electrocardiogram, electromyogram and accelerometry for tonic-clonic seizure detection in children. , 2018, , .		5
117	MicroRNAâ€34a activation in tuberous sclerosis complex during early brain development may lead to impaired corticogenesis. Neuropathology and Applied Neurobiology, 2021, 47, 796-811.	3.2	5
118	An update on congenital melanocytic nevus syndrome: A case report and literature review. Journal of Cutaneous Pathology, 2021, 48, 1497-1503.	1.3	5
119	Semi-Supervised One-Class Transfer Learning for Heart Rate Based Epileptic Seizure Detection. , 0, , .		4
120	Classification as autonomic versus sensory seizures. Epilepsia, 2019, 60, 2003-2005.	5.1	4
121	Patient MW: transient visual hemi-agnosia. Journal of Neurology, 2019, 266, 691-698.	3.6	4
122	Data describing child development at 6 years after maternal cancer diagnosis and treatment during pregnancy. Data in Brief, 2020, 32, 106209.	1.0	4
123	Expanding the clinical spectrum of Fowler syndrome: Three siblings with survival into adulthood and systematic review of the literature. Clinical Genetics, 2020, 98, 423-432.	2.0	4
124	Down-regulation of the brain-specific cell-adhesion molecule contactin-3 in tuberous sclerosis complex during the early postnatal period. Journal of Neurodevelopmental Disorders, 2022, 14, 8.	3.1	4
125	Recommendations for the treatment of epilepsies in general practice in Belgium. Acta Neurologica Belgica, 2008, 108, 118-30.	1.1	4
126	Treatment of Focal-Onset Seizures in Children: Should This Be More Etiology-Driven?. Frontiers in Neurology, 2022, 13, 842276.	2.4	4

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127	Risperidone in the treatment of childhood autistic disorder: an open pilot study. Acta Neuropsychiatrica, 2002, 14, 242-249.	2.1	3
128	Transient hypothyroidism associated with viral Human Parechovirus encephalitis in a newborn. European Journal of Paediatric Neurology, 2015, 19, 706-710.	1.6	3
129	Long-term impact of prenatal exposure to chemotherapy on executive functioning: An ERP study. Clinical Neurophysiology, 2019, 130, 1655-1664.	1.5	3
130	The training and organization of Paediatric Neurology in Europe: Special report of the European Paediatric Neurology Society & Committee of National Advisors. European Journal of Paediatric Neurology, 2020, 28, 6-15.	1.6	3
131	A survey of the European Reference Network EpiCARE on clinical practice for selected rare epilepsies. Epilepsia Open, 2021, 6, 160-170.	2.4	3
132	Evolution of electroencephalogram in infants with tuberous sclerosis complex and neurodevelopmental outcome: a prospective cohort study. Developmental Medicine and Child Neurology, 2022, 64, 495-501.	2.1	3
133	Vaccination and childhood epilepsies. European Journal of Paediatric Neurology, 2022, 36, 57-68.	1.6	3
134	Is it safe for people with epilepsy to donate blood? A systematic review. Epilepsy Research, 2018, 139, 143-149.	1.6	2
135	Paediatric status epilepticus: finally, some evidence-based treatment guidance, but still a long way to go. The Lancet Child and Adolescent Health, 2020, 4, 351-352.	5.6	2
136	Cognitive outcome in drugâ€resistant childhood epilepsy. Developmental Medicine and Child Neurology, 2021, 63, 633-633.	2.1	2
137	PPFIA4 mutation: A second hit in POLG related disease?. Epilepsy and Behavior Reports, 2021, 16, 100455.	1.0	2
138	Long-term effects of cannabinoids on development/behaviour. , 2020, 22, 33-37.		2
139	Detection of epileptic convulsions from accelerometry signals through machine learning approach. , 2014, , .		1
140	Novel treatment approaches and pediatric research networks in status epilepticus. Epilepsy and Behavior, 2019, 101, 106564.	1.7	1
141	Fenfluramine HCl Provides Long-Term Clinically Meaningful Reduction in Seizure Frequency: Results of an Open-Label Extension Study. Epilepsy and Behavior, 2019, 101, 106790.	1.7	1
142	Fenfluramine HCl (FinteplaÂ®) Provides Long-Term Clinically Meaningful Reduction in Seizure Frequency: Results of an Open-Label Extension Study. Neuropediatrics, 2019, 50, .	0.6	1
143	Interictal cardiorespiratory variability in temporal lobe and absence epilepsy in childhood. , 2014, , .		0
144	In response: Vagus nerve stimulation for epilepsy treatment in children. Epilepsia, 2015, 56, 324-325.	5.1	0

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145	Response to the numbering of seizure types. Epilepsia, 2017, 58, 1300-1301.	5.1	0
146	A Conjunctival Vascular Malformation as a Rare Presenting Sign of Wyburnâ€“Mason Syndrome. Journal of Pediatric Neurology, 2018, 16, 239-242.	0.2	0
147	Convulsive status epilepticus in children in Mozambique: Is there a treatment gap?. Journal of International Child Neurology Association, 2021, 1, .	0.0	0
148	Clinical Assessment of Visual Motion Perception in Children With Brain Damage: A Comparison With Base Rates and Control Sample. Frontiers in Human Neuroscience, 2021, 15, 733054.	2.0	0
149	SLC7A3: In Silico Prediction of a Potential New Cause of Childhood Epilepsy. Neuropediatrics, 2022, 53, 046-051.	0.6	0