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	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
2	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
3	Vaccination and childhood epilepsies. European Journal of Paediatric Neurology, 2022, 36, 57-68.	1.6	3
4	Association of Early MRI Characteristics With Subsequent Epilepsy and Neurodevelopmental Outcomes in Children With Tuberous Sclerosis Complex. Neurology, 2022, 98, .	1.1	8
5	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
6	Serotonin receptors in epilepsy: Novel treatment targets?. Epilepsia Open, 2022, 7, 231-246.	2.4	19
7	Treatment of Focal-Onset Seizures in Children: Should This Be More Etiology-Driven?. Frontiers in Neurology, 2022, 13, 842276.	2.4	4
8	SLC7A3: In Silico Prediction of a Potential New Cause of Childhood Epilepsy. Neuropediatrics, 2022, 53, 046-051.	0.6	0
9	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
10	Efficacy and Safety of Fenfluramine for the Treatment of Seizures Associated With Lennox-Gastaut Syndrome. JAMA Neurology, 2022, 79, 554.	9.0	43
11	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
12	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
13	A survey of the European Reference Network EpiCARE on clinical practice for selected rare epilepsies. Epilepsia Open, 2021, 6, 160-170.	2.4	3
14	Cognitive outcome in drugâ€resistant childhood epilepsy. Developmental Medicine and Child Neurology, 2021, 63, 633-633.	2.1	2
15	Early epileptiform EEG activity in infants with tuberous sclerosis complex predicts epilepsy and neurodevelopmental outcomes. Epilepsia, 2021, 62, 1208-1219.	5.1	19
16	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
17	PPFIA4 mutation: A second hit in POLG related disease?. Epilepsy and Behavior Reports, 2021, 16, 100455.	1.0	2
18	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

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19	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
20	The Ketogenic Diet Revisited: Beyond Ketones. Frontiers in Neurology, 2021, 12, 720073.	2.4	10
21	Convulsive status epilepticus in children in Mozambique: Is there a treatment gap?. Journal of International Child Neurology Association, 2021, 1, .	0.0	0
22	An update on congenital melanocytic nevus syndrome: A case report and literature review. Journal of Cutaneous Pathology, 2021, 48, 1497-1503.	1.3	5
23	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
24	Dravet syndrome. Current Opinion in Neurology, 2021, 34, 213-218.	3.6	26
25	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
26	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
27	Use of Zebrafish Models to Boost Research in Rare Genetic Diseases. International Journal of Molecular Sciences, 2021, 22, 13356.	4.1	15
28	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
29	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
30	Why we urgently need improved seizure and epilepsy therapies for children and neonates. Neuropharmacology, 2020, 170, 107854.	4.1	40
31	The concept of disease modification. European Journal of Paediatric Neurology, 2020, 24, 43-46.	1.6	6
32	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
33	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
34	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
35	Data describing child development at 6 years after maternal cancer diagnosis and treatment during pregnancy. Data in Brief, 2020, 32, 106209.	1.0	4
36	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

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37	Child development at 6 years after maternal cancer diagnosis and treatment during pregnancy. European Journal of Cancer, 2020, 138, 57-67.	2.8	31
38	Is autism driven by epilepsy in infants with Tuberous Sclerosis Complex?. Annals of Clinical and Translational Neurology, 2020, 7, 1371-1381.	3.7	23
39	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
40	Prediction of Neurodevelopment in Infants With Tuberous Sclerosis Complex Using Early EEG Characteristics. Frontiers in Neurology, 2020, 11, 582891.	2.4	19
41	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
42	Ketogenic diet for the treatment of pediatric epilepsy: review and meta-analysis. Child's Nervous System, 2020, 36, 1099-1109.	1.1	58
43	Epilepsy and Neurodevelopmental Comorbidities in Tuberous Sclerosis Complex: A Natural History Study. Pediatric Neurology, 2020, 106, 10-16.	2.1	37
44	Antiepileptic Drug Teratogenicity and De Novo Genetic Variation Load. Annals of Neurology, 2020, 87, 897-906.	5.3	9
45	Defining the phenotype of FHF1 developmental and epileptic encephalopathy. Epilepsia, 2020, 61, e71-e78.	5.1	11
46	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
47	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
48	Long-term effects of cannabinoids on development/behaviour. , 2020, 22, 33-37.		2
49	Epilepsy and cannabidiol: a guide to treatment. , 2020, 22, 1-14.		46
50	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
51	Long-term impact of prenatal exposure to chemotherapy on executive functioning: An ERP study. Clinical Neurophysiology, 2019, 130, 1655-1664.	1.5	3
52	Classification as autonomic versus sensory seizures. Epilepsia, 2019, 60, 2003-2005.	5.1	4
53	Novel treatment approaches and pediatric research networks in status epilepticus. Epilepsy and Behavior, 2019, 101, 106564.	1.7	1
54	Drug repurposing for Dravet syndrome in scn1Lab^{â^'/â^'} mutant zebrafish. Epilepsia, 2019, 60, e8-e13.	5.1	31

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55	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
56	2017 International League Against Epilepsy classifications of seizures and epilepsy are steps in the right direction. Epilepsia, 2019, 60, 1040-1044.	5.1	15
57	Diagnostic implications of genetic copy number variation in epilepsy plus. Epilepsia, 2019, 60, 689-706.	5.1	61
58	Genetic Modulation of Neurocognitive Development in Cancer Patients throughout the Lifespan: a Systematic Review. Neuropsychology Review, 2019, 29, 190-219.	4.9	9
59	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
60	Drug Development for Rare Paediatric Epilepsies: Current State and Future Directions. Drugs, 2019, 79, 1917-1935.	10.9	13
61	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
62	Dravet syndrome: Treatment options and management of prolonged seizures. Epilepsia, 2019, 60, S39-S48.	5.1	56
63	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
64	Patient MW: transient visual hemi-agnosia. Journal of Neurology, 2019, 266, 691-698.	3.6	4
65	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
66	Long-term effects of adjunctive perampanel on cognition in adolescents with partial seizures. Epilepsy and Behavior, 2018, 83, 50-58.	1.7	38
67	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
68	Is it safe for people with epilepsy to donate blood? A systematic review. Epilepsy Research, 2018, 139, 143-149.	1.6	2
69	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
70	A Conjunctival Vascular Malformation as a Rare Presenting Sign of Wyburnâ€“Mason Syndrome. Journal of Pediatric Neurology, 2018, 16, 239-242.	0.2	0
71	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
72	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

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73	Management of epilepsy associated with tuberous sclerosis complex: Updated clinical recommendations. European Journal of Paediatric Neurology, 2018, 22, 738-748.	1.6	151
74	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
75	Comparison and combination of electrocardiogram, electromyogram and accelerometry for tonic-clonic seizure detection in children. , 2018, , .		5
76	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
77	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
78	Vagus Nerve Stimulation in children: A focus on intellectual disability. European Journal of Paediatric Neurology, 2017, 21, 427-440.	1.6	26
79	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
80	Instruction manual for the <scp>ILAE</scp> 2017 operational classification of seizure types. Epilepsia, 2017, 58, 531-542.	5.1	699
81	Online Automated Seizure Detection in Temporal Lobe Epilepsy Patients Using Single-lead ECG. International Journal of Neural Systems, 2017, 27, 1750022.	5.2	42
82	Genetic and phenotypic heterogeneity suggest therapeutic implications in SCN2A-related disorders. Brain, 2017, 140, 1316-1336.	7.6	426
83	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
84	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
85	mTOR-related neuropathology in mutant tsc2 zebrafish: Phenotypic, transcriptomic and pharmacological analysis. Neurobiology of Disease, 2017, 108, 225-237.	4.4	29
86	Response to the numbering of seizure types. Epilepsia, 2017, 58, 1300-1301.	5.1	0
87	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
88	Pharmacological Analysis of the Anti-epileptic Mechanisms of Fenfluramine in scn1a Mutant Zebrafish. Frontiers in Pharmacology, 2017, 8, 191.	3.5	96
89	Inborn errors in RNA polymerase III underlie severe varicella zoster virus infections. Journal of Clinical Investigation, 2017, 127, 3543-3556.	8.2	125
90	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

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91	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
92	Gain-of-function FHF1 mutation causes early-onset epileptic encephalopathy with cerebellar atrophy. Neurology, 2016, 86, 2162-2170.	1.1	57
93	Fiveâ€year extended followâ€up status of 10 patients with Dravet syndrome treated with fenfluramine. Epilepsia, 2016, 57, e129-34.	5.1	97
94	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
95	Mutations in SNORD118 cause the cerebral microangiopathy leukoencephalopathy with calcifications and cysts. Nature Genetics, 2016, 48, 1185-1192.	21.4	114
96	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
97	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
98	Serotonergic Modulation as Effective Treatment for Dravet Syndrome in a Zebrafish Mutant Model. ACS Chemical Neuroscience, 2016, 7, 588-598.	3.5	86
99	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
100	Transient hypothyroidism associated with viral Human Parechovirus encephalitis in a newborn. European Journal of Paediatric Neurology, 2015, 19, 706-710.	1.6	3
101	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
102	Interictal cardiorespiratory variability in temporal lobe and absence epilepsy in childhood. Physiological Measurement, 2015, 36, 845-856.	2.1	19
103	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
104	Can ECG monitoring identify seizures?. Journal of Electrocardiology, 2015, 48, 1069-1074.	0.9	25
105	In response: Vagus nerve stimulation for epilepsy treatment in children. Epilepsia, 2015, 56, 324-325.	5.1	0
106	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
107	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
108	Interictal cardiorespiratory variability in temporal lobe and absence epilepsy in childhood. , 2014, , .		0

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109	Detection of epileptic seizures from single lead ECG by means of phase rectified signal averaging. , 2014, 2014, 3789-90.		5
110	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
111	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
112	The challenges and innovations for therapy in children with epilepsy. Nature Reviews Neurology, 2014, 10, 249-260.	10.1	38
113	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
114	Detecting rare events using extreme value statistics applied to epileptic convulsions in children. Artificial Intelligence in Medicine, 2014, 60, 89-96.	6.5	26
115	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
116	Overview of clinical efficacy and risk data of benzodiazepines for prolonged seizures. Epileptic Disorders, 2014, 16, 44-49.	1.3	5
117	Detection of epileptic convulsions from accelerometry signals through machine learning approach. , 2014, , .		1
118	Peri-ictal ECG changes in childhood epilepsy: Implications for detection systems. Epilepsy and Behavior, 2013, 29, 72-76.	1.7	48
119	Long-term home monitoring of hypermotor seizures by patient-worn accelerometers. Epilepsy and Behavior, 2013, 26, 118-125.	1.7	51
120	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
121	Ictal and interictal respiratory changes in temporal lobe and absence epilepsy in childhood. Epilepsy Research, 2013, 106, 410-416.	1.6	14
122	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
123	First line management of prolonged convulsive seizures in children and adults: good practice points. Acta Neurologica Belgica, 2013, 113, 375-380.	1.1	8
124	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
125	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
126	Successful use of fenfluramine as an addâ€on treatment for Dravet syndrome. Epilepsia, 2012, 53, 1131-1139.	5.1	175

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127	Clinical practice. European Journal of Pediatrics, 2011, 170, 413-418.	2.7	20
128	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
129	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
130	Developmental brain alterations in 17 year old boys are related to antenatal maternal anxiety. Clinical Neurophysiology, 2009, 120, 1116-1122.	1.5	64
131	The need for broad spectrum and safe anti-epileptic drugs in childhood epilepsy. Acta Neurologica Belgica, 2009, 109, 167-70.	1.1	7
132	Whatâ€™s new in: â€œGenetics in childhood epilepsyâ€• European Journal of Pediatrics, 2008, 167, 715-722.	2.7	14
133	Malignant migrating partial seizures in Aicardi syndrome. Developmental Medicine and Child Neurology, 2008, 50, 790-792.	2.1	6
134	ERP correlates of complex human decision making in a gambling paradigm: Detection and resolution of conflict. Psychophysiology, 2008, 45, 714-720.	2.4	45
135	Learning Disabilities: Definitions, Epidemiology, Diagnosis, and Intervention Strategies. Pediatric Clinics of North America, 2008, 55, 1259-1268.	1.8	63
136	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
137	Recommendations for the treatment of epilepsies in general practice in Belgium. Acta Neurologica Belgica, 2008, 108, 118-30.	1.1	4
138	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
139	Vagus nerve stimulation for refractory epilepsy: A Belgian multicenter study. European Journal of Paediatric Neurology, 2007, 11, 261-269.	1.6	118
140	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
141	Cognitive side effects of anti-epileptic drugs. Seizure: the Journal of the British Epilepsy Association, 2006, 15, 235-241.	2.0	96
142	Long-term cognitive sequelae of antenatal maternal anxiety: involvement of the orbitofrontal cortex. Neuroscience and Biobehavioral Reviews, 2006, 30, 1078-1086.	6.1	124
143	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
144	Severe Myoclonic Epilepsy in Infancy: Toward an Optimal Treatment. Journal of Child Neurology, 2004, 19, 516-521.	1.4	57

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145	Risperidone in the treatment of childhood autistic disorder: an open pilot study. Acta Neuropsychiatrica, 2002, 14, 242-249.	2.1	3
146	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
147	Cortical malformations: a frequent cause of epilepsy in children. European Journal of Pediatrics, 2000, 159, 555-562.	2.7	20
148	Treatment of a symptomatic posterior fossa subdural effusion in a child. Child's Nervous System, 1999, 15, 90-93.	1.1	7
149	Semi-Supervised One-Class Transfer Learning for Heart Rate Based Epileptic Seizure Detection. , 0, , .		4