Alberto Spalice

List of Publications by Year
in descending order

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183	4,184	109137	174990
papers	citations	35 h-index	52 g-index

194	194	194	4519
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Calcium-channel Blocker Verapamil Administration in Prolonged and Refractory Status Epilepticus. Epilepsia, 2005, 46, 967-969.	2.6	130
2	Earliest Clinical Manifestations and Natural History of Neurofibromatosis Type 2 (NF2) in Childhood: A Study of 24 Patients. Neuropediatrics, 2005, 36, 21-34.	0.3	124
3	Microbiota-gut brain axis involvement in neuropsychiatric disorders. Expert Review of Neurotherapeutics, 2019, 19, 1037-1050.	1.4	116
4	Cytomegalovirus infection and schizencephaly: Case reports. Annals of Neurology, 1998, 43, 123-127.	2.8	104
5	Neuronal migration disorders: clinical, neuroradiologic and genetics aspects. Acta Paediatrica, International Journal of Paediatrics, 2009, 98, 421-433.	0.7	101
6	Ohtahara syndrome with emphasis on recent genetic discovery. Brain and Development, 2012, 34, 459-468.	0.6	89
7	Pharmacotherapy of Spasticity in Children With Cerebral Palsy. Pediatric Neurology, 2006, 34, 1-6.	1.0	86
8	The natural history of spinal neurofibromatosis: a critical review of clinical and genetic features. Clinical Genetics, 2015, 87, 401-410.	1.0	81
9	Clinical spectrum of individuals with pathogenic N F1 missense variants affecting p.Met1149, p.Arg1276, and p.Lys1423: genotypeâ€“phenotype study in neurofibromatosis type 1. Human Mutation, 2020, 41, 299-315.	1.1	80
10	New trends in neuronal migration disorders. European Journal of Paediatric Neurology, 2010, 14, 1-12.	0.7	70
11	Efficacy and safety of levetiracetam: An add-on trial in children with refractory epilepsy. Seizure: the Journal of the British Epilepsy Association, 2005, 14, 248-253.	0.9	68
12	Addition of verapamil in the treatment of severe myoclonic epilepsy in infancy. Epilepsy Research, 2009, 85, 89-95.	0.8	68
13	Should â€œmigralepsyâ€•be considered an obsolete concept? A multicenter retrospective clinical/EEG study and review of the literature. Epilepsy and Behavior, 2011, 21, 52-59.	0.9	65
14	Effectiveness and tolerability of perampanel in children and adolescents with refractory epilepsiesâ€”An Italian observational multicenter study. Epilepsy Research, 2016, 127, 93-100.	0.8	62
15	Lacosamide in pediatric and adult patients: Comparison of efficacy and safety. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 210-216.	0.9	60
16	Gelastic epilepsy: video-EEG, MRI and SPECT characteristics. Brain and Development, 1997, 19, 418-421.	0.6	59
17	Benign convulsions associated with mild gastroenteritis: A multicenter clinical study. Epilepsy Research, 2011, 93, 107-114.	0.8	57
18	Ophthalmological manifestations in segmental neurofibromatosis type 1. British Journal of Ophthalmology, 2004, 88, 1429-1433.	2.1	55

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19	Neurofibromatosis type 1 and infantile spasms. Child's Nervous System, 2009, 25, 211-216.	0.6	55
20	Evaluation and management of nonsyndromic craniosynostosis. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 1185-1194.	0.7	55
21	Rufinamide in children and adults with Lennoxâ€“Gastaut syndrome: First Italian multicenter experience. Seizure: the Journal of the British Epilepsy Association, 2010, 19, 587-591.	0.9	52
22	Peri-ictal and inter-ictal headache in children and adolescents with idiopathic epilepsy: a multicenter cross-sectional study. Child's Nervous System, 2011, 27, 1419-1423.	0.6	50
23	Efficacy of verapamil as an adjunctive treatment in children with drug-resistant epilepsy: A pilot study. Seizure: the Journal of the British Epilepsy Association, 2014, 23, 36-40.	0.9	48
24	Efficacy and Safety of Topiramate in Refractory Epilepsy of Childhood. Journal of Child Neurology, 2005, 20, 893-897.	0.7	46
25	The genetics of monogenic idiopathic epilepsies and epileptic encephalopathies. Seizure: the Journal of the British Epilepsy Association, 2012, 21, 3-11.	0.9	46
26	Benign childhood epilepsy with centrotemporal spikes and the multicomponent model of attention: A matched control study. Epilepsy and Behavior, 2010, 19, 69-77.	0.9	44
27	Efficacy and safety of lacosamide in infants and young children with refractory focal epilepsy. European Journal of Paediatric Neurology, 2014, 18, 55-59.	0.7	44
28	Visual Field Constriction in Children With Epilepsy on Vigabatrin Treatment. Pediatrics, 2000, 106, 838-842.	1.0	44
29	Metabolic epilepsy: An update. Brain and Development, 2013, 35, 827-841.	0.6	43
30	Oxidative Stress and Gut-Derived Lipopolysaccharides in Neurodegenerative Disease: Role of NOX2. Oxidative Medicine and Cellular Longevity, 2020, 2020, 1-7.	1.9	42
31	Infantile spasms in the setting of Sturgeâ€“Weber syndrome. Child's Nervous System, 2009, 25, 111-118.	0.6	41
32	Gaming among Children and Adolescents during the COVID-19 Lockdown: The Role of Parents in Time Spent on Video Games and Gaming Disorder Symptoms. International Journal of Environmental Research and Public Health, 2021, 18, 6642.	1.2	41
33	Rufinamide in refractory childhood epileptic encephalopathies other than Lennox-Gastaut syndrome. European Journal of Neurology, 2011, 18, 246-251.	1.7	40
34	Clinical features of psychogenic non-epileptic seizures in prepubertal and pubertal patients with idiopathic epilepsy. Neurological Sciences, 2009, 30, 319-323.	0.9	38
35	Antiepileptic drug withdrawal in childhood epilepsy: What are the risk factors associated with seizure relapse?. European Journal of Paediatric Neurology, 2012, 16, 599-604.	0.7	38
36	â€œHeadache and epilepsyâ€•â€” How are they connected?. Epilepsy and Behavior, 2013, 26, 386-393.	0.9	38

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37	COVID-19-Related Anosmia: The Olfactory Pathway Hypothesis and Early Intervention. Frontiers in Neurology, 2020, 11, 956.	1.1	38
38	Paroxysmal Tonic Upgaze: Physiopathological Considerations in Three Additional Cases. Journal of Child Neurology, 2000, 15, 15-18.	0.7	37
39	Migraine treatment in developmental age: guidelines update. Journal of Headache and Pain, 2010, 11, 267-276.	2.5	36
40	Morphologic evidence of diffuse vascular damage in human and in the experimental model of ethylmalonic encephalopathy. Journal of Inherited Metabolic Disease, 2012, 35, 451-458.	1.7	35
41	Effect of d-ribose on purine synthesis and neurological symptoms in a patient with adenylosuccinase deficiency. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 1999, 1453, 135-140.	1.8	34
42	Pediatric Autoimmune Neuropsychiatry Disorder Associated with Group a Streptococcal Infection: The Role of Surgical Treatment. International Journal of Immunopathology and Pharmacology, 2014, 27, 371-378.	1.0	33
43	Macrocephaly-capillary malformation syndrome: Description of a case and review of clinical diagnostic criteria. Brain and Development, 2012, 34, 143-147.	0.6	32
44	Efficacy and tolerability of add-on lacosamide in children with Lennox-Gastaut syndrome. Acta Neurologica Scandinavica, 2014, 129, 420-424.	1.0	31
45	Vernal Keratoconjunctivitis: an update focused on clinical grading system. Italian Journal of Pediatrics, 2019, 45, 64.	1.0	31
46	Pharmacotherapy for children and adolescents with epilepsy. Expert Opinion on Pharmacotherapy, 2011, 12, 175-194.	0.9	30
47	Seizures and epilepsy in Sotos syndrome: Analysis of 19 Caucasian patients with longâ€term followâ€up. Epilepsia, 2012, 53, e102-5.	2.6	30
48	Long-term follow-up in children with benign convulsions associated with gastroenteritis. European Journal of Paediatric Neurology, 2014, 18, 572-577.	0.7	30
49	Neuronal migrational disorders in children with epilepsy: MRI, interictal SPECT and EEG comparisons. Brain and Development, 1996, 18, 269-279.	0.6	29
50	Dysembryoplastic Neuroepithelial Tumors: A Prospective Clinicopathologic and Outcome Study of 13 Children. Pediatric Neurology, 2010, 43, 395-402.	1.0	29
51	Epilepsy is a possible feature in Williamsâ€Beuren syndrome patients harboring typical deletions of the 7q11.23 critical region. American Journal of Medical Genetics, Part A, 2016, 170, 148-155.	0.7	29
52	Proton MR spectroscopy in connatal Pelizaeus-Merzbacher disease. Pediatric Radiology, 2000, 30, 171-175.	1.1	28
53	Recent Understanding on Diagnosis and Management of Central Nervous System Vasculitis in Children. Clinical and Developmental Immunology, 2012, 2012, 1-9.	3.3	28
54	Safety and efficacy of topiramate in neonates with hypoxic ischemic encephalopathy treated with hypothermia (NeoNATI). BMC Pediatrics, 2012, 12, 144.	0.7	28

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55	Mean Platelet Volume, Vitamin D and C Reactive Protein Levels in Normal Weight Children with Primary Snoring and Obstructive Sleep Apnea Syndrome. PLoS ONE, 2016, 11, e0152497.	1.1	28
56	The possible use of the L-type calcium channel antagonist verapamil in drug-resistant epilepsy. Expert Review of Neurotherapeutics, 2016, 16, 9-15.	1.4	28
57	Functional neuroradiologic investigations in band heterotopia. Pediatric Neurology, 2001, 24, 159-163.	1.0	27
58	Stroke in children: inherited and acquired factors and ageâ€related variations in the presentation of 48 paediatric patients. Acta Paediatrica, International Journal of Paediatrics, 2009, 98, 1130-1136.	0.7	27
59	Pretransplant conditioning with busulfan and cyclophosphamide in acute leukemia patients: Neurological and electroencephalographic prospective study. Annals of Oncology, 1992, 3, 145-148.	0.6	25
60	Neuronal migrational disorders: diffuse cortical dysplasia or the "double cortex" syndrome. Acta Paediatrica, International Journal of Paediatrics, 1993, 82, 501-503.	0.7	25
61	Electroclinical Features and Long-Term Outcome of Cryptogenic Epilepsy in Children with Down Syndrome. Journal of Pediatrics, 2013, 163, 1754-1758.	0.9	25
62	Seizures in fetal alcohol spectrum disorders: Evaluation of clinical, electroencephalographic, and neuroradiologic features in a pediatric case series. Epilepsia, 2014, 55, e60-6.	2.6	24
63	Epilepsy in the setting of full trisomy 18: A multicenter study on 18 affected children with and without structural brain abnormalities. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2016, 172, 288-295.	0.7	24
64	Epilepsy Course and Developmental Trajectories in STXBP1 -DEE. Neurology: Genetics, 2022, 8, .	0.9	24
65	Epilepsy in patients with Cornelia de Lange syndrome: A clinical series. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 356-359.	0.9	23
66	Cyclic Vomiting Syndrome in Children. Frontiers in Neurology, 2020, 11, 583425.	1.1	23
67	Results From an Italian Expanded Access Program on Cannabidiol Treatment in Highly Refractory Dravet Syndrome and Lennoxâ€“Gastaut Syndrome. Frontiers in Neurology, 2021, 12, 673135.	1.1	23
68	Long term outcome in children affected by absence epilepsy with onset before the age of three years. Epilepsy and Behavior, 2011, 20, 366-369.	0.9	22
69	“Epileptic Encephalopathy” of Infancy and Childhood: Electro-Clinical Pictures and Recent Understandings. Current Neuropharmacology, 2010, 8, 409-421.	1.4	21
70	Surgical timing of craniosynostosis: What to do and when. Journal of Cranio-Maxillo-Facial Surgery, 2014, 42, 513-519.	0.7	21
71	Neonatal Hyperglycemia Related to Parenteral Nutrition Affects Long-Term Neurodevelopment in Preterm Newborn: A Prospective Cohort Study. Nutrients, 2021, 13, 1930.	1.7	21
72	Neuroimaging Changes in Menkes Disease, Part 2. American Journal of Neuroradiology, 2017, 38, 1858-1865.	1.2	20

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73	Cerebral Palsy and Epilepsy in Children: Clinical Perspectives on a Common Comorbidity. Children, 2021, 8, 16.	0.6	20
74	Earlyâ€onset absence epilepsy: <scp>SLC</scp>2<scp>A</scp>1 gene analysis and treatment evolution. European Journal of Neurology, 2013, 20, 856-859.	1.7	19
75	Stroke and migraine is there a possible comorbidity?. Italian Journal of Pediatrics, 2016, 42, 41.	1.0	19
76	Coexistence of childhood absence epilepsy and benign epilepsy with centrotemporal spikes: A case series. European Journal of Paediatric Neurology, 2017, 21, 570-575.	0.7	19
77	Energy-enhanced parenteral nutrition and neurodevelopment of preterm newborns: A cohort study. Nutrition, 2021, 89, 111219.	1.1	19
78	Tensionâ€type headache in paediatric age. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 491-495.	0.7	18
79	Rufinamide for the treatment of refractory epilepsy secondary to neuronal migration disorders. Epilepsy Research, 2014, 108, 542-546.	0.8	18
80	Benign epilepsy of childhood with centrotemporal spikes and unilateral developmental opercular dysplasia. Child's Nervous System, 1994, 10, 264-269.	0.6	17
81	Residual and Persistent Adie's Pupil After Pediatric Ophthalmoplegic Migraine. Pediatric Neurology, 2009, 41, 204-206.	1.0	17
82	Complex malformation (Ruggieriâ€“Happle) phenotype with â€œcutis tricolorâ€•in a 10-year-old girl. Brain and Development, 2012, 34, 869-872.	0.6	17
83	WTX R353X mutation in a family with osteopathia striata and cranial sclerosis (OS-CS): case report and literature review of the disease clinical, genetic and radiological features. Italian Journal of Pediatrics, 2012, 38, 27.	1.0	17
84	Refractory absence seizures: An Italian multicenter retrospective study. European Journal of Paediatric Neurology, 2015, 19, 660-664.	0.7	17
85	Intravenous Immunoglobulin and Interferon: Successful Treatment of Optic Neuritis in Pediatric Multiple Sclerosis. Journal of Child Neurology, 2004, 19, 623-626.	0.7	16
86	Unusual association of SCN2A epileptic encephalopathy with severe cortical dysplasia detected by prenatal MRI. European Journal of Paediatric Neurology, 2017, 21, 587-590.	0.7	16
87	Clonazepam prophylaxis and busulfan-related myoclonic epilepsy in autografted acute leukemia patients. Haematologica, 1995, 80, 532-4.	1.7	16
88	Usefulness of diffusion tensor imaging and fiber tractography in neurological and neurosurgical pediatric diseases. Child's Nervous System, 2010, 26, 995-1002.	0.6	15
89	Sudden benzodiazepine-induced resolution of post-operative pediatric cerebellar mutism syndrome: a clinical-SPECT study. Acta Neurochirurgica, 2017, 159, 475-479.	0.9	15
90	Aromatic L-amino Acid Decarboxylase (AADC) deficiency: results from an Italian modified Delphi consensus. Italian Journal of Pediatrics, 2021, 47, 13.	1.0	15

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91	Clinical dissection of early onset absence epilepsy in children and prognostic implications. Epilepsia, 2013, 54, 1761-1770.	2.6	14
92	Early Myoclonic Encephalopathy in 9q33â€q34 Deletion Encompassing STXBP1 and SPTAN1. Annals of Human Genetics, 2015, 79, 209-217.	0.3	14
93	Clinical spectrum and genotype-phenotype correlations in PRRT2 Italian patients. European Journal of Paediatric Neurology, 2020, 28, 193-197.	0.7	14
94	Diffuse Onset of Ictal Electroencephalography in a Typical Case of Panayiotopoulos Syndrome and Review of the Literature. Journal of Child Neurology, 2009, 24, 472-476.	0.7	13
95	Bilateral (opercular and paracentral lobular) polymicrogyria and neurofibromatosis type 1. American Journal of Medical Genetics, Part A, 2011, 155, 582-585.	0.7	13
96	Effects of the abrupt switch from solution to modified-release granule formulation of valproate. Acta Neurologica Scandinavica, 2012, 125, e14-e18.	1.0	13
97	Severe early onset ethylmalonic encephalopathy with West syndrome. Metabolic Brain Disease, 2015, 30, 1537-1545.	1.4	13
98	Clinical dissection of childhood occipital epilepsy of Gastaut and prognostic implication. European Journal of Neurology, 2016, 23, 241-246.	1.7	13
99	Effects of early energy intake on neonatal cerebral growth of preterm newborn: an observational study. Scientific Reports, 2021, 11, 18457.	1.6	13
100	Seizures in paediatric Chiari type I malformation: the role of singlephoton emission computed tomography. Acta Paediatrica, International Journal of Paediatrics, 2002, 91, 313-317.	0.7	12
101	Bilateral Periventricular Nodular Heterotopia with Amniotic Band Syndrome. Pediatric Neurology, 2007, 36, 407-410.	1.0	12
102	Evaluation of the basal ganglia in neurofibromatosis type 1. Child's Nervous System, 2014, 30, 319-325.	0.6	12
103	Predictors of Evolution Into Multiple Sclerosis After a First Acute Demyelinating Syndrome in Children and Adolescents. Frontiers in Neurology, 2019, 9, 1156.	1.1	12
104	Identification of ultra-rare genetic variants in pediatric acute onset neuropsychiatric syndrome (PANS) by exome and whole genome sequencing. Scientific Reports, 2022, 12, .	1.6	12
105	Positron emission tomography in neuronal ceroid lipofuscinosis (Jansky-Bielschowsky disease): a case report. Brain and Development, 1994, 16, 459-462.	0.6	11
106	Primary CNS demyelinating diseases in childhood: multiple sclerosis. Child's Nervous System, 1996, 12, 149-154.	0.6	11
107	Ictal Single Photon Emission Computed Tomography in Absence Seizures: Apparent Implication of Different Neuronal Mechanisms. Journal of Child Neurology, 2001, 16, 339-344.	0.7	11
108	Epileptic nystagmus: Description of a pediatric case with EEG correlation and SPECT findings. Journal of the Neurological Sciences, 2010, 298, 127-131.	0.3	11

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109	Epilepsy in Menkes disease: An electroclinical long-term study of 28 patients. Epilepsy Research, 2014, 108, 1597-1603.	0.8	11
110	Valproate in adolescents with photosensitive epilepsy with generalized tonicâ€“clonic seizures only. European Journal of Paediatric Neurology, 2014, 18, 13-18.	0.7	11
111	Electroclinical findings and long-term outcomes in epileptic patients with inv dup (15). Acta Neurologica Scandinavica, 2018, 137, 575-581.	1.0	11
112	Risk Factors and Mental Health Promotion Strategies in Children During COVID-19. Frontiers in Public Health, 2020, 8, 580720.	1.3	11
113	Color vision and macular recovery time in epileptic adolescents treated with valproate and carbamazepine. European Journal of Neurology, 2006, 13, 736-741.	1.7	10
114	Long-term outcome of epilepsy in Kabuki syndrome. Seizure: the Journal of the British Epilepsy Association, 2011, 20, 650-654.	0.9	10
115	Cow's milk allergy and rolandic epilepsy: a close relationship?. Archives of Disease in Childhood, 2012, 97, 481.1-481.	1.0	10
116	Panayiotopoulos syndrome with convulsive status epilepticus at the onset: A long-term study. Seizure: the Journal of the British Epilepsy Association, 2014, 23, 728-731.	0.9	10
117	Long-term outcome of epilepsy in patients with Praderâ€“Willi syndrome. Journal of Neurology, 2015, 262, 116-123.	1.8	10
118	Vitamin D levels in children affected by vernal keratoconjunctivitis. Current Medical Research and Opinion, 2017, 33, 269-274.	0.9	10
119	Could Rolandic spikes be a prognostic factor of the neurocognitive outcome of children with BECTS?. Epilepsy and Behavior, 2018, 86, 157-162.	0.9	10
120	The Impact of the COVID-19 Epidemic During the Lockdown on Children With the Pediatric Acute-Onset Neuropsychiatric Syndrome (PANDAS/PANS): The Importance of Environmental Factors on Clinical Conditions. Frontiers in Neurology, 2021, 12, 702356.	1.1	10
121	COVID-19 and abducens nerve palsy in a 9-year-old girlâ€”case report. Italian Journal of Pediatrics, 2022, 48, .	1.0	10
122	Seizures in paediatric Chiari type I malformation: the role of singleâ€photon emission computed tomography. Acta Paediatrica, International Journal of Paediatrics, 2002, 91, 313-317.	0.7	9
123	Genotype-Phenotype Correlations in a Group of 15 SCN1A-Mutated Italian Patients with GEFS+ Spectrum (Seizures plus, Classical and Borderline Severe Myoclonic Epilepsy of Infancy). Journal of Child Neurology, 2010, 25, 1369-1376.	0.7	9
124	Fiber tractography assessment in double cortex syndrome. Child's Nervous System, 2011, 27, 1197-1202.	0.6	9
125	Periventricular Nodular Heterotopia: Report of a Pediatric Series. Journal of Child Neurology, 2002, 17, 300-304.	0.7	8
126	Pigmentary Mosaicism, Subcortical Band Heterotopia, and Brain Cystic Lesions. Pediatric Neurology, 2009, 40, 383-386.	1.0	8

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127	Reflex myoclonic epilepsy in infancy: A multicenter clinical study. Epilepsy Research, 2013, 103, 237-244.	0.8	8
128	Electroclinical features of epilepsy monosomy 1p36 syndrome and their implications. Acta Neurologica Scandinavica, 2018, 138, 523-530.	1.0	8
129	Small 7p22.3 microdeletion: Case report of Snx8 haploinsufficiency and neurological findings. European Journal of Medical Genetics, 2020, 63, 103772.	0.7	8
130	Gelastic seizures not associated with hypothalamic hamartoma: A long-term follow-up study. Epilepsy and Behavior, 2020, 103, 106578.	0.9	8
131	Nasal reactivity evaluation in children with allergic rhinitis receiving grass pollen sublingual immunotherapy. Allergy and Asthma Proceedings, 2020, 41, 357-362.	1.0	8
132	Bilateral perysilvian polymicrogyria in Chiari I malformation. Child's Nervous System, 2006, 22, 1635-1637.	0.6	7
133	Clinical and Pharmacological Aspects of Inflammatory Demyelinating Diseases in Childhood: An Update. Current Neuropharmacology, 2010, 8, 135-148.	1.4	7
134	Early add-on immunoglobulin administration in Rasmussen encephalitis: The hypothesis of neuroimmunomodulation. Medical Hypotheses, 2011, 77, 917-920.	0.8	7
135	Developmental anomalies of the medial septal area: possible implication for limbic epileptogenesis. Child's Nervous System, 2011, 27, 765-770.	0.6	7
136	Electroencephalogram and magnetic resonance imaging comparison as a predicting factor for neurodevelopmental outcome in hypoxic ischemic encephalopathy infant treated with hypothermia. Mental Illness, 2014, 6, 5532.	0.8	7
137	Oxidative stress and gut-derived lipopolysaccharides in children affected by paediatric autoimmune neuropsychiatric disorders associated with streptococcal infections. BMC Pediatrics, 2020, 20, 127.	0.7	7
138	Clinical Features in Patients With PANDAS/PANS and Therapeutic Approaches: A Retrospective Study. Frontiers in Neurology, 2021, 12, 741176.	1.1	7
139	Complex epileptic (Foixâ€“Chavanyâ€“Marie like) syndrome in a child with neurofibromatosis type 1 (NF1) and bilateral (opercular and paracentral) polymicrogyria. Acta Paediatrica, International Journal of Paediatrics, 2009, 98, 760-762.	0.7	6
140	Two siblings with a homozygous MTHFR C677T (G80A-RFC1) mutation and stroke. Child's Nervous System, 2009, 25, 361-365.	0.6	6
141	MTHFR Homozygous Mutation and Additional Risk Factors for Cerebral Infarction in a Large Italian Family. Pediatric Neurology, 2009, 40, 63-67.	1.0	6
142	Dilated Virchow-Robin spaces in children with seizures. A possible correlation?. Medical Hypotheses, 2020, 136, 109481.	0.8	6
143	Pediatrics for Disability: A Comprehensive Approach to Children with Syndromic Psychomotor Delay. Current Pediatric Reviews, 2022, 18, 110-120.	0.4	6
144	Chiari Type I Malformation, Syncope, Headache, Hypoglycemia and Hepatic Steatosis in an 8-Year Old Girl: A Causal Association?. Mental Illness, 2010, 2, e8.	0.8	5

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145	Longâ€term prognosis of patients with <scp>E</scp>hlersâ€<scp>D</scp>anlos syndrome and epilepsy. Epilepsia, 2014, 55, 1213-1219.	2.6	5
146	Spinal neurofibromatosis in a family with classical neurofibromatosis type 1 and a novel NF1 gene mutation. Journal of Clinical Neuroscience, 2014, 21, 328-330.	0.8	5
147	Focus on Cardiologic Findings in 30 Children With PANS/PANDAS: An Italian Single-Center Observational Study. Frontiers in Pediatrics, 2019, 7, 395.	0.9	5
148	Teaching NeuroImages: Schizencephaly in fetal alcohol syndrome. Neurology, 2011, 77, e96-e96.	1.5	4
149	Posterior fossa malformations and sex chromosomes anomalies. Report of a case with XYY syndrome and overview of known associations. European Journal of Pediatrics, 2013, 172, 1267-1270.	1.3	4
150	The crucial role of FBXO28 in the pathogenesis of the 1q41q42 microdeletion syndrome. American Journal of Medical Genetics, Part A, 2016, 170, 3041-3042.	0.7	4
151	EpiNet as a way of involving more physicians and patients in epilepsy research: Validation study and accreditation process. Epilepsia Open, 2017, 2, 20-31.	1.3	4
152	Eye movement desensitisation and reprocessing (EMDR) treatment associated with parent management training (PMT) for the acute symptoms in a patient with PANDAS syndrome: a case report. Italian Journal of Pediatrics, 2019, 45, 74.	1.0	4
153	Pediatric Autoimmune Neuropsychiatric Disorder Associated with Streptococcal Infection (PANDAS): Clinical Manifestations, IVIG Treatment Outcomes, Results from a Cohort of Italian Patients. Neuropsychiatry, 2018, 08, .	0.4	4
154	Stupor and Fast Activity on Electroencephalography in a Child Treated With Valproate. Pediatric Neurology, 2009, 41, 53-56.	1.0	3
155	The role of cytomegalovirus in schizencephaly. American Journal of Medical Genetics, Part A, 2011, 155, 1768-1768.	0.7	3
156	Teaching Neuro Images : Acute necrotizing encephalopathy during novel influenza A (H1N1) virus infection. Neurology, 2011, 77, e121.	1.5	3
157	Unilateral Lisch Nodules in a 47-year-old Woman Without Other Stigmata of Neurofibromatosis Type I: An Example of Segmental Neurofibromatosis?. Ophthalmic Genetics, 2013, 34, 178-179.	0.5	3
158	Photosensitivity as an early marker of epileptic and developmental encephalopathies. Epilepsia, 2018, 59, 1086-1087.	2.6	3
159	A new clinical feature associated with familial early-onset of dystonic-guttural tics: An unusual diagnosis of PANDAS. Journal of Pediatric Neurosciences, 2014, 9, 79.	0.2	3
160	Pediatric cerebellar stroke associated with elevated titer of antibodies to Î²2-glycoprotein. Medical Hypotheses, 2011, 76, 831-833.	0.8	2
161	Cardiovascular Risks of Ketogenic Diet for Glut-1 Deficiency. Pediatric Neurology Briefs, 2018, 32, 8.	0.2	2
162	Ictal Single Photon Emission Computed Tomography in Absence Seizures: Apparent Implication of Different Neuronal Mechanisms. Journal of Child Neurology, 2001, 16, 339.	0.7	2

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163	Novel missense mutation (L1917P) involving sac-domain of NSD1 gene in a patient with Sotos syndrome. Journal of Genetics, 2011, 90, 147-150.	0.4	1
164	Immunotherapy in Rasmussenâ€™s encephalitis: when should it be taken into account?. Neurological Sciences, 2013, 34, 1247-1248.	0.9	1
165	Spinal neurofibromatosis in children. Journal of Pediatrics, 2013, 162, 217.	0.9	1
166	Comorbidity of migraine in children presenting with epilepsy to a tertiary care center. Neurology, 2013, 80, 421-421.	1.5	1
167	Myoclonic status and central fever in Angelman syndrome due to paternal uniparental disomy. Journal of Neurogenetics, 2015, 29, 178-182.	0.6	1
168	Neuronal heterotopia. Neurology, 1995, 45, 1945-1945.	1.5	0
169	Displasie corticali: Nuove acquisizioni eziopatogenetiche. The Neuroradiology Journal, 1999, 12, 59-61.	0.1	0
170	Methylenetetrahydrofolate reductase homozygous mutation in a young boy with cerebellar infarction. Mental Illness, 2009, 1, e4.	0.8	0
171	Ageâ€related variation in the presentation of childhood stroke varies with inclusion criteria: Authorâ€™s reply. Acta Paediatrica, International Journal of Paediatrics, 2010, 99, 7-7.	0.7	0
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