Dimitrios I Zafeiriou

List of Publications by Year
in descending order

182	4,676	147801 31	133252 59
papers	citations	h-index	g-index

185	185	185	6439
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Leber congenital amaurosis: Comprehensive survey of the genetic heterogeneity, refinement of the clinical definition, and genotype-phenotype correlations as a strategy for molecular diagnosis. Human Mutation, 2004, 23, 306-317.	2.5	313
2	Tyrosine hydroxylase deficiency: a treatable disorder of brain catecholamine biosynthesis. Brain, 2010, 133, 1810-1822.	7.6	268
3	Exome Sequencing and the Management of Neurometabolic Disorders. New England Journal of Medicine, 2016, 374, 2246-2255.	27.0	254
4	Primitive reflexes and postural reactions in the neurodevelopmental examination. Pediatric Neurology, 2004, 31, 1-8.	2.1	166
5	Childhood autism and associated comorbidities. Brain and Development, 2007, 29, 257-272.	1.1	153
6	Intracerebral gene therapy in children with mucopolysaccharidosis type IIIB syndrome: an uncontrolled phase 1/2 clinical trial. Lancet Neurology, The, 2017, 16, 712-720.	10.2	149
7	Incidence and natural history of mucopolysaccharidosis type III in France and comparison with United Kingdom and Greece. American Journal of Medical Genetics, Part A, 2011, 155, 58-68.	1.2	133
8	Sepiapterin reductase deficiency: A Treatable Mimic of Cerebral Palsy. Annals of Neurology, 2012, 71, 520-530.	5.3	125
9	Brain and Spinal MR Imaging Findings in Mucopolysaccharidoses: A Review. American Journal of Neuroradiology, 2013, 34, 5-13.	2.4	112
10	An overview of L-2-hydroxyglutarate dehydrogenase gene (L2HGDH) variants: a genotype-phenotype study. Human Mutation, 2010, 31, 380-390.	2.5	108
11	Characteristics and Prognosis of Epilepsy in Children With Cerebral Palsy. Journal of Child Neurology, 1999, 14, 289-294.	1.4	94
12	<scp>l</scp>-2-Hydroxyglutaric Aciduria: Pattern of MR Imaging Abnormalities in 56 Patients. Radiology, 2009, 251, 856-865.	7.3	90
13	Obstetrical Brachial Plexus Palsy. Pediatric Neurology, 2008, 38, 235-242.	2.1	87
14	Autism spectrum disorders: The quest for genetic syndromes. American Journal of Medical Genetics Part B: Neuropsychiatric Genetics, 2013, 162, 327-366.	1.7	84
15	Management goals for type 1 Gaucher disease: An expert consensus document from the European working group on Gaucher disease. Blood Cells, Molecules, and Diseases, 2018, 68, 203-208.	1.4	82
16	Childhood Autism and Auditory System Abnormalities. Pediatric Neurology, 2010, 42, 309-314.	2.1	75
17	Pediatric Ischemic Stroke: An Infrequent Complication of <scp>SARSâ€CoV</scp>â€2. Annals of Neurology, 2021, 89, 657-665.	5.3	74
18	The Serotonergic System: Its Role in Pathogenesis and Early Developmental Treatment of Autism. Current Neuropharmacology, 2009, 7, 150-157.	2.9	64

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19	Update on transcobalamin deficiency: clinical presentation, treatment and outcome. Journal of Inherited Metabolic Disease, 2014, 37, 461-473.	3.6	59
20	Serial magnetic resonance imaging findings in mucopolysaccharidosis IIIB (Sanfilippo's syndrome B). Brain and Development, 2001, 23, 385-389.	1.1	56
21	Ethylmalonic Encephalopathy: Clinical and Biochemical Observations. Neuropediatrics, 2007, 38, 78-82.	0.6	47
22	Diagnosis of tuberous sclerosis complex in the fetus. European Journal of Paediatric Neurology, 2018, 22, 1027-1034.	1.6	46
23	Developing treatment options for metachromatic leukodystrophy. Molecular Genetics and Metabolism, 2012, 105, 56-63.	1.1	41
24	Neurophysiologic and intellectual evaluation of beta-thalassemia patients. Brain and Development, 2006, 28, 14-18.	1.1	40
25	Clinical features, laboratory findings and differential diagnosis of benign acute childhood myositis. Acta Paediatrica, International Journal of Paediatrics, 2000, 89, 1493-1494.	1.5	40
26	Pontocerebellar hypoplasia type 2: Variability in clinical and imaging findings. European Journal of Paediatric Neurology, 2007, 11, 146-152.	1.6	39
27	Endothelial activation and inflammation biomarkers in children and adolescents with sickle cell disease. International Journal of Hematology, 2013, 98, 158-163.	1.6	39
28	Xeroderma Pigmentosum Group G with Severe Neurological Involvement and Features of Cockayne Syndrome in Infancy. Pediatric Research, 2001, 49, 407-412.	2.3	36
29	Neurological complications in Î²-thalassemia. Brain and Development, 2006, 28, 477-481.	1.1	36
30	Prospective follow-up of primitive reflex profiles in high-risk infants: Clues to an early diagnosis of cerebral palsy. Pediatric Neurology, 1995, 13, 148-152.	2.1	35
31	Congenital Horner's syndrome associated with cervical neuroblastoma. European Journal of Paediatric Neurology, 2006, 10, 90-92.	1.6	34
32	3-Hydroxy-3-Methylglutaryl Coenzyme A Lyase Deficiency with Reversible White Matter Changes after Treatment. Pediatric Neurology, 2007, 37, 47-50.	2.1	33
33	Tyrosine hydroxylase deficiency with severe clinical course. Molecular Genetics and Metabolism, 2009, 97, 18-20.	1.1	33
34	Central nervous system abnormalities in asymptomatic young patients with S?-thalassemia. Annals of Neurology, 2004, 55, 835-839.	5.3	32
35	MR Spectroscopy and Serial Magnetic Resonance Imaging in a Patient with Mitochondrial Cystic Leukoencephalopathy due to Complex I Deficiency andNDUFV1Mutations and Mild Clinical Course. Neuropediatrics, 2008, 39, 172-175.	0.6	30
36	Stroke and Hypertension in Children and Adolescents. Journal of Child Neurology, 2017, 32, 408-417.	1.4	30

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37	A review of co-occurrence of autism spectrum disorder and Tourette syndrome. Research in Autism Spectrum Disorders, 2016, 24, 39-51.	1.5	29
38	Mortality After Pediatric Arterial Ischemic Stroke. Pediatrics, 2018, 141, .	2.1	29
39	Eponym. European Journal of Pediatrics, 2010, 169, 411-414.	2.7	28
40	Bayley-III scales at 12 months of corrected age in preterm infants: Patterns of developmental performance and correlations to environmental and biological influences. Research in Developmental Disabilities, 2015, 45-46, 110-119.	2.2	28
41	Identification of feeding risk factors for impaired nutrition status in paediatric patients with cerebral palsy. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, 649-654.	1.5	27
42	Early infantile Krabbe disease: deceptively normal magnetic resonance imaging and serial neurophysiological studies. Brain and Development, 1997, 19, 488-491.	1.1	26
43	Possible Genotype-Phenotype Correlations in Children with Mild Clinical Course of Canavan Disease. Neuropediatrics, 2005, 36, 252-255.	0.6	26
44	l-2-Hydroxyglutaric aciduria presenting with severe autistic features. Brain and Development, 2008, 30, 305-307.	1.1	26
45	Serial MRI and neurophysiological studies in late-infantile Krabbe disease. Pediatric Neurology, 1996, 15, 240-244.	2.1	25
46	Neurophysiologic Evaluation of Long-Term Desferrioxamine Therapy in Beta-Thalassemia Patients. Pediatric Neurology, 1998, 18, 420-424.	2.1	25
47	Clinical and Laboratory Data in a Sample of Greek Children with Autism Spectrum Disorders. Journal of Autism and Developmental Disorders, 2012, 42, 1470-1476.	2.7	25
48	Arterial Ischemic Stroke Secondary to Cardiac Disease in Neonates and Children. Pediatric Neurology, 2019, 100, 35-41.	2.1	25
49	Clinical and neurophysiological characteristics of congenital myasthenic syndromes presenting in early infancy. Brain and Development, 2004, 26, 47-52.	1.1	24
50	Spectrum ofSPATA7mutations in Leber congenital amaurosis and delineation of the associated phenotype. Human Mutation, 2010, 31, E1241-E1250.	2.5	24
51	Deficiency in complex IV (cytochrome c oxidase) of the respiratory chain, presenting as a leukodystrophy in two siblings with Leigh syndrome. Brain and Development, 1995, 17, 117-121.	1.1	23
52	Atypical and Variable Clinical Presentation of Glutaric Aciduria Type I. Neuropediatrics, 2000, 31, 303-306.	0.6	23
53	Two Greek siblings with sepiapterin reductase deficiency. Molecular Genetics and Metabolism, 2008, 94, 403-409.	1.1	23
54	Somatosensory Evoked Potentials in Children With Bilateral Spastic Cerebral Palsy. Pediatric Neurology, 2011, 44, 177-182.	2.1	23

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55	Protracted course of N-acetylaspartic aciduria in two non-Jewish siblings: identical clinical and magnetic resonance imaging findings. Brain and Development, 1999, 21, 205-208.	1.1	22
56	Niemann-pick type C disease associated with peripheral neuropathy. Pediatric Neurology, 2003, 29, 242-244.	2.1	22
57	Association between iron deficiency and febrile seizures. European Journal of Paediatric Neurology, 2015, 19, 591-596.	1.6	21
58	Insights into the expanding phenotypic spectrum of inherited disorders of biogenic amines. Nature Communications, 2021, 12, 5529.	12.8	21
59	Marinesco SjÃ¶gren Syndrome with Rhabdomyolysis. A New Subtype of the Disease. Neuropediatrics, 1998, 29, 97-101.	0.6	20
60	Neurophysiology and MRI in late-infantile metachromatic leukodystrophy. Pediatric Neurology, 1999, 21, 843-846.	2.1	20
61	Subclinical central nervous system involvement and thrombophilic status in young thalassemia intermedia patients of Greek origin. Blood Coagulation and Fibrinolysis, 2012, 23, 195-202.	1.0	20
62	Clinical course and seizure outcome of idiopathic childhood epilepsy: determinants of early and long-term prognosis. BMC Neurology, 2013, 13, 206.	1.8	20
63	Moyamoya syndrome and neurofibromatosis type 1. Italian Journal of Pediatrics, 2014, 40, 59.	2.6	20
64	Dyslipidemia in Children With Arterial Ischemic Stroke: Prevalence and Risk Factors. Pediatric Neurology, 2018, 78, 46-54.	2.1	20
65	Prenatal aspects in spinal muscular atrophy: From early detection to early presymptomatic intervention. European Journal of Paediatric Neurology, 2018, 22, 944-950.	1.6	20
66	Mutated SUCLG1 causes mislocalization of SUCLG2 protein, morphological alterations of mitochondria and an early-onset severe neurometabolic disorder. Molecular Genetics and Metabolism, 2019, 126, 43-52.	1.1	20
67	Seizures and Outcome One Year After Neonatal and Childhood Cerebral Sinovenous Thrombosis. Pediatric Neurology, 2020, 105, 21-26.	2.1	20
68	Plasmalogen levels in Gaucher disease. Blood Cells, Molecules, and Diseases, 2008, 41, 196-199.	1.4	19
69	Premonitory Urges and Their Link With Tic Severity in Children and Adolescents With Tic Disorders. Frontiers in Psychiatry, 2019, 10, 569.	2.6	19
70	Single dose immunoglobulin therapy for childhood Guillain-BarrÃ©syndrome. Brain and Development, 1997, 19, 323-325.	1.1	18
71	Diverse Clinical and Genetic Aspects of Craniofrontonasal Syndrome. Pediatric Neurology, 2011, 44, 83-87.	2.1	18
72	Castlemanâ€™s Disease Presenting as a Goiter in a Child. Hormone Research in Paediatrics, 2003, 59, 42-42.	1.8	17

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73	Cerebellar Agenesis and Diabetes Insipidus. Neuropediatrics, 2004, 35, 364-367.	0.6	17
74	Association of brain-derived neurotrophic factor (BDNF) and elongator protein complex 4 (ELP4) polymorphisms with benign epilepsy with centrotemporal spikes in a Greek population. Epilepsy Research, 2014, 108, 1734-1739.	1.6	17
75	Severe clinical presentation in monozygotic twins with 10p15.3 microdeletion syndrome. American Journal of Medical Genetics, Part A, 2014, 164, 764-768.	1.2	17
76	Using postural reactions as a screening test to identify high-risk infants for cerebral palsy: a prospective study. Brain and Development, 1998, 20, 307-311.	1.1	16
77	Familial moyamoya disease in a Greek family. Brain and Development, 2003, 25, 288-290.	1.1	16
78	Pandemic influenza A (H1N1) 2009-associated hemolytic uremic syndrome. Pediatric Nephrology, 2011, 26, 143-144.	1.7	16
79	Extracellular matrix components: An intricate network of possible biomarkers for lysosomal storage disorders?. FEBS Letters, 2013, 587, 1258-1267.	2.8	16
80	Neurological complications in childhood nephrotic syndrome: A systematic review. European Journal of Paediatric Neurology, 2019, 23, 384-391.	1.6	16
81	Adrenocorticotropic hormone and vigabatrin treatment of children with infantile spasms underlying cerebral palsy. Brain and Development, 1996, 18, 450-452.	1.1	15
82	Superior sagittal sinus thrombosis in steroid-resistant nephrotic syndrome. Pediatric Neurology, 2005, 32, 282-284.	2.1	15
83	Hemolytic Anemia Presenting With Idiopathic Intracranial Hypertension. Pediatric Neurology, 2008, 38, 53-54.	2.1	15
84	PEX1 deficiency presenting as Leber congenital amaurosis. Pediatric Neurology, 2004, 31, 146-149.	2.1	14
85	Nephrocalcinosis and Renal Failure in Lesch-Nyhan Syndrome: Report of Two Familial Cases and Review of the Literature. Urology, 2016, 97, 194-196.	1.0	14
86	Fetal stroke and cerebrovascular disease. European Journal of Paediatric Neurology, 2018, 22, 989-1005.	1.6	14
87	Reflex myoclonic epilepsy in infancy: a benign age-dependent idiopathic startle epilepsy. Epileptic Disorders, 2003, 5, 121-2.	1.3	14
88	L-2-Hydroxyglutaric aciduria presenting as status epilepticus. Brain and Development, 2001, 23, 255-257.	1.1	13
89	Visual Function in Preterm Infants without Major Retinopathy of Prematurity or Neurological Complications. American Journal of Perinatology, 2012, 29, 747-754.	1.4	13
90	Comparative Study of Refractive Errors, Strabismus, Microsaccades, and Visual Perception Between Preterm and Full-Term Children With Infantile Cerebral Palsy. Journal of Child Neurology, 2015, 30, 972-975.	1.4	13

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91	Impact of ambulatory SBP and overweight on executive function performance in children and adolescents. Journal of Hypertension, 2020, 38, 1123-1130.	0.5	13
92	Brain <scp>MR</scp> patterns in inherited disorders of monoamine neurotransmitters: An analysis of 70 patients. Journal of Inherited Metabolic Disease, 2021, 44, 1070-1082.	3.6	13
93	International Prevalence and Mechanisms of SARS-CoV-2 in Childhood Arterial Ischemic Stroke During the COVID-19 Pandemic. Stroke, 2022, 53, 2497-2503.	2.0	13
94	Plantar grasp reflex in high-risk infants during the first year of life. Pediatric Neurology, 2000, 22, 75-76.	2.1	12
95	Pontocerebellar Hypoplasia in Extreme Prematurity: Clinical and Neuroimaging Findings. Pediatric Neurology, 2013, 48, 48-51.	2.1	12
96	The Spectrum of Niemann-Pick Type C Disease in Greece. JIMD Reports, 2017, 36, 41-48.	1.5	12
97	Neurological outcome at 6 and 12 months corrected age in hospitalised late preterm infants -a prospective study. European Journal of Paediatric Neurology, 2018, 22, 602-609.	1.6	12
98	Benign congenital hemifacial spasm. Pediatric Neurology, 1997, 17, 174-176.	2.1	11
99	Russellâ€™s diencephalic syndrome. Neurology, 2001, 57, 932-932.	1.1	11
100	Serial magnetic resonance imaging and neurophysiological studies in multiple sulphatase deficiency. European Journal of Paediatric Neurology, 2008, 12, 190-194.	1.6	11
101	Episodic ataxia type 2 showing ictal hyperhidrosis with hypothermia and interictal chronic diarrhea due to a novel CACNA1A mutation. European Journal of Paediatric Neurology, 2009, 13, 191-193.	1.6	11
102	Sleep disorders and executive function in children and adolescents with chronic kidney disease. Sleep Medicine, 2019, 55, 33-39.	1.6	11
103	Pregabalin adjunctive therapy for focal onset seizures in children 1 month to <4 years of age: A doubleâ€blind, placeboâ€controlled, videoâ€electroencephalographic trial. Epilepsia, 2020, 61, 617-626.	5.1	11
104	Clinical features, laboratory findings and differential diagnosis of benign acute childhood myositis. Acta Paediatrica, International Journal of Paediatrics, 2000, 89, 1493-1494.	1.5	11
105	Hypoparathyroidism and intracranial calcifications in Î²-thalassemia major. Journal of Pediatrics, 2001, 138, 411.	1.8	9
106	Hemimegalencephaly and tuberous sclerosis complex: A rare yet challenging association. European Journal of Paediatric Neurology, 2021, 30, 58-65.	1.6	9
107	Intracerebral Gene Therapy in Four Children with Sanfilippo B Syndrome: 5.5-Year Follow-Up Results. Human Gene Therapy, 2021, 32, 1251-1259.	2.7	9
108	Brainstem Auditory Evoked Potentials in Boys with Autism: Still Searching for the Hidden Truth. Iranian Journal of Child Neurology, 2015, 9, 21-8.	0.3	9

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109	Screening for minor changes in the distal part of the human dystrophin gene in Greek DMD/BMD patients. European Journal of Human Genetics, 1999, 7, 179-187.	2.8	8
110	Utility of brainstem auditory evoked potentials in children with spastic cerebral palsy. Acta Paediatrica, International Journal of Paediatrics, 2000, 89, 194-197.	1.5	8
111	Hypoglossal-Nerve Palsy. New England Journal of Medicine, 2004, 350, e4.	27.0	8
112	Thelarche variant in a girl with Angelman syndrome. Brain and Development, 2004, 26, 339-341.	1.1	8
113	Self-mutilation and mental retardation: clues to congenital insensitivity to pain with anhidrosis. Journal of Pediatrics, 2004, 144, 284.	1.8	8
114	Differential Expression of Matrix Metalloproteinases in the Serum of Patients with Mucopolysaccharidoses. JIMD Reports, 2011, 3, 59-66.	1.5	8
115	Multiple Endocrine Disorders Associated With Adrenomyeloneuropathy and a Novel Mutation of the ABCD1 Gene. Pediatric Neurology, 2014, 50, 622-624.	2.1	8
116	Screening for TSC1 and TSC2 mutations using NGS in Greek children with tuberous sclerosis syndrome. European Journal of Paediatric Neurology, 2018, 22, 419-426.	1.6	8
117	Matrix metalloproteinase â˜2, â˜9 and arterial stiffness in children and adolescents: The role of chronic kidney disease, diabetes, and hypertension. International Journal of Cardiology: Hypertension, 2020, 4, 100025.	2.2	8
118	Hypertension and childhood stroke. Pediatric Nephrology, 2021, 36, 809-823.	1.7	8
119	Brittle Hair, Photosensitivity, Brain Hypomyelination and Immunodeficiency: Clues to Trichothiodystrophy. Indian Journal of Pediatrics, 2017, 84, 89-90.	0.8	7
120	Assessment of intellectual impairment, healthâ€related quality of life, and behavioral phenotype in patients with neurotransmitter related disorders: Data from the <scp>iNTD</scp> registry. Journal of Inherited Metabolic Disease, 2021, 44, 1489-1502.	3.6	7
121	Three Individuals with PURA Syndrome in a Cohort of Patients with Neuromuscular Disease. Neuropediatrics, 2021, 52, 390-393.	0.6	7
122	Malignant hyperthermia and Marinesco-SjÃ¶gren syndrome. Lancet, The, 1991, 338, 1603.	13.7	6
123	Plantar Response Profile of High-Risk Infants at One Year of Life. Journal of Child Neurology, 1999, 14, 514-517.	1.4	6
124	Test of Everyday Attention for Children (TEA-Ch): Greek Normative Data and Discriminative Validity for Children with Combined Type of Attention Deficit-Hyperactivity Disorder. Developmental Neuropsychology, 2019, 44, 189-202.	1.4	6
125	Protracted course of N-acetylaspartic aciduria in two non-Jewish siblings: identical clinical and magnetic resonance imaging findings. Brain and Development, 1999, 21, 205-8.	1.1	6
126	Childhood steroid-responsive painful opthalmoplegia: Clues to opthalmoplegic migraine. Journal of Pediatrics, 2006, 149, 881.e1-881.e2.	1.8	5

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127	A patient with Lemierre syndrome. European Journal of Pediatrics, 2010, 169, 491-493.	2.7	5
128	Early-onset diabetes mellitus and neurodevelopmental retardation: the first Greek case of Wolcott-Rallison syndrome. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 967-970.	0.9	5
129	Growth hormone replacement therapy in Costello syndrome. Growth Hormone and IGF Research, 2014, 24, 271-275.	1.1	5
130	Fetal ventriculomegaly: What we have and what is still missing. European Journal of Paediatric Neurology, 2018, 22, 898-899.	1.6	5
131	Benign epilepsy with centrotemporal spikes: Relationship between type of seizures and response to medication in a Greek population. Journal of Neurosciences in Rural Practice, 2015, 06, 545-548.	0.8	5
132	Pseudoaneurysm of the Popliteal Artery Complicated by Peroneal Mononeuropathy in a 4-year-old Child: Report of a Case. Surgery Today, 2007, 37, 798-801.	1.5	4
133	MYH9-related Disorders. Journal of Pediatric Hematology/Oncology, 2012, 34, 412-415.	0.6	4
134	Recurrent episodes of rhabdomyolysis in pontocerebellar hypoplasia type 2. Neuromuscular Disorders, 2013, 23, 116-119.	0.6	4
135	Multiple Coronary Artery Microfistulas in a Girl with Kleefstra Syndrome. Case Reports in Genetics, 2016, 2016, 1-5.	0.2	4
136	Neuroradiological, neurophysiological and molecular findings in infantile Krabbe disease: two case reports. Balkan Journal of Medical Genetics, 2016, 19, 85-90.	0.5	4
137	An emerging cause of concern in Europe: Zika virus, the developing CNS and the pediatric neurologist. European Journal of Paediatric Neurology, 2016, 20, 497-499.	1.6	4
138	Uric Acid Associates With Executive Function in Children and Adolescents With Hypertension. Hypertension, 2021, 77, 1737-1744.	2.7	4
139	Is the prevalence of thyroid disease higher in children receiving antiepileptic medication? A systematic review and meta-analysis. Seizure: the Journal of the British Epilepsy Association, 2022, 94, 117-125.	2.0	4
140	Targeted Genotyping of MIS-C Patients Reveals a Potential Alternative Pathway Mediated Complement Dysregulation during COVID-19 Infection. Current Issues in Molecular Biology, 2022, 44, 2811-2824.	2.4	4
141	Immune and Neural Status of Thalassemic Patients Receiving Deferiprone or Combined Deferiprone and Deferoxamine Chelation Treatment. Hemoglobin, 2008, 32, 35-40.	0.8	3
142	Parental reports of health-related quality of life in greek children with neurofibromatosis type 1. Journal of Pediatrics, 2009, 155, 453.	1.8	3
143	Noninvasive ultra high-frequency (1kHz) oscillationsâ€™ recording: High-fidelity over somatosensory cortex. Clinical Neurophysiology, 2012, 123, 2323-2324.	1.5	3
144	Paroxysmal tonic upward gaze of childhood â€œplusâ€: An oculomotor channelopathy?. European Journal of Paediatric Neurology, 2015, 19, 278-279.	1.6	3

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145	Diagnosis of tuberous sclerosis complex in a patient referred for uncontrolled hypertension and renal dysfunction. Journal of Hypertension, 2017, 35, 2109-2114.	0.5	3
146	Central SBP and executive function in children and adolescents with primary and secondary hypertension. Journal of Hypertension, 2020, 38, 2176-2184.	0.5	3
147	The landscape of Mucopolysaccharidosis in Southern and Eastern European countries: a survey from 19 specialistic centers. Orphanet Journal of Rare Diseases, 2022, 17, 136.	2.7	3
148	Integrative Approach to Predict Severity in Nonketotic Hyperglycinemia. Annals of Neurology, 2022, 92, 292-303.	5.3	3
149	Prospective study of ankle clonus at the first year of life. Brain and Development, 1997, 19, 440.	1.1	2
150	A case presenting with delayed motor milestones. Neuromuscular Disorders, 2005, 15, 817-818.	0.6	2
151	Unraveling the brainstem mysteries in late-preterm infants. Clinical Neurophysiology, 2012, 123, 852-853.	1.5	2
152	Incontinentia Pigmenti: A Skin, Brain, and Eye Matter. Journal of Pediatrics, 2013, 163, 1520.	1.8	2
153	Alexander Disease. Journal of Pediatrics, 2013, 162, 648.	1.8	2
154	<scp>S</scp>tÃ¼veâ€“<scp>W</scp>iedemann syndrome in a neonate. Pediatrics International, 2015, 57, 302-304.	0.5	2
155	Investigation of the motor system in two siblings with Canavanâ€™s disease: a combined transcranial magnetic stimulation (TMS) â€“ diffusion tensor imaging (DTI) study. Metabolic Brain Disease, 2017, 32, 307-310.	2.9	2
156	Application of high-resolution array comparative genomic hybridization in children with unknown syndromic microcephaly. Pediatric Research, 2017, 82, 253-260.	2.3	2
157	Arterial Stiffness in a Toddler with Neurofibromatosis Type 1 and Refractory Hypertension. Case Reports in Pediatrics, 2018, 2018, 1-4.	0.4	2
158	Hemolytic Uremic Syndrome Due to Methylmalonic Acidemia and Homocystinuria in an Infant: A Case Report and Literature Review. Children, 2021, 8, 112.	1.5	2
159	The Complex Interplay of Cortex, Cerebellum, and Age in a Cohort of Pediatric Patients With Tuberous Sclerosis Complex. Pediatric Neurology, 2021, 123, 43-49.	2.1	2
160	Consensus statement on enzyme replacement therapy for mucopolysaccharidosis IVA in Central and South-Eastern European countries. Orphanet Journal of Rare Diseases, 2022, 17, 190.	2.7	2
161	Single High-Dose Immunoglobulin Therapy for Childhood Guillain-BarrÃ© Syndrome. Journal of Child Neurology, 1999, 14, 480-481.	1.4	1
162	Ulnar hypoplasia and neurofibromatosis type I. Journal of Pediatrics, 2004, 145, 859.	1.8	1

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163	High-fidelity over the somatosensory cortex revisited: Back to basics. Clinical Neurophysiology, 2015, 126, 223-224.	1.5	1
164	Neurophysiologic evaluation of infants with congenital hypothyroidism before and after treatment. Acta Neurologica Belgica, 2015, 115, 129-136.	1.1	1
165	Neurological Involvement in a Thalassemic Child Receiving Deferiprone. Journal of Pediatric Hematology/Oncology, 2016, 38, 406-406.	0.6	1
166	Response to â€œLeigh-like syndrome with mild mtDNA depletion due to the SUCLG1 variant c.626C>Aâ€• Molecular Genetics and Metabolism Reports, 2019, 18, 10.	1.1	1
167	Gene therapy for spinal muscular atrophy: Solomon's consensus in Covid times. European Journal of Paediatric Neurology, 2020, 28, 2-3.	1.6	1
168	Subjective sleepâ€related breathing disorders and executive function in children with intermittent or mild persistent asthma. Clinical Respiratory Journal, 2021, 15, 794-799.	1.6	1
169	Phenotype assessment in neurologically impaired paediatric patients: Impact of a nutrition intervention protocol. Clinical Nutrition, 2021, 40, 5734-5741.	5.0	1
170	Mild myopathic phenotype in a patient with homozygous c.416C > T mutation in gene. Acta Myologica, 2020, 39, 94-97.	1.5	1
171	Blood pressure and glomerular filtration rate in youth with tuberous sclerosis complex. European Journal of Pediatrics, 2022, 181, 1465-1472.	2.7	1
172	Response:. Pediatric Neurology, 2008, 39, 371.	2.1	0
173	Flash VEP findings in children with bilateral spastic cerebral palsy: Correlation with clinico-radiological parameters. Journal of Pediatric Neurology, 2015, 06, 329-339.	0.2	0
174	Cranial nerve palsies as presenting feature of acute lymphoblastic leukemia. Journal of Pediatric Neurology, 2015, 11, 055-057.	0.2	0
175	Cerebral embolism secondary to atrial myxoma in a child. Journal of Pediatric Neurology, 2015, 10, 317-318.	0.2	0
176	The brainstem in late preterm birth: Born small-for-gestational-age is the â€œtip of the icebergâ€• Clinical Neurophysiology, 2016, 127, 3178-3179.	1.5	0
177	Clinical Aspects of Pediatric Brain Tumors. Journal of Pediatric Neuroradiology, 2016, 05, 046-048.	0.1	0
178	Correspondence regarding the systematic review entitled â€˜Evidence-based treatment of multicystic dysplastic kidney: a systematic reviewâ€™. Journal of Pediatric Urology, 2019, 15, 291-292.	1.1	0
179	Age of first diagnosis in children with cerebral palsy - the need for new developmental screening tests. Acta Paediatrica, International Journal of Paediatrics, 1998, 87, 814-814.	1.5	0
180	Two hands are better than one: Mind the grasp. European Journal of Paediatric Neurology, 2022, 37, A2.	1.6	0

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