Piero Pavone

List of Publications by Year
in descending order

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300	5,774	117625	138484
papers	citations	34 h-index	58 g-index

319	319	319	6138
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	A Genomewide Screen for Autism Susceptibility Loci. American Journal of Human Genetics, 2001, 69, 327-340.	6.2	287
2	Hepatic echinococcosis: Clinical and therapeutic aspects. World Journal of Gastroenterology, 2012, 18, 1448.	3.3	253
3	Neuropsychological assessment in children with absence epilepsy. Neurology, 2001, 56, 1047-1051.	1.1	168
4	Infantile spasms syndrome, West syndrome and related phenotypes: What we know in 2013. Brain and Development, 2014, 36, 739-751.	1.1	159
5	Worldwide distribution and broader clinical spectrum of muscle-eye-brain disease. Human Molecular Genetics, 2003, 12, 527-534.	2.9	133
6	Anti-brain antibodies in PANDAS versus uncomplicated streptococcal infection. Pediatric Neurology, 2004, 30, 107-110.	2.1	114
7	The neurology of coeliac disease in childhood: what is the evidence? A systematic review and metaâ€analysis. Developmental Medicine and Child Neurology, 2010, 52, 700-707.	2.1	92
8	Ohtahara syndrome with emphasis on recent genetic discovery. Brain and Development, 2012, 34, 459-468.	1.1	89
9	Septo-Optic Dysplasia Complex: A Heterogeneous Malformation Syndrome. Pediatric Neurology, 2006, 34, 66-71.	2.1	81
10	The natural history of spinal neurofibromatosis: a critical review of clinical and genetic features. Clinical Genetics, 2015, 87, 401-410.	2.0	81
11	Acute Disseminated Encephalomyelitis: A Long-Term Prospective Study and Meta-Analysis. Neuropediatrics, 2010, 41, 246-255.	0.6	75
12	West syndrome: a comprehensive review. Neurological Sciences, 2020, 41, 3547-3562.	1.9	73
13	Epilepsy is not a Prominent Feature of Primary Autism. Neuropediatrics, 2004, 35, 207-210.	0.6	71
14	Low Prevalence of Neurologic and Psychiatric Manifestations in Children with Gluten Sensitivity. Journal of Pediatrics, 2008, 152, 244-249.e1.	1.8	71
15	Hydranencephaly: cerebral spinal fluid instead of cerebral mantles. Italian Journal of Pediatrics, 2014, 40, 79.	2.6	71
16	Topical Review: Autoimmune Neuropsychiatric Disorders Associated With Streptococcal Infection: Sydenham Chorea, PANDAS, and PANDAS Variants. Journal of Child Neurology, 2006, 21, 727-736.	1.4	67
17	Congenital muscular dystrophy: from muscle to brain. Italian Journal of Pediatrics, 2016, 42, 78.	2.6	64
18	Temporal Triangular Alopecia in Association With Mental Retardation and Epilepsy in a Mother and Daughter. Archives of Dermatology, 2000, 136, 426-427.	1.4	64

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19	Pediatric autoimmune encephalitis. Journal of Pediatric Neurosciences, 2017, 12, 130.	0.3	61
20	Lacosamide in pediatric and adult patients: Comparison of efficacy and safety. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 210-216.	2.0	60
21	Epilepsy and innate immune system: A possible immunogenic predisposition and related therapeutic implications. Human Vaccines and Immunotherapeutics, 2015, 11, 2021-2029.	3.3	59
22	Ataxia in children: early recognition and clinical evaluation. Italian Journal of Pediatrics, 2017, 43, 6.	2.6	59
23	Silent COVID-19: what your skin can reveal. Lancet Infectious Diseases, The, 2021, 21, 24-25.	9.1	56
24	Ophthalmological manifestations in segmental neurofibromatosis type 1. British Journal of Ophthalmology, 2004, 88, 1429-1433.	3.9	55
25	Headache in Pediatric Patients With Celiac Disease and Its Prevalence as a Diagnostic Clue. Journal of Pediatric Gastroenterology and Nutrition, 2009, 49, 202-207.	1.8	55
26	Neurofibromatosis type 1 and infantile spasms. Child's Nervous System, 2009, 25, 211-216.	1.1	55
27	Natural history of neurofibromatosis type 2 with onset before the age of 1Âyear. Neurogenetics, 2013, 14, 89-98.	1.4	55
28	Nevus vascularis mixtus (cutaneous vascular twin nevi) associated with intracranial vascular malformation of the Dykeâ€“Davidoffâ€“Masson type in two patients. American Journal of Medical Genetics, Part A, 2012, 158A, 2870-2880.	1.2	54
29	Inherited human IRAK-1 deficiency selectively impairs TLR signaling in fibroblasts. Proceedings of the National Academy of Sciences of the United States of America, 2017, 114, E514-E523.	7.1	49
30	Increased Antistreptococcal Antibody Titers and Antiâ€”Basal Ganglia Antibodies in Patients With Tourette Syndrome: Controlled Cross-Sectional Study. Journal of Child Neurology, 2006, 21, 747-753.	1.4	48
31	Targeting inflammation as a therapeutic strategy for drug-resistant epilepsies. Human Vaccines and Immunotherapeutics, 2014, 10, 868-875.	3.3	47
32	Wide spectrum of congenital anomalies including choanal atresia, malformed extremities, and brain and spinal malformations in a girl with a de novo 5.6â€Mb deletion of 13q12.11â€“13q12.13. American Journal of Medical Genetics, Part A, 2014, 164, 1734-1743.	1.2	46
33	Ketogenic diet for infants with epilepsy: A literature review. Epilepsy and Behavior, 2020, 112, 107361.	1.7	46
34	A clinical review on megalencephaly. Medicine (United States), 2017, 96, e6814.	1.0	44
35	The role of polytherapy in the management of epilepsy: suggestions for rational antiepileptic drug selection. Expert Review of Neurotherapeutics, 2020, 20, 167-173.	2.8	43
36	Infantile spasms in the setting of Sturgeâ€“Weber syndrome. Child's Nervous System, 2009, 25, 111-118.	1.1	41

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37	Primary Headaches in Children: Clinical Findings and the Association with other Conditions. International Journal of Immunopathology and Pharmacology, 2012, 25, 1083-1091.	2.1	41
38	Epilepsy in Children: From Diagnosis to Treatment with Focus on Emergency. Journal of Clinical Medicine, 2019, 8, 39.	2.4	40
39	The immunomodulatory effect of probiotics beyond atopy: an update. Journal of Asthma, 2014, 51, 320-332.	1.7	39
40	Lack of progression of brain atrophy in Aicardi-GoutiÃ¨res syndrome. Pediatric Neurology, 2001, 24, 300-302.	2.1	38
41	Congenital generalized hypertrichosis: the skin as a clue to complex malformation syndromes. Italian Journal of Pediatrics, 2015, 41, 55.	2.6	38
42	A Case of COVID-19 with Late-Onset Rash and Transient Loss of Taste and Smell in a 15-Year-Old Boy. American Journal of Case Reports, 2020, 21, e925813.	0.8	38
43	Hypomelanosis of Ito: a round on the frequency and type of epileptic complications. Neurological Sciences, 2015, 36, 1173-1180.	1.9	37
44	STROKE IN TWO CHILDREN WITH MYCOPLASMA PNEUMONIAE INFECTION A CAUSAL OR CASUAL RELATIONSHIP?. Pediatric Infectious Disease Journal, 2005, 24, 843-845.	2.0	36
45	Congenital talipes equinovarus: an epidemiological study in Sicily. Monthly Notices of the Royal Astronomical Society: Letters, 2012, 83, 294-298.	3.3	36
46	Randomized Comparison of Helmet CPAP Versus High-Flow Nasal Cannula Oxygen in Pediatric Respiratory Distress. Respiratory Care, 2017, 62, 1036-1042.	1.6	35
47	Hypertrichosis Cubiti (Hairy Elbow Syndrome): A Clue to a Malformation Syndrome. Journal of Pediatric Endocrinology and Metabolism, 2005, 18, 1019-25.	0.9	34
48	West Syndrome Treatment: New Roads for an Old Syndrome. Frontiers in Neurology, 2013, 4, 113.	2.4	34
49	Marin-Amat Syndrome: Case Report and Review of the Literature. Journal of Child Neurology, 1999, 14, 266-268.	1.4	33
50	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.	2.1	33
51	Total Hemi-overgrowth in Pigmentary Mosaicism of the (Hypomelanosis of) Ito Type. Medicine (United) Tj ETQq1 1 0.784314 rgBT /Ov	1.0	33
52	Levetiracetam in neonatal seizures as first-line treatment: A prospective study. Journal of Pediatric Neurosciences, 2017, 12, 24.	0.3	33
53	Unilateral Eye Blinking Arising From the Ictal Ipsilateral Occipital Area. Clinical EEG and Neuroscience, 2016, 47, 243-246.	1.7	32
54	Prognostic Challenges of SCN1A Genetic Mutations: Report on Two Children with Mild Features. Journal of Pediatric Neurology, 2016, 14, 082-088.	0.2	32

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55	Acquired Peripheral Neuropathy: A Report on 20 Children. International Journal of Immunopathology and Pharmacology, 2012, 25, 513-517.	2.1	31
56	Diagnosis and management of catamenial seizures: a review. International Journal of Women's Health, 2012, 4, 535.	2.6	31
57	Recurrent peripheral neuropathy in a girl with celiac disease. Journal of Neurology, Neurosurgery and Psychiatry, 2000, 68, 104-105.	1.9	30
58	Hemihydranencephaly: living with half brain dysfunction. Italian Journal of Pediatrics, 2013, 39, 3.	2.6	30
59	Clinical guidelines in pediatric headache: evaluation of quality using the AGREE II instrument. Journal of Headache and Pain, 2014, 15, 57.	6.0	30
60	Protective effects of the sigma agonist Pre-084 in the rat retina. British Journal of Ophthalmology, 2007, 91, 1382-1384.	3.9	29
61	Dysembryoplastic Neuroepithelial Tumors: A Prospective Clinicopathologic and Outcome Study of 13 Children. Pediatric Neurology, 2010, 43, 395-402.	2.1	29
62	A neurocutaneous phenotype with paired hypo- and hyperpigmented macules, microcephaly and stunted growth as prominent features. European Journal of Medical Genetics, 2016, 59, 283-289.	1.3	29
63	A Probable Topiramate-induced Limbs Paraesthesia and Rigid Fingers Flexion. Current Drug Safety, 2018, 13, 131-136.	0.6	29
64	The usefulness of immunotherapy in pediatric neurodegenerative disorders: A systematic review of literature data. Human Vaccines and Immunotherapeutics, 2015, 11, 2749-2763.	3.3	28
65	Dynamic and Static Splinting for Treatment of Developmental Dysplasia of the Hip: A Systematic Review. Children, 2021, 8, 104.	1.5	28
66	SARS-CoV-2 related paediatric acute-onset neuropsychiatric syndrome. The Lancet Child and Adolescent Health, 2021, 5, e19-e21.	5.6	28
67	White matter changes mimicking a leukodystrophy in a patient with Mucopolysaccharidosis: characterization by MRI. Journal of the Neurological Sciences, 2002, 195, 171-175.	0.6	27
68	Clinical Heterogeneity in Familial Congenital Ptosis: Analysis of Fourteen Cases in One Family Over Five Generations. Pediatric Neurology, 2005, 33, 251-254.	2.1	27
69	Late Primary Unilateral Thalamic Hemorrhage in Infancy: Report of Two Cases. Neuropediatrics, 1999, 30, 264-267.	0.6	26
70	Neurological Manifestations in Individuals with Pure Cutaneous or Syndromic (Ruggieri-Happle) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 14	0.6	26
71	Electroclinical Features and Long-Term Outcome of Cryptogenic Epilepsy in Children with Down Syndrome. Journal of Pediatrics, 2013, 163, 1754-1758.	1.8	25
72	Expression of the OAS Gene Family Is Highly Modulated in Subjects Affected by Juvenile Dermatomyositis, Resembling an Immune Response to a dsRNA Virus Infection. International Journal of Molecular Sciences, 2018, 19, 2786.	4.1	25

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73	Gelastic seizures due to hypothalamic hamartoma: Rapid resolution after endoscopic tumor disconnection. Neurocase, 2013, 19, 458-461.	0.6	24
74	Cannabidiol Treatment for Refractory Epilepsies in Pediatrics. Frontiers in Pharmacology, 2020, 11, 586110.	3.5	23
75	Cyclic Vomiting Syndrome in Children. Frontiers in Neurology, 2020, 11, 583425.	2.4	23
76	Speech rehabilitation in dysarthria after stroke: a systematic review of the studies. European Journal of Physical and Rehabilitation Medicine, 2020, 56, 547-562.	2.2	23
77	Diabetes insipidus in neurobrucellosis. Clinical Neurology and Neurosurgery, 2000, 102, 163-165.	1.4	22
78	Microcephaly, sensorineural deafness and Currarino triad with duplicationâ€“deletion of distal 7q. European Journal of Pediatrics, 2010, 169, 475-481.	2.7	22
79	Pediatric status epilepticus: improved management with new drug therapies?. Expert Opinion on Pharmacotherapy, 2017, 18, 789-798.	1.8	22
80	Outbreak of COVID-19 infection in children: fear and serenity. European Review for Medical and Pharmacological Sciences, 2020, 24, 4572-4575.	0.7	22
81	Congenital idiopathic talipes equinovarus: an evaluation in infants treated by the Ponseti method. European Review for Medical and Pharmacological Sciences, 2013, 17, 2675-9.	0.7	22
82	Focal Neurological Deficits in Children with Î²-Thalassemia Major. Neuropediatrics, 1999, 30, 45-48.	0.6	21
83	Molecular Mechanism Involved in the Pathogenesis of Early-Onset Epileptic Encephalopathy. Frontiers in Molecular Neuroscience, 2019, 12, 118.	2.9	21
84	The role of micronutrients in the diet of HIV-1-infected individuals. Frontiers in Bioscience - Elite, 2012, E4, 2442-2456.	1.8	21
85	Hemihydranencephaly: Case Report and Literature Review. Journal of Child Neurology, 2001, 16, 218-221.	1.4	20
86	Congenital muscular dystrophy with adducted thumbs, ptosis, external ophthalmoplegia, mental retardation and cerebellar hypoplasia: a novel form of CMD. Neuromuscular Disorders, 2002, 12, 623-630.	0.6	20
87	Haemophilic arthropathy: A narrative review on the use of intraâ€articular drugs for arthritis. Haemophilia, 2019, 25, 919-927.	2.1	20
88	Short-Term Neurodevelopmental Outcome in Term Neonates Treated with Phenobarbital versus Levetiracetam: A Single-Center Experience. Behavioural Neurology, 2019, 2019, 1-8.	2.1	20
89	Oxidative Stress in Preterm Infants: Overview of Current Evidence and Future Prospects. Pharmaceuticals, 2020, 13, 145.	3.8	20
90	Cerebral Palsy and Epilepsy in Children: Clinical Perspectives on a Common Comorbidity. Children, 2021, 8, 16.	1.5	20

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91	Noninvasive prenatal diagnosis of chromosomal aneuploidies by isolation and analysis of fetal cells from maternal blood. American Journal of Medical Genetics Part A, 2001, 101, 262-267.	2.4	19
92	Acute hyperkinetic movement disorders in Italian paediatric emergency departments. Archives of Disease in Childhood, 2018, 103, 790-794.	1.9	19
93	Benign and severe early-life seizures: a round in the first year of life. Italian Journal of Pediatrics, 2018, 44, 54.	2.6	19
94	Congenital Insensitivity to Pain with Anhidrosis (NTRK1 Mutation) and Early Onset Renal Disease: Clinical Report on Three Sibs with a 25-Year Follow-Up in One of Them. Neuropediatrics, 2005, 36, 270-273.	0.6	18
95	Otologic findings in children with gastroesophageal reflux. International Journal of Pediatric Otorhinolaryngology, 2007, 71, 1693-1697.	1.0	18
96	The aristaless Arx gene one gene for many interneuronopathies. Frontiers in Bioscience - Elite, 2010, E2, 701-710.	1.8	18
97	Clinical Course of N-Methyl-D-Aspartate Receptor Encephalitis and the Effectiveness of Cyclophosphamide Treatment. Journal of Pediatric Neurology, 2017, 15, 084-089.	0.2	18
98	Interstitial 16p13.3 microduplication: Case report and critical review of genotypeâ€“phenotype correlation. European Journal of Medical Genetics, 2012, 55, 747-752.	1.3	17
99	Noninvasive ventilation for acute respiratory distress in children with central nervous system disorders. Respiratory Medicine, 2013, 107, 1370-1375.	2.9	17
100	Refractory absence seizures: An Italian multicenter retrospective study. European Journal of Paediatric Neurology, 2015, 19, 660-664.	1.6	17
101	Aripiprazole for the treatment of irritability and aggression in children and adolescents affected by autism spectrum disorders. Expert Review of Neurotherapeutics, 2016, 16, 867-874.	2.8	17
102	Pyridoxine Add-On Treatment for the Control of Behavioral Adverse Effects Induced by Levetiracetam in Children: A Case-Control Prospective Study. Annals of Pharmacotherapy, 2018, 52, 645-649.	1.9	17
103	COVID-19 Pandemic Outbreak and its Psychological Impact on Patients with Rare Lysosomal Diseases. Journal of Clinical Medicine, 2020, 9, 2716.	2.4	17
104	Recurrent Hopkin's syndrome: A case report and review of the literature. Journal of the Neurological Sciences, 2010, 297, 89-91.	0.6	16
105	Rotavirus and Celiac Disease. American Journal of Gastroenterology, 2007, 102, 1831-1831.	0.4	15
106	Epileptic seizures as a manifestation of cowâ€™s milk allergy: a studied relationship and description of our pediatric experience. Expert Review of Clinical Immunology, 2014, 10, 1597-1609.	3.0	15
107	Rapid Spontaneous Resolution of Fibromatosis Colli in a 3-Week-Old Girl. Case Reports in Otolaryngology, 2014, 2014, 1-4.	0.2	15
108	Ptosis in childhood. Medicine (United States), 2018, 97, e12124.	1.0	15

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109	Biological Drugs in Guillain-BarrÃ© Syndrome: An Update. Current Neuropharmacology, 2017, 15, 938-950.	2.9	15
110	Autoimmune Hepatitis Associated with Celiac Disease in Childhood: Report of Two Cases. Journal of Gastroenterology and Hepatology (Australia), 2003, 18, 1324-1327.	2.8	14
111	Peripheral Neuropathy as First Sign of Ulcerative Colitis in a Child. Journal of Clinical Gastroenterology, 2004, 38, 115-117.	2.2	14
112	Reflex myoclonic epilepsy in infancy: a critical review. Epileptic Disorders, 2013, 15, 114-122.	1.3	14
113	Clinical dissection of early onset absence epilepsy in children and prognostic implications. Epilepsia, 2013, 54, 1761-1770.	5.1	14
114	Liver transplantation in a child with celiac disease. Journal of Gastroenterology and Hepatology (Australia), 2005, 20, 956-960.	2.8	13
115	Anton-Babinski syndrome in a child with early-stage adrenoleukodystrophy. European Journal of Neurology, 2007, 14, e11-e12.	3.3	13
116	Callosal anomalies with interhemispheric cyst: Expanding the phenotype. Acta Paediatrica, International Journal of Paediatrics, 2007, 94, 1066-1072.	1.5	13
117	Clinical dissection of childhood occipital epilepsy of Gastaut and prognostic implication. European Journal of Neurology, 2016, 23, 241-246.	3.3	13
118	Acetaminophen administration in pediatric age: an observational prospective cross-sectional study. Italian Journal of Pediatrics, 2016, 42, 20.	2.6	13
119	PRRT2 gene variant in a child with dysmorphic features, congenital microcephaly, and severe epileptic seizures: genotype-phenotype correlation?. Italian Journal of Pediatrics, 2019, 45, 159.	2.6	13
120	Genotype-phenotype variable correlation in Wilson disease: clinical history of two sisters with the similar genotype. BMC Medical Genetics, 2020, 21, 128.	2.1	13
121	Treating the symptom or treating the disease in neonatal seizures: a systematic review of the literature. Italian Journal of Pediatrics, 2021, 47, 85.	2.6	13
122	Clinical heterogeneity in eyelid myoclonia, with absences, and epilepsy. European Journal of Pediatrics, 2002, 161, 175-177.	2.7	12
123	Klippelâ€Trenaunay syndrome in a boy with concomitant ipsilateral overgrowth and undergrowth. American Journal of Medical Genetics, Part A, 2014, 164, 1262-1267.	1.2	12
124	Microcephaly/Trigonocephaly, Intellectual Disability, Autism Spectrum Disorder, and Atypical Dysmorphic Features in a Boy with Xp22.31 Duplication. Molecular Syndromology, 2018, 9, 253-258.	0.8	12
125	Grisel Syndrome in Pediatric Age: A Single-Center Italian Experience and Review of the Literature. World Neurosurgery, 2019, 125, 374-382.	1.3	12
126	N-BiPAP vs n-CPAP in term neonate with respiratory distress syndrome. Early Human Development, 2020, 142, 104965.	1.8	12

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127	Fetal Cells in Maternal Blood: A Six-Fold Increase in Women who have Undergone Amniocentesis and Carry a Fetus with Down Syndrome: A Multicenter Study. Neuropediatrics, 2004, 35, 321-324.	0.6	11
128	Valproate in adolescents with photosensitive epilepsy with generalized tonicâ€“clonic seizures only. European Journal of Paediatric Neurology, 2014, 18, 13-18.	1.6	11
129	Sodium metabisulphite allergy with multiple food and drug hypersensitivities in a five-year-old child: A case report and literature review. Allergologia Et Immunopathologia, 2015, 43, 106-108.	1.7	11
130	Electroclinical findings and long-term outcomes in epileptic patients with inv dup (15). Acta Neurologica Scandinavica, 2018, 137, 575-581.	2.1	11
131	A New Patient with Potockiâ€“Lupski Syndrome: A Literature Review. Journal of Pediatric Genetics, 2018, 07, 029-034.	0.7	11
132	ENT involvement and orobuccal movements' disorders in Pandas patients: assessment and rehabilitations tools. European Review for Medical and Pharmacological Sciences, 2019, 23, 4110-4117.	0.7	11
133	Fetal Nucleated Red Blood Cell Counts in Peripheral Blood of Mothers Bearing Down Syndrome Fetus. Neuropediatrics, 2001, 32, 147-149.	0.6	10
134	A Complex Brain Malformation Syndrome with Rhombencephalosynapsis, Preaxial Hexadactyly plus Facial and Skull Anomalies. Neuropediatrics, 2005, 36, 279-283.	0.6	10
135	Neonatal onset of hot water reflex seizures in monozygotic twins subsequently manifesting episodes of alternating hemiplegia. Epilepsy Research, 2008, 78, 225-231.	1.6	10
136	Long-term outcome of epilepsy in Kabuki syndrome. Seizure: the Journal of the British Epilepsy Association, 2011, 20, 650-654.	2.0	10
137	Polysomnographic evaluation of nonâ€invasive ventilation in children with neuromuscular disease. Respirology, 2014, 19, 80-84.	2.3	10
138	Propranolol: Effectiveness and Failure in Infantile Cutaneous Hemangiomas. Drug Safety - Case Reports, 2015, 2, 6.	0.9	10
139	Incidence of Mediterranean Spotted Fever in Sicilian children: a clinical-epidemiological observational retrospective study from 1987 to 2010. International Journal of Infectious Diseases, 2015, 31, 35-40.	3.3	10
140	Therapeutic approaches to pediatric pseudotumor cerebri: New insights from literature data. International Journal of Immunopathology and Pharmacology, 2017, 30, 94-97.	2.1	10
141	Ictal blinking, an under-recognized phenomenon: our experience and literature review. Neuropsychiatric Disease and Treatment, 2017, Volume 13, 1435-1439.	2.2	10
142	Serum concentrations of perfluorinated compounds among children living in Sicily (Italy). Toxicology Letters, 2018, 298, 186-193.	0.8	10
143	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.	2.4	10
144	Liver infection and COVID-19: the electron microscopy proof and revision of the literature. European Review for Medical and Pharmacological Sciences, 2021, 25, 2146-2151.	0.7	10

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145	Headache, migraine and obesity: an overview on plausible links. Journal of Biological Regulators and Homeostatic Agents, 2016, 30, 333-8.	0.7	10
146	Tuberous breast deformity in an adolescent girl with Hurler-Scheie syndrome. European Journal of Pediatrics, 2000, 159, 936-937.	2.7	9
147	Symptomatic hypocalcemia in an epileptic child treated with valproic acid plus lamotrigine: a case report. Cases Journal, 2009, 2, 7394.	0.4	9
148	Apneic crises: A clue for MECP2 testing in severe neonatal hypotonia-respiratory failure. European Journal of Paediatric Neurology, 2012, 16, 744-748.	1.6	9
149	Clinical spectrum of woolly hair: indications for cerebral involvement. Italian Journal of Pediatrics, 2017, 43, 99.	2.6	9
150	Long-term follow-up and novel genotype-phenotype analysis of monozygotic twins with ATP1A3 mutation in Alternating Hemiplegia of Childhood-2. European Journal of Medical Genetics, 2020, 63, 103957.	1.3	9
151	Chilblainsâ€like lesions and SARSâ€CoV â€2 in children: An overview in therapeutic approach. Dermatologic Therapy, 2021, 34, e14502.	1.7	9
152	Intronic Variant in CNTNAP2 Gene in a Boy With Remarkable Conduct Disorder, Minor Facial Features, Mild Intellectual Disability, and Seizures. Frontiers in Pediatrics, 2020, 8, 550.	1.9	9
153	A nationwide study on Sydenham's chorea: Clinical features, treatment and prognostic factors. European Journal of Paediatric Neurology, 2022, 36, 1-6.	1.6	9
154	Tracheostomy in childhood: new causes for an old strategy. European Review for Medical and Pharmacological Sciences, 2012, 16, 1719-22.	0.7	9
155	Arthrogryposis Multiplex Congenita and Pituitary Ectopia. A Case Report. Neuropediatrics, 2000, 31, 325-327.	0.6	8
156	Neonatal onset of bath-induced alternating hemiplegia of childhood. European Journal of Paediatric Neurology, 2010, 14, 192-193.	1.6	8
157	Primary headache: Role of investigations in a cohort of young children and adolescents. Pediatrics International, 2011, 53, 964-967.	0.5	8
158	Reflex myoclonic epilepsy in infancy: A multicenter clinical study. Epilepsy Research, 2013, 103, 237-244.	1.6	8
159	Spinal Neurofibromatosis with Central Nervous System Involvement in a Set of Twin Girls and a Boy: Further Expansion of the Phenotype. Neuropediatrics, 2013, 44, 239-244.	0.6	8
160	Noninvasive ventilation in pediatric emergency care: a literature review and description of our experience. Expert Review of Respiratory Medicine, 2013, 7, 545-552.	2.5	8
161	Vitamin D3 an ever green molecule. Frontiers in Bioscience - Scholar, 2013, S5, 247-260.	2.1	8
162	Optic Neuropathy, Secondary to Ethmoiditis, and Onodi Cell Inflammation during Childhood: A Case Report and Review of the Literature. Neuropediatrics, 2019, 50, 341-345.	0.6	8

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163	A Systematic Review of Pharmacologic and Rehabilitative Treatment of Small Fiber Neuropathies. Diagnostics, 2020, 10, 1022.	2.6	8
164	A novel GABRB3 variant in Dravet syndrome: Case report and literature review. Molecular Genetics & Genomic Medicine, 2020, 8, e1461.	1.2	8
165	Arthroereisis in juvenile flexible flatfoot: Which device should we implant? A systematic review of literature published in the last 5 years. World Journal of Orthopedics, 2021, 12, 433-444.	1.8	8
166	Efficacy and safety of omalizumab in paediatric age: an update of literature data. Journal of Biological Regulators and Homeostatic Agents, 2016, 30, 579-84.	0.7	8
167	Callosal anomalies with interhemispheric cyst: Expanding the phenotype. Acta Paediatrica, International Journal of Paediatrics, 2005, 94, 1066-1072.	1.5	7
168	Lissencephalic syndromes brain and beyond. Frontiers in Bioscience - Scholar, 2010, S2, 85-95.	2.1	7
169	An 11-Year Follow-up Study of Neonatal-Onset, Bath-Induced Alternating Hemiplegia of Childhood in Twins. Journal of Child Neurology, 2012, 27, 657-662.	1.4	7
170	Generalised epilepsy with febrile seizures plus (GEFS+): molecular analysis in a restricted area. Child's Nervous System, 2012, 28, 141-145.	1.1	7
171	Cerebral White Matter Lesions and Dysmorphisms: Signs Suggestive of 6p25 Deletion Syndromeâ€"Literature Review. Journal of Pediatric Genetics, 2019, 08, 205-211.	0.7	7
172	Griselâ€™s syndrome caused by Mycoplasma pneumoniae infection: a case report and review of the literature. Child's Nervous System, 2019, 35, 523-527.	1.1	7
173	Vitamin B12 Deficiency and West Syndrome: An Uncommon but Preventable Cause of Neurological Disorder. Report on Three Cases, One of Them with Late Onset during Vitamin B12 Treatment. Neuropediatrics, 2021, 52, 333-336.	0.6	7
174	The Evolution of the Role of Imaging in the Diagnosis of Craniosynostosis: A Narrative Review. Children, 2021, 8, 727.	1.5	7
175	Congenital myasthenic syndromes: Clinical and molecular report on 7 Sicilian patients. Journal of Pediatric Neurosciences, 2013, 8, 19.	0.3	7
176	Fever-Associated Seizures or Epilepsy: An Overview of Old and Recent Literature Acquisitions. Frontiers in Pediatrics, 2022, 10, 858945.	1.9	7
177	Two siblings with a homozygous MTHFR C677T (G80A-RFC1) mutation and stroke. Child's Nervous System, 2009, 25, 361-365.	1.1	6
178	Sublingual immunotherapy in preschool children: an update. Expert Review of Clinical Immunology, 2013, 9, 385-390.	3.0	6
179	Autonomic Dysfunction Manifesting With Asymmetric Face Flushing and Paroxysmal Nonconvulsive Episodes. Journal of Child Neurology, 2013, 28, 1673-1676.	1.4	6
180	Nevus Sebaceous and Its Association With Neurologic Involvement. Seminars in Pediatric Neurology, 2015, 22, 302-309.	2.0	6

#	Article	IF	Citations
181	Resuming the obsolete term â€œsmall headâ€ when microcephaly occurs without cognitive impairment. Neurological Sciences, 2017, 38, 1723-1725.	1.9	6
182	Diagnosis, Rehabilitation and Preventive Strategies for Pudendal Neuropathy in Cyclists, A Systematic Review. Journal of Functional Morphology and Kinesiology, 2021, 6, 42.	2.4	6
183	The Gutâ€“brain axis: A new pathogenic view of neurologic symptoms â€“ Description of a pediatric case. Journal of Pediatric Neurosciences, 2017, 12, 105.	0.3	6
184	Obsessive-Compulsive Disorder in PANS/PANDAS in Children: In Search of a Qualified Treatmentâ€”A Systematic Review and Metanalysis. Children, 2022, 9, 155.	1.5	6
185	Diffuse Polymicrogyria Associated With Congenital Hydrocephalus, Craniosynostosis, Severe Mental Retardation, and Minor Facial and Genital Anomalies. Journal of Child Neurology, 2000, 15, 493-495.	1.4	5
186	Holoprosencephaly (Lobar Form) Associated With Bilateral Vocal Cord Palsy. Journal of Child Neurology, 2001, 16, 932-934.	1.4	5
187	CNS findings in three cases of septo-optic dysplasia, including one with semilobar holoprosencephaly. American Journal of Medical Genetics, Part A, 2005, 136A, 357-357.	1.2	5
188	Natalizumab in multiple sclerosis: discontinuation, progressive multifocal leukoencephalopathy and possible use in children. Expert Review of Neurotherapeutics, 2015, 15, 1321-1341.	2.8	5
189	Legumes steam allergy in childhood: Update of the reported cases. Allergologia Et Immunopathologia, 2015, 43, 196-202.	1.7	5
190	Cervical neurenteric cyst and Klippel-Feil syndrome: An abrupt onset of myelopathic signs in a young patient. Journal of Pediatric Surgery Case Reports, 2017, 24, 12-16.	0.2	5
191	Chromosome 2p15-p16.1 microduplication in a boy with congenital anomalies: Is it a distinctive syndrome?. European Journal of Medical Genetics, 2019, 62, 47-54.	1.3	5
192	Severe Psychotic Symptoms in Youth with PANS/PANDAS: Case-Series. Journal of Child and Adolescent Psychopharmacology, 2020, 30, 567-571.	1.3	5
193	Electroclinical pattern and epilepsy evolution in an infant with Millerâ€“Dieker syndrome. Journal of Pediatric Neurosciences, 2018, 13, 302.	0.3	5
194	A measles outbreak in Catania, Sicily: the importance of high vaccination coverage and early notification of cases for health and economic reasons. Infezioni in Medicina, 2014, 22, 222-6.	1.1	5
195	Autoimmune encephalitis and CSF anti-AMPA GluR3 antibodies in childhood: a case report and literature review. Neurological Sciences, 2022, 43, 5237-5241.	1.9	5
196	Extraordinary intrathecal bone reaction in Î²-thalassaemia intermedia. Lancet, The, 1999, 354, 922.	13.7	4
197	Long-Term Survival in a Child With Arthrogryposis Multiplex Congenita and Spinal Muscular Atrophy. Journal of Child Neurology, 2001, 16, 934-936.	1.4	4
198	Blink reflex abnormalities in children with Tourette syndrome. European Journal of Neurology, 2006, 13, 869-873.	3.3	4

#	Article	IF	Citations
199	Polyneuritis Cranialis: Full Recovery after Intravenous Immunoglobulins. Pediatric Neurology, 2007, 37, 209-211.	2.1	4
200	Incidence and causes of neonatal hyperbilirubinemia in a center of Catania. Therapeutics and Clinical Risk Management, 2009, 5, 247.	2.0	4
201	Leiomyosarcoma of the Parotid Gland in an HIV-Positive Patient: Therapeutic Approach, Clinical Course and Review of the Literature. Journal of Chemotherapy, 2009, 21, 215-218.	1.5	4
202	First case of dizygous twins with Xâ€linked Î±â€thalassemia/mental retardation syndrome showing wide clinical variability. Pediatrics International, 2010, 52, e72-5.	0.5	4
203	Spine and brain malformations in a patient obligate carrier of MTHFR with autism and mental retardation. Clinical Neurology and Neurosurgery, 2012, 114, 1280-1282.	1.4	4
204	7q31.32 partial duplication: First report of a child with dysmorphism, autistic spectrum disorder, moderate intellectual disability and, epilepsy. Literature review. Epilepsy Research, 2019, 158, 106223.	1.6	4
205	Congenital Genetic Microcephaly: Clinical Diagnostic Approach. Journal of Pediatric Neurology, 2020, 18, 131-134.	0.2	4
206	Perinatal Femoral Fracture: A Ten-Year Observational Case Series Study. Children, 2020, 7, 156.	1.5	4
207	Deciphering the Invdupdel(8p) Genotypeâ€“Phenotype Correlation: Our Opinion. Brain Sciences, 2020, 10, 451.	2.3	4
208	Gray matter heterotopia: clinical and neuroimaging report on 22 children. Acta Neurologica Belgica, 2022, 122, 153-162.	1.1	4
209	Nervous system involvement in clinical peripheral inflammation: A description of three pediatric cases. Journal of Pediatric Neurosciences, 2016, 11, 277.	0.3	4
210	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
211	Otorrhea in Kawasaki disease diagnosis complicated by an EBV infection: coincidental disease or a true association. European Review for Medical and Pharmacological Sciences, 2013, 17, 989-93.	0.7	4
212	Bickerstaff's brainstem encephalitis (BBE) in childhood: rapid resolution after intravenous immunoglobulins treatment. European Review for Medical and Pharmacological Sciences, 2014, 18, 2496-9.	0.7	4
213	Preventive strategies, exercises and rehabilitation of hand neuropathy in cyclists: A systematic review. Journal of Hand Therapy, 2022, 35, 164-173.	1.5	4
214	Vein of Galen Malformation and Infantile Spasms. Journal of Child Neurology, 1999, 14, 196-198.	1.4	3
215	Colobomatous microphthalmia, microcephaly with cerebellar hypoplasia: Association or new syndrome?. , 2000, 92, 278-280.		3
216	Congenital Lymphedema-lymphangiectasia Associated With Scrotal Angiokeratoma (Fordyce Type) and Hearing Impairment. Journal of Clinical Gastroenterology, 2008, 42, 715-719.	2.2	3

#	Article	IF	Citations
217	Ganglioneuroblastoma-associated Vitamin D Deficiency Rickets. Journal of Pediatric Hematology/Oncology, 2009, 31, 502-504.	0.6	3
218	Recurrent obstructive hydrocephalus in a 4-month-old infant. Child's Nervous System, 2010, 26, 133-136.	1.1	3
219	Intravenous Immunoglobulin Therapy When Plasmapheresis Fails in Thrombotic Thrombocytopenic Purpura Associated with Severe ADAMTS 13 Deficiency in Childhood: A Case Report. European Journal of Inflammation, 2013, 11, 291-296.	0.5	3
220	Usefulness of video-EEG in the paediatric emergency department. Expert Review of Neurotherapeutics, 2014, 14, 769-785.	2.8	3
221	Diagnostic Clue in a Neonate with Amniotic Band Sequence. Case Reports in Pediatrics, 2020, 2020, 1-5.	0.4	3
222	Primary Microcephaly with Novel Variant of MCPH1 Gene in Twins: Both Manifesting in Childhood at the Same Time with Hashimoto's Thyroiditis. Journal of Pediatric Genetics, 2020, 09, 177-182.	0.7	3
223	PANS/PANDAS: Clinical Experience in IVIG Treatment and State of the Art in Rehabilitation Approaches. NeuroSci, 2020, 1, 75-84.	1.2	3
224	Poland Syndrome with Atypical Malformations Associated to a de novo 1.5â€‰Mb Xp22.31 Duplication. Neuropediatrics, 2020, 51, 359-363.	0.6	3
225	Aarskog-Scott syndrome: clinical and molecular characterisation of a family with the coexistence of a novel FGD1 mutation and 16p13.11-p12.3 microduplication. BMJ Case Reports, 2020, 13, e235183.	0.5	3
226	Chromosome 15q BP3 to BP5 deletion is a likely locus for speech delay and language impairment: Report on a fourâ€member family and an unrelated boy. Molecular Genetics & Genomic Medicine, 2020, 8, e1109.	1.2	3
227	Ocular Motor Paroxysmal Events in Neonates and Infants: A Review of the Literature. Pediatric Neurology, 2021, 117, 4-9.	2.1	3
228	Single and in combination antiepileptic drug therapy in children with epilepsy: how to use it. AIMS Medical Science, 2021, 8, 138-146.	0.4	3
229	Neurotropism of SARS-CoV 2 and others Coronavirus in Children: Mechanisms and Clinical Manifestations. Eurasian Journal of Medicine and Oncology, 2021, , .	1.0	3
230	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.	0.3	3
231	Idiopathic intracranial hypertension in a paediatric population: a retrospective observational study on epidemiology, symptoms and treatment. Journal of Biological Regulators and Homeostatic Agents, 2017, 31, 195-200.	0.7	3
232	Benign Acute Childhood Myositis: Our Experience on Clinical Evaluation. Neuropediatrics, 2022, 53, 418-422.	0.6	3
233	Primary headache in childhood associated with psychiatric disturbances: an update. European Review for Medical and Pharmacological Sciences, 2020, 24, 6893-6898.	0.7	3
234	Bickerstaff's brainstem encephalitis in childhood: a literature overview. European Review for Medical and Pharmacological Sciences, 2020, 24, 12802-12807.	0.7	3

#	Article	IF	Citations
235	Lobar Holoprosencephaly. Journal of Child Neurology, 2002, 17, 543-544.	1.4	2
236	Microcephaly associated with Legg-CalvÃ¨-Perthes disease in two siblings. Neurological Sciences, 2012, 33, 1401-1405.	1.9	2
237	Acute Glomerulonephritis in a Child withChlamydia pneumoniaeInfection: A Case Report. Case Reports in Medicine, 2013, 2013, 1-4.	0.7	2
238	Congenital familial myasthenic syndromes: disease and course in an affected dizygotic twin pair. BMJ Case Reports, 2013, 2013, bcr2012007651-bcr2012007651.	0.5	2
239	Down syndrome and arterial ischemic stroke in childhood: A potential immunologic link with selective IgG4 subclass deficiency. European Journal of Paediatric Neurology, 2014, 18, 520-525.	1.6	2
240	Three Cases of Palatal Tics and Gilles De La Tourette Syndrome. Journal of Child Neurology, 2015, 30, 1199-1203.	1.4	2
241	A girl with a 14.7â€‰Mb 3q26.32â€“q28 duplication: a new report of 3q duplication syndrome and a literature review. Clinical Dysmorphology, 2016, 25, 121-127.	0.3	2
242	Giant Melanocytic Nevi and Soft Tissue Undergrowth in the Left Leg: Pathogenetic Hypothesis. Mental Illness, 2019, 11, 8184.	0.8	2
243	Clinicoradiographic data and management of children with Chiari malformation type 1 and 1.5: an Italian case series. Acta Neurologica Belgica, 2021, 121, 1547-1554.	1.1	2
244	Advanced olfactory neuroblastoma in a teenager: a clinical case and short review of literature. Child's Nervous System, 2020, 36, 485-489.	1.1	2
245	On clinical findings of Bickerstaffâ€™s brainstem encephalitis in childhood. Journal of Integrative Neuroscience, 2021, 20, 509.	1.7	2
246	Non-Epileptic Paroxysmal Events: Clinical features and diagnostic differences with epileptic seizures. A Single Tertiary Centre Study. Clinical Neurology and Neurosurgery, 2021, 207, 106739.	1.4	2
247	Chromosome 15q BP4-BP5 Deletion in a Girl with Nocturnal Frontal Lobe Epilepsy, Migraine, Circumscribed Hypertrichosis, and Language Impairment. Journal of Epilepsy Research, 2020, 10, 84-91.	0.4	2
248	A Selective Mutism Arising from First Language Attrition, Successfully Treated with Paroxetine-CBT Combination Treatment. Psychiatry Investigation, 2015, 12, 569.	1.6	2
249	Blepharoptosis in children: our experience at the light of literature. Clinica Terapeutica, 2010, 161, 241-3.	0.3	2
250	Alternating Hemiplegia of Childhood: neurological comorbidities and intrafamilial variability. Italian Journal of Pediatrics, 2022, 48, 29.	2.6	2
251	Neonatal ischemic limb lesions: From etiology to topical nitroglycerine. A case series analysis. Dermatologic Therapy, 2022, 35, e15426.	1.7	2
252	Thymosin beta-4 prenatal administration improves fetal development and halts side effects due to preterm delivery. European Review for Medical and Pharmacological Sciences, 2021, 25, 431-437.	0.7	2

#	Article	IF	Citations
253	Pseudohypoparathyroidism Ia With Evans Syndrome. Journal of Pediatric Hematology/Oncology, 2008, 30, 628-630.	0.6	1
254	A Syndrome with Coarse Face, Mental Retardation and Unusual Stereotyped Movements. Neuropediatrics, 2009, 40, 186-188.	0.6	1
255	Longâ€term neurological outcome of a sextuplet pregnancy. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 777-779.	1.5	1
256	Cutaneous and Leptomeningeal Hemangiomas With Impressive Benign Evolution. Pediatric Neurology, 2013, 48, 73-75.	2.1	1
257	Single-Epileptic Spasms with or without Hypsarrhythmia: A Study of 16 Patients. Journal of Pediatric Epilepsy, 2017, 06, 149-155.	0.2	1
258	Focal Neuropathy Mimicking Focal Dystonia in a Child: Diagnostic and Rehabilitative Tools. Journal of Functional Morphology and Kinesiology, 2019, 4, 54.	2.4	1
259	Previously Unreported <i>COL7A1</i> Mutation in a Somali Patient with Dystrophic Epidermolysis Bullosa. Molecular Syndromology, 2019, 10, 332-338.	0.8	1
260	Arterial ischemic stroke (AIS) in childhood: clinical report from a single control center. Child's Nervous System, 2019, 35, 283-293.	1.1	1
261	Malformations of Cortical Development, Cognitive Involvementand Epilepsy: A Single Institution Experience in 19 Young Patients. Children, 2021, 8, 637.	1.5	1
262	Pulse Oximetry Saturation (Spo 2) Monitoring in the Neonatal Intensive Care Unit (NICU): The Challenge for Providers. Advances in Neonatal Care, 2022, 22, 231-238.	1.1	1
263	Anti-MOG Antibody Syndrome and Cerebral Sinovenous Thrombosis: A Causeâ€"Effect Hypothesis. Journal of Pediatric Neurology, 2021, 19, 127-131.	0.2	1
264	Child Abuse Syndrome (CAS): A Newly Recognized Distinct Entity. The Open Neurology Journal, 2016, 10, 30-31.	0.4	1
265	Mild Hypoxicâ€"Ischemic Encephalopathy: Can Neurophysiological Monitoring Predict Unfavorable Neurological Outcome? A Systematic Review and Meta-analysis. American Journal of Perinatology, 2021, , .	1.4	1
266	Neurodevelopmental outcomes of neonatal nonâ€epileptic paroxysmal events: a prospective study. Developmental Medicine and Child Neurology, 2021, 63, 343-348.	2.1	1
267	A Novel 4q32.3 Deletion in a Boy: Additional Signs and the Role of MARCH1. Journal of Pediatric Genetics, 2021, 10, 259-265.	0.7	1
268	Acute asthma in children: treatment in emergency. European Review for Medical and Pharmacological Sciences, 2011, 15, 711-6.	0.7	1
269	Comment on safety and efficacy of oral lopinavir/ritonavir in pediatric patients with coronavirus disease: a nationwide comparative analysis. European Review for Medical and Pharmacological Sciences, 2021, 25, 2473-2474.	0.7	1
270	Reply to Dr Hahn. Journal of Child Neurology, 2003, 18, 152-152.	1.4	0

#	Article	IF	Citations
271	Chronic Yersinia Enterocolitis Infection Presenting as Intestinal Obstruction. Infectious Diseases in Clinical Practice, 2007, 15, 68-69.	0.3	0
272	Voluntary selective big toe dorsal flection: pseudo-Babinski phenomenon?. Neurological Sciences, 2008, 29, 495-496.	1.9	0
273	Chediak-Steinbrinck-Higashi Syndrome. , 2009, , 314-314.		0
274	A boy born with multiple lesions of atrophoderma. Acta Paediatrica, International Journal of Paediatrics, 2010, 99, 460-463.	1.5	0
275	Neonatal Autoimmune Hypothyroidism: A Patient Report. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 957-61.	0.9	0
276	Erratum to â€œRecurrent Hopkin's syndrome: A case report and review of the literatureâ€•[J Neurol Sci 297 (2010) 89â€“91]. Journal of the Neurological Sciences, 2011, 303, 150.	0.6	0
277	Non-Epileptik Psikojen NÃ¶betler: Klinik Pratikte Ã‡Ã¶zÃ¼lmesi Zor TanÄ±sal Bir Problem. Noropsikiyatri Arsivi, 2012, 49, 243-247.	0.7	0
278	Peripheral neuropathy in a child with Mycoplasma pneumoniae infections. Journal of Pediatric Infectious Diseases, 2015, 02, 163-166.	0.2	0
279	Asthma and hypoxia. Italian Journal of Pediatrics, 2015, 41, .	2.6	0
280	Seizures Are Not a Prevalent Feature of Wyburn-Mason Syndrome. Journal of Pediatric Epilepsy, 2016, 05, 111-114.	0.2	0
281	Holoprosencephaly and Recurrent Episodes of Severe Neurogenic Hypernatremia. Journal of Pediatric Neurology, 2016, 14, 025-030.	0.2	0
282	Array-CGH in pediatric neurology: A prospective observational study. European Journal of Inflammation, 2016, 14, 196-199.	0.5	0
283	The Role of Dendritic Cells in Central Nervous System Autoimmunity: Focusing on Multiple Sclerosis and Emerging Therapeutics Targeting Dendritic Cells. Journal of Pediatric Biochemistry, 2017, 06, 121-135.	0.2	0
284	Adaptive psychological structure in childhood hearing impairment: audiological correlations. Acta Otorhinolaryngologica Italica, 2017, 37, 175-179.	1.5	0
285	P137 Abdominal and neurologic pain in Harderoporphyria: a case report. Digestive and Liver Disease, 2018, 50, e405-e406.	0.9	0
286	Neuroaspergillosis as the Presenting Sign of a Chronic Granulomatous Disease. Pediatric Neurology, 2020, 102, 79-80.	2.1	0
287	Acute Motor Axonal Neuropathy in a 5-Month-Old Child. Journal of Pediatric Neurology, 2020, 18, 171-174.	0.2	0
288	Aicardiâ€“GoutiÃ¨res Syndrome Type 2: A Report on Two Cases with Different Phenotypes Caused by RNASEH2B Gene Mutations. Journal of Pediatric Neurology, 2020, 18, 206-209.	0.2	0

#	Article	IF	Citations
289	Secondary Scoliosis as a Complication of Acute Transverse Myelitis in a Child. Journal of Functional Morphology and Kinesiology, 2020, 5, 39.	2.4	0
290	Pure Interstitial 7q21.3-q 31.1 Duplication: A Rare Segmental Genomic Aneuploidy: Case Report and Review of Cases with Distal and Similar Segment Involved. Global Medical Genetics, 2021, 08, 123-128.	0.9	0
291	Atypical Presentation of Herpes Simplex Virus 1 Encephalitis in Pediatric Age. Open Access Macedonian Journal of Medical Sciences, 2021, 9, 204-208.	0.2	0
292	Hemihydranencephaly: Case Report and Literature Review. Journal of Child Neurology, 2001, 16, 212.	1.4	0
293	A mild phenotype associated with a de novo microdeletion 10q23.1-q23.2: a new patient with a novel feature. BMJ Case Reports, 2016, 2016, bcr2016214388.	0.5	0
294	Non-syndromic megalencephaly and epilepsy: Our findings. Brain and Nerves, 2020, 5, .	0.3	0
295	Celiac disease and headache in children: a narrative state of the art. Acta Biomedica, 2020, 91, e2020056.	0.3	0
296	Colobomatous microphthalmia, microcephaly with cerebellar hypoplasia: association or new syndrome?. American Journal of Medical Genetics Part A, 2000, 92, 278-80.	2.4	0
297	Parasympathetic nervous system involvement in food allergy: description of a paediatric case. Journal of Biological Regulators and Homeostatic Agents, 2016, 30, 1137-1140.	0.7	0
298	Congenital muscular dystrophy and epilepsy: a prospective case series of pediatric patients. Journal of Biological Regulators and Homeostatic Agents, 2016, 30, 1217-1221.	0.7	0
299	Peripheral Neuropathies: State of the Art on Modern Use of Electrodiagnostic Examination in Guillain-BarrÃ¨ Syndrome and Rehabilitative Implications in Children. Acta Scientific Paediatrics, 0, , 03-08.	0.1	0
300	Concomitant MPZ and MFN2 Gene Variants and Charcot Marie Tooth Disease in a Boy: Clinical and Genetic Analysisâ€"Literature Review. Case Reports in Pediatrics, 2022, 2022, 1-6.	0.4	0