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	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
2	Gray matter heterotopia: clinical and neuroimaging report on 22 children. Acta Neurologica Belgica, 2022, 122, 153-162.	1.1	4
3	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
4	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
5	Alternating Hemiplegia of Childhood: neurological comorbidities and intrafamilial variability. Italian Journal of Pediatrics, 2022, 48, 29.	2.6	2
6	Benign Acute Childhood Myositis: Our Experience on Clinical Evaluation. Neuropediatrics, 2022, 53, 418-422.	0.6	3
7	Neonatal ischemic limb lesions: From etiology to topical nitroglycerine. A case series analysis. Dermatologic Therapy, 2022, 35, e15426.	1.7	2
8	Preventive strategies, exercises and rehabilitation of hand neuropathy in cyclists: A systematic review. Journal of Hand Therapy, 2022, 35, 164-173.	1.5	4
9	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
10	Fever-Associated Seizures or Epilepsy: An Overview of Old and Recent Literature Acquisitions. Frontiers in Pediatrics, 2022, 10, 858945.	1.9	7
11	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
12	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
13	Silent COVID-19: what your skin can reveal. Lancet Infectious Diseases, The, 2021, 21, 24-25.	9.1	56
14	Ocular Motor Paroxysmal Events in Neonates and Infants: A Review of the Literature. Pediatric Neurology, 2021, 117, 4-9.	2.1	3
15	Chilblainsâ€like lesions and SARSâ€CoV â€2 in children: An overview in therapeutic approach. Dermatologic Therapy, 2021, 34, e14502.	1.7	9
16	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
17	On clinical findings of Bickerstaffâ€™s brainstem encephalitis in childhood. Journal of Integrative Neuroscience, 2021, 20, 509.	1.7	2
18	Neurotropism of SARS-CoV 2 and others Coronavirus in Children: Mechanisms and Clinical Manifestations. Eurasian Journal of Medicine and Oncology, 2021, , .	1.0	3

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19	Dynamic and Static Splinting for Treatment of Developmental Dysplasia of the Hip: A Systematic Review. Children, 2021, 8, 104.	1.5	28
20	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
21	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
22	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
23	SARS-CoV-2 related paediatric acute-onset neuropsychiatric syndrome. The Lancet Child and Adolescent Health, 2021, 5, e19-e21.	5.6	28
24	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
25	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
26	Malformations of Cortical Development, Cognitive Involvementand Epilepsy: A Single Institution Experience in 19 Young Patients. Children, 2021, 8, 637.	1.5	1
27	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
28	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
29	The Evolution of the Role of Imaging in the Diagnosis of Craniosynostosis: A Narrative Review. Children, 2021, 8, 727.	1.5	7
30	Anti-MOG Antibody Syndrome and Cerebral Sinovenous Thrombosis: A Causeâ€“Effect Hypothesis. Journal of Pediatric Neurology, 2021, 19, 127-131.	0.2	1
31	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
32	Cerebral Palsy and Epilepsy in Children: Clinical Perspectives on a Common Comorbidity. Children, 2021, 8, 16.	1.5	20
33	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
34	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
35	Neurodevelopmental outcomes of neonatal nonâ€epileptic paroxysmal events: a prospective study. Developmental Medicine and Child Neurology, 2021, 63, 343-348.	2.1	1
36	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

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37	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
38	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
39	Neuroaspergillosis as the Presenting Sign of a Chronic Granulomatous Disease. Pediatric Neurology, 2020, 102, 79-80.	2.1	0
40	Congenital Genetic Microcephaly: Clinical Diagnostic Approach. Journal of Pediatric Neurology, 2020, 18, 131-134.	0.2	4
41	Acute Motor Axonal Neuropathy in a 5-Month-Old Child. Journal of Pediatric Neurology, 2020, 18, 171-174.	0.2	0
42	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
43	Perinatal Femoral Fracture: A Ten-Year Observational Case Series Study. Children, 2020, 7, 156.	1.5	4
44	Ketogenic diet for infants with epilepsy: A literature review. Epilepsy and Behavior, 2020, 112, 107361.	1.7	46
45	Severe Psychotic Symptoms in Youth with PANS/PANDAS: Case-Series. Journal of Child and Adolescent Psychopharmacology, 2020, 30, 567-571.	1.3	5
46	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
47	Diagnostic Clue in a Neonate with Amniotic Band Sequence. Case Reports in Pediatrics, 2020, 2020, 1-5.	0.4	3
48	A Systematic Review of Pharmacologic and Rehabilitative Treatment of Small Fiber Neuropathies. Diagnostics, 2020, 10, 1022.	2.6	8
49	Deciphering the Invdupdel(8p) Genotypeâ€“Phenotype Correlation: Our Opinion. Brain Sciences, 2020, 10, 451.	2.3	4
50	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
51	COVID-19 Pandemic Outbreak and its Psychological Impact on Patients with Rare Lysosomal Diseases. Journal of Clinical Medicine, 2020, 9, 2716.	2.4	17
52	A novel GABRB3 variant in Dravet syndrome: Case report and literature review. Molecular Genetics & Genomic Medicine, 2020, 8, e1461.	1.2	8
53	West syndrome: a comprehensive review. Neurological Sciences, 2020, 41, 3547-3562.	1.9	73
54	Cannabidiol Treatment for Refractory Epilepsies in Pediatrics. Frontiers in Pharmacology, 2020, 11, 586110.	3.5	23

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55	Cyclic Vomiting Syndrome in Children. Frontiers in Neurology, 2020, 11, 583425.	2.4	23
56	PANS/PANDAS: Clinical Experience in IVIG Treatment and State of the Art in Rehabilitation Approaches. NeuroSci, 2020, 1, 75-84.	1.2	3
57	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
58	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
59	Secondary Scoliosis as a Complication of Acute Transverse Myelitis in a Child. Journal of Functional Morphology and Kinesiology, 2020, 5, 39.	2.4	0
60	Oxidative Stress in Preterm Infants: Overview of Current Evidence and Future Prospects. Pharmaceuticals, 2020, 13, 145.	3.8	20
61	Poland Syndrome with Atypical Malformations Associated to a de novo 1.5â€‰Mb Xp22.31 Duplication. Neuropediatrics, 2020, 51, 359-363.	0.6	3
62	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
63	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
64	N-BiPAP vs n-CPAP in term neonate with respiratory distress syndrome. Early Human Development, 2020, 142, 104965.	1.8	12
65	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
66	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
67	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
68	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
69	Outbreak of COVID-19 infection in children: fear and serenity. European Review for Medical and Pharmacological Sciences, 2020, 24, 4572-4575.	0.7	22
70	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
71	Non-syndromic megalencephaly and epilepsy: Our findings. Brain and Nerves, 2020, 5, .	0.3	0
72	Celiac disease and headache in children: a narrative state of the art. Acta Biomedica, 2020, 91, e2020056.	0.3	0

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73	Primary headache in childhood associated with psychiatric disturbances: an update. European Review for Medical and Pharmacological Sciences, 2020, 24, 6893-6898.	0.7	3
74	Bickerstaff's brainstem encephalitis in childhood: a literature overview. European Review for Medical and Pharmacological Sciences, 2020, 24, 12802-12807.	0.7	3
75	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
76	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
77	Focal Neuropathy Mimicking Focal Dystonia in a Child: Diagnostic and Rehabilitative Tools. Journal of Functional Morphology and Kinesiology, 2019, 4, 54.	2.4	1
78	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
79	Haemophilic arthropathy: A narrative review on the use of intraâ€articular drugs for arthritis. Haemophilia, 2019, 25, 919-927.	2.1	20
80	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
81	Giant Melanocytic Nevi and Soft Tissue Undergrowth in the Left Leg: Pathogenetic Hypothesis. Mental Illness, 2019, 11, 8184.	0.8	2
82	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
83	Molecular Mechanism Involved in the Pathogenesis of Early-Onset Epileptic Encephalopathy. Frontiers in Molecular Neuroscience, 2019, 12, 118.	2.9	21
84	Grisel Syndrome in Pediatric Age: A Single-Center Italian Experience and Review of the Literature. World Neurosurgery, 2019, 125, 374-382.	1.3	12
85	Previously Unreported <i>COL7A1</i> Mutation in a Somali Patient with Dystrophic Epidermolysis Bullosa. Molecular Syndromology, 2019, 10, 332-338.	0.8	1
86	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
87	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
88	Arterial ischemic stroke (AIS) in childhood: clinical report from a single control center. Child's Nervous System, 2019, 35, 283-293.	1.1	1
89	Epilepsy in Children: From Diagnosis to Treatment with Focus on Emergency. Journal of Clinical Medicine, 2019, 8, 39.	2.4	40
90	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

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91	Acute hyperkinetic movement disorders in Italian paediatric emergency departments. Archives of Disease in Childhood, 2018, 103, 790-794.	1.9	19
92	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
93	Electroclinical findings and long-term outcomes in epileptic patients with inv dup (15). Acta Neurologica Scandinavica, 2018, 137, 575-581.	2.1	11
94	A New Patient with Potockiâ€“Lupski Syndrome: A Literature Review. Journal of Pediatric Genetics, 2018, 07, 029-034.	0.7	11
95	Ptosis in childhood. Medicine (United States), 2018, 97, e12124.	1.0	15
96	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
97	P137 Abdominal and neurologic pain in Harderoporphyria: a case report. Digestive and Liver Disease, 2018, 50, e405-e406.	0.9	0
98	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
99	Serum concentrations of perfluorinated compounds among children living in Sicily (Italy). Toxicology Letters, 2018, 298, 186-193.	0.8	10
100	Benign and severe early-life seizures: a round in the first year of life. Italian Journal of Pediatrics, 2018, 44, 54.	2.6	19
101	A Probable Topiramate-induced Limbs Paraesthesia and Rigid Fingers Flexion. Current Drug Safety, 2018, 13, 131-136.	0.6	29
102	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
103	Electroclinical pattern and epilepsy evolution in an infant with Millerâ€“Dieker syndrome. Journal of Pediatric Neurosciences, 2018, 13, 302.	0.3	5
104	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
105	Ataxia in children: early recognition and clinical evaluation. Italian Journal of Pediatrics, 2017, 43, 6.	2.6	59
106	Randomized Comparison of Helmet CPAP Versus High-Flow Nasal Cannula Oxygen in Pediatric Respiratory Distress. Respiratory Care, 2017, 62, 1036-1042.	1.6	35
107	Pediatric status epilepticus: improved management with new drug therapies?. Expert Opinion on Pharmacotherapy, 2017, 18, 789-798.	1.8	22
108	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

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109	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
110	Therapeutic approaches to pediatric pseudotumor cerebri: New insights from literature data. International Journal of Immunopathology and Pharmacology, 2017, 30, 94-97.	2.1	10
111	Resuming the obsolete term â€œsmall headâ€ when microcephaly occurs without cognitive impairment. Neurological Sciences, 2017, 38, 1723-1725.	1.9	6
112	A clinical review on megalencephaly. Medicine (United States), 2017, 96, e6814.	1.0	44
113	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
114	Adaptive psychological structure in childhood hearing impairment: audiological correlations. Acta Otorhinolaryngologica Italica, 2017, 37, 175-179.	1.5	0
115	Ictal blinking, an under-recognized phenomenon: our experience and literature review. Neuropsychiatric Disease and Treatment, 2017, Volume 13, 1435-1439.	2.2	10
116	Clinical spectrum of woolly hair: indications for cerebral involvement. Italian Journal of Pediatrics, 2017, 43, 99.	2.6	9
117	Single-Epileptic Spasms with or without Hypsarrhythmia: A Study of 16 Patients. Journal of Pediatric Epilepsy, 2017, 06, 149-155.	0.2	1
118	Biological Drugs in Guillain-BarrÃ© Syndrome: An Update. Current Neuropharmacology, 2017, 15, 938-950.	2.9	15
119	Levetiracetam in neonatal seizures as first-line treatment: A prospective study. Journal of Pediatric Neurosciences, 2017, 12, 24.	0.3	33
120	Pediatric autoimmune encephalitis. Journal of Pediatric Neurosciences, 2017, 12, 130.	0.3	61
121	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
122	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
123	Unilateral Eye Blinking Arising From the Ictal Ipsilateral Occipital Area. Clinical EEG and Neuroscience, 2016, 47, 243-246.	1.7	32
124	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
125	Total Hemi-overgrowth in Pigmentary Mosaicism of the (Hypomelanosis of) Ito Type. Medicine (United) Tj ETQq1 1 0.784314 rgBT /Ov	1.0	33
126	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

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127	Clinical dissection of childhood occipital epilepsy of Gastaut and prognostic implication. European Journal of Neurology, 2016, 23, 241-246.	3.3	13
128	Seizures Are Not a Prevalent Feature of Wyburn-Mason Syndrome. Journal of Pediatric Epilepsy, 2016, 05, 111-114.	0.2	0
129	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
130	Holoprosencephaly and Recurrent Episodes of Severe Neurogenic Hypernatremia. Journal of Pediatric Neurology, 2016, 14, 025-030.	0.2	0
131	Prognostic Challenges of SCN1A Genetic Mutations: Report on Two Children with Mild Features. Journal of Pediatric Neurology, 2016, 14, 082-088.	0.2	32
132	Congenital muscular dystrophy: from muscle to brain. Italian Journal of Pediatrics, 2016, 42, 78.	2.6	64
133	Array-CGH in pediatric neurology: A prospective observational study. European Journal of Inflammation, 2016, 14, 196-199.	0.5	0
134	Acetaminophen administration in pediatric age: an observational prospective cross-sectional study. Italian Journal of Pediatrics, 2016, 42, 20.	2.6	13
135	Child Abuse Syndrome (CAS): A Newly Recognized Distinct Entity. The Open Neurology Journal, 2016, 10, 30-31.	0.4	1
136	Nervous system involvement in clinical peripheral inflammation: A description of three pediatric cases. Journal of Pediatric Neurosciences, 2016, 11, 277.	0.3	4
137	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
138	Headache, migraine and obesity: an overview on plausible links. Journal of Biological Regulators and Homeostatic Agents, 2016, 30, 333-8.	0.7	10
139	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
140	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
141	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
142	Peripheral neuropathy in a child with Mycoplasma pneumoniae infections. Journal of Pediatric Infectious Diseases, 2015, 02, 163-166.	0.2	0
143	Congenital generalized hypertrichosis: the skin as a clue to complex malformation syndromes. Italian Journal of Pediatrics, 2015, 41, 55.	2.6	38
144	Asthma and hypoxia. Italian Journal of Pediatrics, 2015, 41, .	2.6	0

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145	Nevus Sebaceous and Its Association With Neurologic Involvement. Seminars in Pediatric Neurology, 2015, 22, 302-309.	2.0	6
146	Hypomelanosis of Ito: a round on the frequency and type of epileptic complications. Neurological Sciences, 2015, 36, 1173-1180.	1.9	37
147	Refractory absence seizures: An Italian multicenter retrospective study. European Journal of Paediatric Neurology, 2015, 19, 660-664.	1.6	17
148	Propranolol: Effectiveness and Failure in Infantile Cutaneous Hemangiomas. Drug Safety - Case Reports, 2015, 2, 6.	0.9	10
149	The natural history of spinal neurofibromatosis: a critical review of clinical and genetic features. Clinical Genetics, 2015, 87, 401-410.	2.0	81
150	Epilepsy and innate immune system: A possible immunogenic predisposition and related therapeutic implications. Human Vaccines and Immunotherapeutics, 2015, 11, 2021-2029.	3.3	59
151	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
152	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
153	Natalizumab in multiple sclerosis: discontinuation, progressive multifocal leukoencephalopathy and possible use in children. Expert Review of Neurotherapeutics, 2015, 15, 1321-1341.	2.8	5
154	Legumes steam allergy in childhood: Update of the reported cases. Allergologia Et Immunopathologia, 2015, 43, 196-202.	1.7	5
155	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
156	Three Cases of Palatal Tics and Gilles De La Tourette Syndrome. Journal of Child Neurology, 2015, 30, 1199-1203.	1.4	2
157	A Selective Mutism Arising from First Language Attrition, Successfully Treated with Paroxetine-CBT Combination Treatment. Psychiatry Investigation, 2015, 12, 569.	1.6	2
158	Targeting inflammation as a therapeutic strategy for drug-resistant epilepsies. Human Vaccines and Immunotherapeutics, 2014, 10, 868-875.	3.3	47
159	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
160	Usefulness of video-EEG in the paediatric emergency department. Expert Review of Neurotherapeutics, 2014, 14, 769-785.	2.8	3
161	Polysomnographic evaluation of nonâ€invasive ventilation in children with neuromuscular disease. Respirology, 2014, 19, 80-84.	2.3	10
162	Clinical guidelines in pediatric headache: evaluation of quality using the AGREE II instrument. Journal of Headache and Pain, 2014, 15, 57.	6.0	30

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163	Hydranencephaly: cerebral spinal fluid instead of cerebral mantles. Italian Journal of Pediatrics, 2014, 40, 79.	2.6	71
164	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
165	Rapid Spontaneous Resolution of Fibromatosis Colli in a 3-Week-Old Girl. Case Reports in Otolaryngology, 2014, 2014, 1-4.	0.2	15
166	Infantile spasms syndrome, West syndrome and related phenotypes: What we know in 2013. Brain and Development, 2014, 36, 739-751.	1.1	159
167	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
168	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
169	The immunomodulatory effect of probiotics beyond atopy: an update. Journal of Asthma, 2014, 51, 320-332.	1.7	39
170	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
171	Valproate in adolescents with photosensitive epilepsy with generalized tonicâ€“clonic seizures only. European Journal of Paediatric Neurology, 2014, 18, 13-18.	1.6	11
172	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
173	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
174	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
175	Hemihydranencephaly: living with half brain dysfunction. Italian Journal of Pediatrics, 2013, 39, 3.	2.6	30
176	Reflex myoclonic epilepsy in infancy: a critical review. Epileptic Disorders, 2013, 15, 114-122.	1.3	14
177	Sublingual immunotherapy in preschool children: an update. Expert Review of Clinical Immunology, 2013, 9, 385-390.	3.0	6
178	Noninvasive ventilation for acute respiratory distress in children with central nervous system disorders. Respiratory Medicine, 2013, 107, 1370-1375.	2.9	17
179	Reflex myoclonic epilepsy in infancy: A multicenter clinical study. Epilepsy Research, 2013, 103, 237-244.	1.6	8
180	Electroclinical Features and Long-Term Outcome of Cryptogenic Epilepsy in Children with Down Syndrome. Journal of Pediatrics, 2013, 163, 1754-1758.	1.8	25

#	Article	IF	Citations
181	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
182	Natural history of neurofibromatosis type 2 with onset before the age of 1Âyear. Neurogenetics, 2013, 14, 89-98.	1.4	55
183	Gelastic seizures due to hypothalamic hamartoma: Rapid resolution after endoscopic tumor disconnection. Neurocase, 2013, 19, 458-461.	0.6	24
184	Cutaneous and Leptomeningeal Hemangiomas With Impressive Benign Evolution. Pediatric Neurology, 2013, 48, 73-75.	2.1	1
185	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
186	Acute Glomerulonephritis in a Child withChlamydia pneumoniaeInfection: A Case Report. Case Reports in Medicine, 2013, 2013, 1-4.	0.7	2
187	Autonomic Dysfunction Manifesting With Asymmetric Face Flushing and Paroxysmal Nonconvulsive Episodes. Journal of Child Neurology, 2013, 28, 1673-1676.	1.4	6
188	Clinical dissection of early onset absence epilepsy in children and prognostic implications. Epilepsia, 2013, 54, 1761-1770.	5.1	14
189	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
190	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
191	Congenital familial myasthenic syndromes: disease and course in an affected dizygotic twin pair. BMJ Case Reports, 2013, 2013, bcr2012007651-bcr2012007651.	0.5	2
192	West Syndrome Treatment: New Roads for an Old Syndrome. Frontiers in Neurology, 2013, 4, 113.	2.4	34
193	Vitamin D3 an ever green molecule. Frontiers in Bioscience - Scholar, 2013, S5, 247-260.	2.1	8
194	Congenital myasthenic syndromes: Clinical and molecular report on 7 Sicilian patients. Journal of Pediatric Neurosciences, 2013, 8, 19.	0.3	7
195	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
196	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
197	Non-Epileptik Psikojen NÃ¶betler: Klinik Pratikte Ã‡Ã¶zÃ¼lmesi Zor TanÄ±sal Bir Problem. Noropsikiyatri Arsivi, 2012, 49, 243-247.	0.7	0
198	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

#	Article	IF	Citations
199	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
200	Acquired Peripheral Neuropathy: A Report on 20 Children. International Journal of Immunopathology and Pharmacology, 2012, 25, 513-517.	2.1	31
201	Hepatic echinococcosis: Clinical and therapeutic aspects. World Journal of Gastroenterology, 2012, 18, 1448.	3.3	253
202	Microcephaly associated with Legg-CalvÃ¨-Perthes disease in two siblings. Neurological Sciences, 2012, 33, 1401-1405.	1.9	2
203	Congenital talipes equinovarus: an epidemiological study in Sicily. Monthly Notices of the Royal Astronomical Society: Letters, 2012, 83, 294-298.	3.3	36
204	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
205	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
206	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
207	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
208	Diagnosis and management of catamenial seizures: a review. International Journal of Women's Health, 2012, 4, 535.	2.6	31
209	Ohtahara syndrome with emphasis on recent genetic discovery. Brain and Development, 2012, 34, 459-468.	1.1	89
210	Generalised epilepsy with febrile seizures plus (GEFS+): molecular analysis in a restricted area. Child's Nervous System, 2012, 28, 141-145.	1.1	7
211	The role of micronutrients in the diet of HIV-1-infected individuals. Frontiers in Bioscience - Elite, 2012, E4, 2442-2456.	1.8	21
212	Tracheostomy in childhood: new causes for an old strategy. European Review for Medical and Pharmacological Sciences, 2012, 16, 1719-22.	0.7	9
213	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
214	Long-term outcome of epilepsy in Kabuki syndrome. Seizure: the Journal of the British Epilepsy Association, 2011, 20, 650-654.	2.0	10
215	Primary headache: Role of investigations in a cohort of young children and adolescents. Pediatrics International, 2011, 53, 964-967.	0.5	8
216	Longâ€term neurological outcome of a sextuplet pregnancy. Acta Paediatrica, International Journal of Paediatrics, 2011, 100, 777-779.	1.5	1

#	Article	IF	Citations
217	Acute asthma in children: treatment in emergency. European Review for Medical and Pharmacological Sciences, 2011, 15, 711-6.	0.7	1
218	Recurrent obstructive hydrocephalus in a 4-month-old infant. Child's Nervous System, 2010, 26, 133-136.	1.1	3
219	Microcephaly, sensorineural deafness and Currarino triad with duplicationâ€“deletion of distal 7q. European Journal of Pediatrics, 2010, 169, 475-481.	2.7	22
220	Neonatal onset of bath-induced alternating hemiplegia of childhood. European Journal of Paediatric Neurology, 2010, 14, 192-193.	1.6	8
221	A boy born with multiple lesions of atrophoderma. Acta Paediatrica, International Journal of Paediatrics, 2010, 99, 460-463.	1.5	0
222	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
223	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
224	The aristaless Arx gene one gene for many interneuronopathies. Frontiers in Bioscience - Elite, 2010, E2, 701-710.	1.8	18
225	Lissencephalic syndromes brain and beyond. Frontiers in Bioscience - Scholar, 2010, S2, 85-95.	2.1	7
226	Acute Disseminated Encephalomyelitis: A Long-Term Prospective Study and Meta-Analysis. Neuropediatrics, 2010, 41, 246-255.	0.6	75
227	Neonatal Autoimmune Hypothyroidism: A Patient Report. Journal of Pediatric Endocrinology and Metabolism, 2010, 23, 957-61.	0.9	0
228	Neurological Manifestations in Individuals with Pure Cutaneous or Syndromic (Ruggieri-Happle) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 30	0.6	26
229	Recurrent Hopkin's syndrome: A case report and review of the literature. Journal of the Neurological Sciences, 2010, 297, 89-91.	0.6	16
230	Dysembryoplastic Neuroepithelial Tumors: A Prospective Clinicopathologic and Outcome Study of 13 Children. Pediatric Neurology, 2010, 43, 395-402.	2.1	29
231	Blepharoptosis in children: our experience at the light of literature. Clinica Terapeutica, 2010, 161, 241-3.	0.3	2
232	Incidence and causes of neonatal hyperbilirubinemia in a center of Catania. Therapeutics and Clinical Risk Management, 2009, 5, 247.	2.0	4
233	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
234	A Syndrome with Coarse Face, Mental Retardation and Unusual Stereotyped Movements. Neuropediatrics, 2009, 40, 186-188.	0.6	1

#	Article	IF	Citations
235	Infantile spasms in the setting of Sturgeâ€“Weber syndrome. Child's Nervous System, 2009, 25, 111-118.	1.1	41
236	Neurofibromatosis type 1 and infantile spasms. Child's Nervous System, 2009, 25, 211-216.	1.1	55
237	Two siblings with a homozygous MTHFR C677T (G80A-RFC1) mutation and stroke. Child's Nervous System, 2009, 25, 361-365.	1.1	6
238	Chediak-Steinbrinck-Higashi Syndrome. , 2009, , 314-314.		0
239	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
240	Ganglioneuroblastoma-associated Vitamin D Deficiency Rickets. Journal of Pediatric Hematology/Oncology, 2009, 31, 502-504.	0.6	3
241	Symptomatic hypocalcemia in an epileptic child treated with valproic acid plus lamotrigine: a case report. Cases Journal, 2009, 2, 7394.	0.4	9
242	Voluntary selective big toe dorsal flection: pseudo-Babinski phenomenon?. Neurological Sciences, 2008, 29, 495-496.	1.9	0
243	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
244	Low Prevalence of Neurologic and Psychiatric Manifestations in Children with Gluten Sensitivity. Journal of Pediatrics, 2008, 152, 244-249.e1.	1.8	71
245	Pseudohypoparathyroidism Ia With Evans Syndrome. Journal of Pediatric Hematology/Oncology, 2008, 30, 628-630.	0.6	1
246	Congenital Lymphedema-lymphangiectasia Associated With Scrotal Angiokeratoma (Fordyce Type) and Hearing Impairment. Journal of Clinical Gastroenterology, 2008, 42, 715-719.	2.2	3
247	Rotavirus and Celiac Disease. American Journal of Gastroenterology, 2007, 102, 1831-1831.	0.4	15
248	Protective effects of the sigma agonist Pre-084 in the rat retina. British Journal of Ophthalmology, 2007, 91, 1382-1384.	3.9	29
249	Chronic Yersinia Enterocolitis Infection Presenting as Intestinal Obstruction. Infectious Diseases in Clinical Practice, 2007, 15, 68-69.	0.3	0
250	Polyneuritis Cranialis: Full Recovery after Intravenous Immunoglobulins. Pediatric Neurology, 2007, 37, 209-211.	2.1	4
251	Otologic findings in children with gastroesophageal reflux. International Journal of Pediatric Otorhinolaryngology, 2007, 71, 1693-1697.	1.0	18
252	Anton-Babinski syndrome in a child with early-stage adrenoleukodystrophy. European Journal of Neurology, 2007, 14, e11-e12.	3.3	13

#	Article	IF	Citations
253	Callosal anomalies with interhemispheric cyst: Expanding the phenotype. Acta Paediatrica, International Journal of Paediatrics, 2007, 94, 1066-1072.	1.5	13
254	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
255	Septo-Optic Dysplasia Complex: A Heterogeneous Malformation Syndrome. Pediatric Neurology, 2006, 34, 66-71.	2.1	81
256	Blink reflex abnormalities in children with Tourette syndrome. European Journal of Neurology, 2006, 13, 869-873.	3.3	4
257	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
258	STROKE IN TWO CHILDREN WITH MYCOPLASMA PNEUMONIAE INFECTION A CAUSAL OR CASUAL RELATIONSHIP?. Pediatric Infectious Disease Journal, 2005, 24, 843-845.	2.0	36
259	Liver transplantation in a child with celiac disease. Journal of Gastroenterology and Hepatology (Australia), 2005, 20, 956-960.	2.8	13
260	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
261	A Complex Brain Malformation Syndrome with Rhombencephalosynapsis, Preaxial Hexadactyly plus Facial and Skull Anomalies. Neuropediatrics, 2005, 36, 279-283.	0.6	10
262	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
263	Hypertrichosis Cubiti (Hairy Elbow Syndrome): A Clue to a Malformation Syndrome. Journal of Pediatric Endocrinology and Metabolism, 2005, 18, 1019-25.	0.9	34
264	Callosal anomalies with interhemispheric cyst: Expanding the phenotype. Acta Paediatrica, International Journal of Paediatrics, 2005, 94, 1066-1072.	1.5	7
265	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
266	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
267	Epilepsy is not a Prominent Feature of Primary Autism. Neuropediatrics, 2004, 35, 207-210.	0.6	71
268	Ophthalmological manifestations in segmental neurofibromatosis type 1. British Journal of Ophthalmology, 2004, 88, 1429-1433.	3.9	55
269	Anti-brain antibodies in PANDAS versus uncomplicated streptococcal infection. Pediatric Neurology, 2004, 30, 107-110.	2.1	114
270	Peripheral Neuropathy as First Sign of Ulcerative Colitis in a Child. Journal of Clinical Gastroenterology, 2004, 38, 115-117.	2.2	14

#	Article	IF	Citations
271	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
272	Worldwide distribution and broader clinical spectrum of muscle-eye-brain disease. Human Molecular Genetics, 2003, 12, 527-534.	2.9	133
273	Reply to Dr Hahn. Journal of Child Neurology, 2003, 18, 152-152.	1.4	0
274	Lobar Holoprosencephaly. Journal of Child Neurology, 2002, 17, 543-544.	1.4	2
275	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
276	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
277	Clinical heterogeneity in eyelid myoclonia, with absences, and epilepsy. European Journal of Pediatrics, 2002, 161, 175-177.	2.7	12
278	A Genomewide Screen for Autism Susceptibility Loci. American Journal of Human Genetics, 2001, 69, 327-340.	6.2	287
279	Lack of progression of brain atrophy in Aicardi-GoutiÃ¨res syndrome. Pediatric Neurology, 2001, 24, 300-302.	2.1	38
280	Neuropsychological assessment in children with absence epilepsy. Neurology, 2001, 56, 1047-1051.	1.1	168
281	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
282	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
283	Holoprosencephaly (Lobar Form) Associated With Bilateral Vocal Cord Palsy. Journal of Child Neurology, 2001, 16, 932-934.	1.4	5
284	Hemihydranencephaly: Case Report and Literature Review. Journal of Child Neurology, 2001, 16, 218-221.	1.4	20
285	Fetal Nucleated Red Blood Cell Counts in Peripheral Blood of Mothers Bearing Down Syndrome Fetus. Neuropediatrics, 2001, 32, 147-149.	0.6	10
286	Hemihydranencephaly: Case Report and Literature Review. Journal of Child Neurology, 2001, 16, 212.	1.4	0
287	Colobomatous microphthalmia, microcephaly with cerebellar hypoplasia: Association or new syndrome?. , 2000, 92, 278-280.		3
288	Tuberous breast deformity in an adolescent girl with Hurler-Scheie syndrome. European Journal of Pediatrics, 2000, 159, 936-937.	2.7	9

#	Article	IF	Citations
289	Arthrogryposis Multiplex Congenita and Pituitary Ectopia. A Case Report. Neuropediatrics, 2000, 31, 325-327.	0.6	8
290	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
291	Recurrent peripheral neuropathy in a girl with celiac disease. Journal of Neurology, Neurosurgery and Psychiatry, 2000, 68, 104-105.	1.9	30
292	Diabetes insipidus in neurobrucellosis. Clinical Neurology and Neurosurgery, 2000, 102, 163-165.	1.4	22
293	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
294	Colobomatous microphthalmia, microcephaly with cerebellar hypoplasia: association or new syndrome?. American Journal of Medical Genetics Part A, 2000, 92, 278-80.	2.4	0
295	Late Primary Unilateral Thalamic Hemorrhage in Infancy: Report of Two Cases. Neuropediatrics, 1999, 30, 264-267.	0.6	26
296	Focal Neurological Deficits in Children with Î²-Thalassemia Major. Neuropediatrics, 1999, 30, 45-48.	0.6	21
297	Vein of Galen Malformation and Infantile Spasms. Journal of Child Neurology, 1999, 14, 196-198.	1.4	3
298	Extraordinary intrathecal bone reaction in Î²-thalassaemia intermedia. Lancet, The, 1999, 354, 922.	13.7	4
299	Marin-Amat Syndrome: Case Report and Review of the Literature. Journal of Child Neurology, 1999, 14, 266-268.	1.4	33
300	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