Neil R Friedman

List of Publications by Year
in descending order

47 papers	2,684 citations	257450 24 h-index	243625 44 g-index
47 all docs	47 docs citations	47 times ranked	4093 citing authors

#	Article	IF	Citations
1	Deletion of conserved nonâ€coding sequences downstream from NKX2â€1: A novel diseaseâ€causing mechanism for benign hereditary chorea. Molecular Genetics & Genomic Medicine, 2021, 9, e1647.	1.2	3
2	Child Neurology: Genetically determined dystonias with childhood onset. Neurology, 2020, 94, 892-895.	1.1	3
3	Arterial Ischemic Stroke Secondary to Cardiac Disease in Neonates and Children. Pediatric Neurology, 2019, 100, 35-41.	2.1	25
4	Dyslipidemia in Children With Arterial Ischemic Stroke: Prevalence and Risk Factors. Pediatric Neurology, 2018, 78, 46-54.	2.1	20
5	Focal Cerebral Arteriopathy of Childhood. Stroke, 2018, 49, 2590-2596.	2.0	46
6	Clinical and Imaging Characteristics of Arteriopathy Subtypes in Children with Arterial Ischemic Stroke: Results of the VIPS Study. American Journal of Neuroradiology, 2017, 38, 2172-2179.	2.4	89
7	Muscle biopsy findings in a child with NALCN gene mutation. Journal of Clinical Neuroscience, 2016, 34, 222-223.	1.5	6
8	Inflammatory Biomarkers in Childhood Arterial Ischemic Stroke. Stroke, 2016, 47, 2221-2228.	2.0	38
9	Congenital glioneuronal tumor with neuropil-like islands. Journal of Clinical Neuroscience, 2016, 24, 156-157.	1.5	4
10	Risk of Recurrent Arterial Ischemic Stroke in Childhood. Stroke, 2016, 47, 53-59.	2.0	138
11	Arterial Tortuosity: An Imaging Biomarker of Childhood Stroke Pathogenesis?. Stroke, 2016, 47, 1265-1270.	2.0	22
12	Beta Blockade as Treatment for Intracranial Infantile Hemangioma: Case Report and Literature Review. Pediatric Neurology, 2016, 59, 13-17.	2.1	11
13	The Way Forward: Challenges and Opportunities in Pediatric Stroke. Pediatric Neurology, 2016, 56, 3-7.	2.1	10
14	Preparing for a â€œPediatric Stroke Alertâ€. Pediatric Neurology, 2016, 56, 18-24.	2.1	39
15	Pediatric cardiovascular disease and stroke. Journal of Pediatric Neurology, 2015, 08, 259-265.	0.2	1
16	Stroke in Children With Cardiac Disease: Report From the International Pediatric Stroke Study Group Symposium. Pediatric Neurology, 2015, 52, 5-15.	2.1	55
17	Enterovirus D68: A clinically important respiratory enterovirus. Cleveland Clinic Journal of Medicine, 2015, 82, 26-31.	1.3	23
18	Arteriopathy Diagnosis in Childhood Arterial Ischemic Stroke. Stroke, 2014, 45, 3597-3605.	2.0	130

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19	Emergence of the Primary Pediatric Stroke Center. Stroke, 2014, 45, 2018-2023.	2.0	108
20	â€˜Double troubleâ€™: Diagnostic challenges in Duchenne muscular dystrophy in patients with an additional hereditary skeletal dysplasia. Neuromuscular Disorders, 2013, 23, 955-961.	0.6	7
21	Neonatal Stroke and Progressive Leukoencephalopathy in a Child With an ACTA2 Mutation. Journal of Child Neurology, 2013, 28, 531-534.	1.4	25
22	Cerebral Vasculopathy in Children With Neurofibromatosis Type 1. Journal of Child Neurology, 2013, 28, 95-101.	1.4	51
23	Fetal magnetic resonance imaging in hydranencephaly. Journal of Paediatrics and Child Health, 2013, 49, 335-336.	0.8	13
24	Tuberculous Radiculomyelitis Presenting in a Toddler With Lower Extremity Weakness and Seizure. Pediatric Infectious Disease Journal, 2013, 32, 919-921.	2.0	3
25	Fever, Lethargy, and Leg Weakness in a 9-Month-Old Boy. Clinical Pediatrics, 2012, 51, 808-811.	0.8	0
26	Authorsâ€™ Reply in Response to the Correspondence About the Article â€œHirayama Disease in Children From North Americaâ€•Published in Journal of Child Neurology Dec 2011 Issue. Journal of Child Neurology, 2012, 27, 817-818.	1.4	0
27	Pitt-Hopkins Syndrome in a Boy With Charcot Marie Tooth Disease Type 1A: A Rare Co-occurrence of 2 Genetic Disorders. Journal of Child Neurology, 2012, 27, 1602-1606.	1.4	8
28	Phenotypic Heterogeneity of Genomic Disorders and Rare Copy-Number Variants. New England Journal of Medicine, 2012, 367, 1321-1331.	27.0	519
29	Ophthalmologic Abnormalities in Mowat-Wilson Syndrome and a Mutation inZEB2. Ophthalmic Genetics, 2012, 33, 159-160.	1.2	21
30	Concurrent Validity and Reliability of Retrospective Scoring of the Pediatric National Institutes of Health Stroke Scale. Stroke, 2012, 43, 341-345.	2.0	46
31	Phosphoribosylpyrophosphate synthetase superactivity and recurrent infections is caused by a p.Val142Leu mutation in PRSâ€‰. American Journal of Medical Genetics, Part A, 2012, 158A, 455-460.	1.2	26
32	Case series: 2q33.1 microdeletion syndrome--further delineation of the phenotype. Journal of Medical Genetics, 2011, 48, 290-298.	3.2	59
33	Hirayama Disease in Children From North America. Journal of Child Neurology, 2011, 26, 1542-1547.	1.4	37
34	Reversible Cerebral Vasoconstriction Syndrome. Journal of Child Neurology, 2011, 26, 1580-1584.	1.4	23
35	Interrater Reliability of the Pediatric National Institutes of Health Stroke Scale (PedNIHSS) in a Multicenter Study. Stroke, 2011, 42, 613-617.	2.0	135
36	Small vessel childhood primary angiitis of the CNS: first steps toward a standardised treatment regimen. Lancet Neurology, The, 2010, 9, 1042-1044.	10.2	1

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37	Genetically Confirmed CADASIL in a Pediatric Patient. Pediatrics, 2010, 126, e1603-e1607.	2.1	16
38	Predictors of Cerebral Arteriopathy in Children With Arterial Ischemic Stroke. Circulation, 2009, 119, 1417-1423.	1.6	314
39	Utility of cystatin C to monitor renal function in duchenne muscular dystrophy. Muscle and Nerve, 2009, 40, 438-442.	2.2	84
40	Pediatric Stroke: Past, Present and Future. Advances in Pediatrics, 2009, 56, 271-299.	1.4	24
41	Pituitary stalk duplication in association with moya moya disease and bilateral morning glory disc anomaly â€“ broadening the clinical spectrum of midline defects. Journal of Neurology, 2008, 255, 885-890.	3.6	20
42	Increased MECP2 gene copy number as the result of genomic duplication in neurodevelopmentally delayed males. Genetics in Medicine, 2006, 8, 784-792.	2.4	245
43	Horizontal gaze palsy with progressive scoliosis can result from compound heterozygous mutations in ROBO3. Journal of Medical Genetics, 2005, 43, e11-e11.	3.2	70
44	Oxidative phosphorylation analysis: assessing the integrated functional activity of human skeletal muscle mitochondriaâ€"case studies. Mitochondrion, 2004, 4, 377-385.	3.4	63
45	Cardiac transplantation in twins with autosomal dominant Emery-Dreifuss muscular dystrophy. Journal of Heart and Lung Transplantation, 2004, 23, 496-498.	0.6	20
46	Metabolic myopathies: a clinical approach; part II. Pediatric Neurology, 2000, 22, 171-181.	2.1	32
47	Metabolic myopathies: a clinical approach; part I. Pediatric Neurology, 2000, 22, 87-97.	2.1	51