Partha S Ghosh

List of Publications by Year
in descending order

44	332	933447 10	888059 17
papers	citations	h-index	g-index

44	44	44	583
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Electromyography in infants: experience from a pediatric neuromuscular center. Acta Neurologica Belgica, 2022, 122, 1195-1200.	1.1	2
2	6-month old with new-onset weakness â€“ consider infant botulism. Archives of Disease in Childhood, 2022, 107, 380-380.	1.9	0
3	â€œSpokes of Wheelâ€•in Muscle Biopsy. Journal of Clinical Neuromuscular Disease, 2022, 23, 160-161.	0.7	0
4	Unilateral Cavovarus Foot From Sciatic Nerve Perineurioma. Neurology India, 2022, 70, 178.	0.4	0
5	Congenital Myasthenic Syndrome From a Single Center: Phenotypic and Genotypic features. Journal of Child Neurology, 2021, 36, 610-617.	1.4	12
6	Neuroimage: the crescent moon sign of Hirayama disease. Acta Neurologica Belgica, 2021, , 1.	1.1	3
7	Developing outcome measures of disease activity in pediatric myasthenia. Muscle and Nerve, 2021, 63, 751-757.	2.2	0
8	Postsurgical Inflammatory Sciatic Neuropathy. Journal of Clinical Neuromuscular Disease, 2021, 22, 237-238.	0.7	1
9	Aphasia and Weakness in a Child Think Beyond Stroke. Journal of Clinical Neuromuscular Disease, 2021, 22, 233-234.	0.7	0
10	BET1 variants establish impaired vesicular transport as a cause for muscular dystrophy with epilepsy. EMBO Molecular Medicine, 2021, 13, e13787.	6.9	9
11	Myasthenia Gravis in a Child After Graft-versus-Host Disease. Journal of Clinical Neuromuscular Disease, 2021, 23, 53-54.	0.7	0
12	A Boy with Herculean Appearance. Journal of Pediatrics, 2021, , .	1.8	0
13	Acute Neuromuscular Disorders in the Pediatric Intensive Care Unit. Journal of Child Neurology, 2020, 35, 17-24.	1.4	12
14	Response to â€œThe Spectrum of Neuromuscular Disorders Admitted to a Pediatric Intensive Care Unit Is Broader Than Anticipatedâ€• Journal of Child Neurology, 2020, 35, 302-303.	1.4	0
15	Update on Muscular Dystrophies with Focus on Novel Treatments and Biomarkers. Current Neurology and Neuroscience Reports, 2020, 20, 14.	4.2	22
16	Clinical Reasoning: A 10-year-old girl with muscle stiffness. Neurology, 2020, 95, e773-e778.	1.1	1
17	Clinical Reasoning: Seven-year-old girl with progressive gait difficulties. Neurology, 2020, 94, 364-367.	1.1	1
18	Clinical Reasoning: Pes cavus and neuropathy. Neurology, 2019, 93, e823-e826.	1.1	1

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19	Clinical Reasoning: A teenager with left arm weakness. Neurology, 2018, 90, e907-e910.	1.1	0
20	Arthrogryposis Multiplex Congenita. , 2018, , 874-886.e4.		1
21	Neuralgic amyotrophy in children. Muscle and Nerve, 2018, 57, 932-936.	2.2	13
22	Spectrum of Neuromuscular Disorders With HyperCKemia From a Tertiary Care Pediatric Neuromuscular Center. Journal of Child Neurology, 2018, 33, 389-396.	1.4	12
23	Loss of tubulin deglutamylase <scp>CCP</scp> 1 causes infantileâ€onset neurodegeneration. EMBO Journal, 2018, 37, .	7.8	86
24	Clinical Reasoning: A child with arthrogryposis. Neurology, 2018, 91, e995-e998.	1.1	1
25	Clinical Reasoning: Young adult with dysphagia and severe weight loss. Neurology, 2018, 91, e1083-e1086.	1.1	0
26	Spectrum of Nondystrophic Skeletal Muscle Channelopathies in Children. Pediatric Neurology, 2017, 70, 26-33.	2.1	12
27	Schwartz-Jampel Syndrome. Pediatric Neurology, 2017, 68, 77-78.	2.1	5
28	Clinical Reasoning: A child with delayed motor milestones and ptosis. Neurology, 2017, 88, e158-e163.	1.1	1
29	Reliability of a novel ultrasound system for grayâ€scale analysis of muscle. Muscle and Nerve, 2017, 56, 408-412.	2.2	15
30	Brain involvement in Charcotâ€“Marieâ€“Tooth disease due to ganglioside-induced differentiation associated-protein 1 mutation. Neuromuscular Disorders, 2017, 27, 848-851.	0.6	0
31	Clinical Reasoning: A tale of a hypotonic infant. Neurology, 2016, 87, e11-6.	1.1	0
32	Reply. Muscle and Nerve, 2016, 54, 343-344.	2.2	0
33	Clinical Reasoning: A 2-year-old child with acute flaccid paralysis. Neurology, 2016, 87, e149-e154.	1.1	1
34	Teaching Neuro Images : Medullary lesions causing dysphagia in Leigh/MELAS overlap syndrome. Neurology, 2016, 87, e18-9.	1.1	2
35	Myotonic Dystrophy Type 1: A Neurological Cause of Dysphagia. Pediatric Neurology, 2016, 57, 105-106.	2.1	1
36	Camptocormia as presenting manifestation of a spectrum of myopathic disorders. Muscle and Nerve, 2015, 52, 1008-1012.	2.2	34

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37	Recurrent Right-Sided Ptosis in a Child. JAMA Pediatrics, 2015, 169, 693.	6.2	8
38	Clinical and laboratory findings of 21 patients with radiation-induced myopathy. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 152-158.	1.9	32
39	Use of Clinical and Electrical Myotonia to Differentiate Childhood Myopathies. Journal of Child Neurology, 2015, 30, 1300-1306.	1.4	4
40	Pearls & Oy-sters: HyperCKemia with limb-girdle weakness. Neurology, 2014, 83, e209-12.	1.1	3
41	Clinical Reasoning: A 38-year-old woman with childhood-onset weakness. Neurology, 2014, 83, e81-4.	1.1	0
42	Neurologic Complications Following Pediatric Renal Transplantation. Journal of Child Neurology, 2014, 29, 793-798.	1.4	8
43	Inclusionâ€body myositis presenting with facial diplegia. Muscle and Nerve, 2014, 49, 287-289.	2.2	13
44	Diagnostic Yield of Electromyography in Children With Myopathic Disorders. Pediatric Neurology, 2014, 51, 215-219.	2.1	16