# 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 | Loss of tubulin deglutamylase <scp>CCP</scp> 1 causes infantileâ€onset neurodegeneration. EMBO Journal, 2018, 37, . | 7.8 | 86 |
| 2 | Camptocormia as presenting manifestation of a spectrum of myopathic disorders. Muscle and Nerve, 2015, 52, 1008-1012. | 2.2 | 34 |
| 3 | Clinical and laboratory findings of 21 patients with radiation-induced myopathy. Journal of Neurology, Neurosurgery and Psychiatry, 2015, 86, 152-158. | 1.9 | 32 |
| 4 | Update on Muscular Dystrophies with Focus on Novel Treatments and Biomarkers. Current Neurology and Neuroscience Reports, 2020, 20, 14. | 4.2 | 22 |
| 5 | Diagnostic Yield of Electromyography in Children With Myopathic Disorders. Pediatric Neurology, 2014, 51, 215-219. | 2.1 | 16 |
| 6 | Reliability of a novel ultrasound system for grayâ€scale analysis of muscle. Muscle and Nerve, 2017, 56, 408-412. | 2.2 | 15 |
| 7 | Inclusionâ€body myositis presenting with facial diplegia. Muscle and Nerve, 2014, 49, 287-289. | 2.2 | 13 |
| 8 | Neuralgic amyotrophy in children. Muscle and Nerve, 2018, 57, 932-936. | 2.2 | 13 |
| 9 | Spectrum of Nondystrophic Skeletal Muscle Channelopathies in Children. Pediatric Neurology, 2017, 70, 26-33. | 2.1 | 12 |
| 10 | Spectrum of Neuromuscular Disorders With HyperCKemia From a Tertiary Care Pediatric Neuromuscular Center. Journal of Child Neurology, 2018, 33, 389-396. | 1.4 | 12 |
| 11 | Acute Neuromuscular Disorders in the Pediatric Intensive Care Unit. Journal of Child Neurology, 2020, 35, 17-24. | 1.4 | 12 |
| 12 | Congenital Myasthenic Syndrome From a Single Center: Phenotypic and Genotypic features. Journal of Child Neurology, 2021, 36, 610-617. | 1.4 | 12 |
| 13 | BET1 variants establish impaired vesicular transport as a cause for muscular dystrophy with epilepsy. EMBO Molecular Medicine, 2021, 13, e13787. | 6.9 | 9 |
| 14 | Neurologic Complications Following Pediatric Renal Transplantation. Journal of Child Neurology, 2014, 29, 793-798. | 1.4 | 8 |
| 15 | Recurrent Right-Sided Ptosis in a Child. JAMA Pediatrics, 2015, 169, 693. | 6.2 | 8 |
| 16 | Schwartz-Jampel Syndrome. Pediatric Neurology, 2017, 68, 77-78. | 2.1 | 5 |
| 17 | Use of Clinical and Electrical Myotonia to Differentiate Childhood Myopathies. Journal of Child Neurology, 2015, 30, 1300-1306. | 1.4 | 4 |
| 18 | Pearls & Oy-sters: HyperCKemia with limb-girdle weakness. Neurology, 2014, 83, e209-12. | 1.1 | 3 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Neuroimage: the crescent moon sign of Hirayama disease. Acta Neurologica Belgica, 2021, , 1. | 1.1 | 3 |
| 20 | Teaching Neuro Images : Medullary lesions causing dysphagia in Leigh/MELAS overlap syndrome. Neurology, 2016, 87, e18-9. | 1.1 | 2 |
| 21 | Electromyography in infants: experience from a pediatric neuromuscular center. Acta Neurologica Belgica, 2022, 122, 1195-1200. | 1.1 | 2 |
| 22 | Clinical Reasoning: A 2-year-old child with acute flaccid paralysis. Neurology, 2016, 87, e149-e154. | 1.1 | 1 |
| 23 | Myotonic Dystrophy Type 1: A Neurological Cause of Dysphagia. Pediatric Neurology, 2016, 57, 105-106. | 2.1 | 1 |
| 24 | Clinical Reasoning: A child with delayed motor milestones and ptosis. Neurology, 2017, 88, e158-e163. | 1.1 | 1 |
| 25 | Arthrogryposis Multiplex Congenita. , 2018, , 874-886.e4. | | 1 |
| 26 | Clinical Reasoning: A child with arthrogryposis. Neurology, 2018, 91, e995-e998. | 1.1 | 1 |
| 27 | Clinical Reasoning: Pes cavus and neuropathy. Neurology, 2019, 93, e823-e826. | 1.1 | 1 |
| 28 | Clinical Reasoning: A 10-year-old girl with muscle stiffness. Neurology, 2020, 95, e773-e778. | 1.1 | 1 |
| 29 | Clinical Reasoning: Seven-year-old girl with progressive gait difficulties. Neurology, 2020, 94, 364-367. | 1.1 | 1 |
| 30 | Postsurgical Inflammatory Sciatic Neuropathy. Journal of Clinical Neuromuscular Disease, 2021, 22, 237-238. | 0.7 | 1 |
| 31 | Clinical Reasoning: A 38-year-old woman with childhood-onset weakness. Neurology, 2014, 83, e81-4. | 1.1 | 0 |
| 32 | Clinical Reasoning: A tale of a hypotonic infant. Neurology, 2016, 87, e11-6. | 1.1 | 0 |
| 33 | Reply. Muscle and Nerve, 2016, 54, 343-344. | 2.2 | 0 |
| 34 | 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 |
| 35 | Clinical Reasoning: A teenager with left arm weakness. Neurology, 2018, 90, e907-e910. | 1.1 | 0 |
| 36 | Clinical Reasoning: Young adult with dysphagia and severe weight loss. Neurology, 2018, 91, e1083-e1086. | 1.1 | 0 |

| # | Article | IF | Citations |
|---|---|---|---|
| 37 | 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 |
| 38 | Developing outcome measures of disease activity in pediatric myasthenia. Muscle and Nerve, 2021, 63, 751-757. | 2.2 | 0 |
| 39 | Aphasia and Weakness in a Child Think Beyond Stroke. Journal of Clinical Neuromuscular Disease, 2021, 22, 233-234. | 0.7 | 0 |
| 40 | 6-month old with new-onset weakness â€“ consider infant botulism. Archives of Disease in Childhood, 2022, 107, 380-380. | 1.9 | 0 |
| 41 | Myasthenia Gravis in a Child After Graft-versus-Host Disease. Journal of Clinical Neuromuscular Disease, 2021, 23, 53-54. | 0.7 | 0 |
| 42 | â€œSpokes of Wheelâ€•in Muscle Biopsy. Journal of Clinical Neuromuscular Disease, 2022, 23, 160-161. | 0.7 | 0 |
| 43 | A Boy with Herculean Appearance. Journal of Pediatrics, 2021, , . | 1.8 | 0 |
| 44 | Unilateral Cavovarus Foot From Sciatic Nerve Perineurioma. Neurology India, 2022, 70, 178. | 0.4 | 0 |