

Sindhu Ramchandren

List of Publications by Year in descending order

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Version: 2024-02-01

10
papers

469
citations

1307594

7
h-index

1372567

10
g-index

10
all docs

10
docs citations

10
times ranked

709
citing authors

#	ARTICLE	IF	CITATIONS
1	Development and Validation of the Pediatric Charcot-Marie-Tooth Disease Quality of Life Outcome Measure. <i>Annals of Neurology</i> , 2021, 89, 369-379.	5.3	13
2	A longitudinal study of CMT1A using Rasch analysis based CMT neuropathy and examination scores. <i>Neurology</i> , 2020, 94, e884-e896.	1.1	29
3	Natural history of Charcot-Marie-Tooth disease type 2A: a large international multicentre study. <i>Brain</i> , 2020, 143, 3589-3602.	7.6	39
4	Charcot-Marie-Tooth Disease and Other Genetic Polyneuropathies. <i>CONTINUUM Lifelong Learning in Neurology</i> , 2017, 23, 1360-1377.	0.8	25
5	Knowledge of carrier status and barriers to testing among mothers of sons with Duchenne or Becker muscular dystrophy. <i>Neuromuscular Disorders</i> , 2016, 26, 860-864.	0.6	10
6	Outdated risk assessment in a family with Duchenne dystrophy: Implications for duty to reassess. <i>Neurology: Genetics</i> , 2016, 2, e103.	1.9	3
7	CMT subtypes and disease burden in patients enrolled in the Inherited Neuropathies Consortium natural history study: a cross-sectional analysis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 873-878.	1.9	249
8	Defining disability: development and validation of a mobility-Disability Severity Index (mDSI) in Charcot-Marie-tooth disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 635-639.	1.9	7
9	Genotype-phenotype characteristics and baseline natural history of heritable neuropathies caused by mutations in the <i>MPZ</i> gene. <i>Brain</i> , 2015, 138, 3180-3192.	7.6	80
10	Headache in a patient with Klinefelter's syndrome and hyperostosis frontalis interna. <i>Journal of Headache and Pain</i> , 2007, 8, 342-344.	6.0	14