Vamshi K Rao

List of Publications by Year
in descending order

22 papers	2,272 citations	759055 12 h-index	713332 21 g-index
23 all docs	23 docs citations	23 times ranked	3131 citing authors

#	Article	IF	Citations
1	Palliative Care in Duchenne Muscular Dystrophy: Goals of Care Discussions and Beyond. Muscle and Nerve, 2022, , .	1.0	1
2	An expanded access program of risdiplam for patients with Type 1 or 2 spinal muscular atrophy. Annals of Clinical and Translational Neurology, 2022, 9, 810-818.	1.7	18
3	Long-Term Functional Efficacy and Safety of Viltolarsen in Patients with Duchenne Muscular Dystrophy. Journal of Neuromuscular Diseases, 2022, 9, 493-501.	1.1	31
4	Combination molecular therapies for type 1 spinal muscular atrophy. Muscle and Nerve, 2020, 62, 550-554.	1.0	51
5	Safety, Tolerability, and Efficacy of Viltolarsen in Boys With Duchenne Muscular Dystrophy Amenable to Exon 53 Skipping. JAMA Neurology, 2020, 77, 982.	4.5	169
6	Transforaminal Intrathecal Access for Injection of Nusinersen in Adult and Pediatric Patients with Spinal Muscular Atrophy. Journal of Pediatric Neurology, 2020, 18, 088-094.	0.0	2
7	Spinal muscular atrophy care in the COVIDâ€19 pandemic era. Muscle and Nerve, 2020, 62, 46-49.	1.0	31
8	The care of patients with Duchenne, Becker, and other muscular dystrophies in the <scp>COVID</scp>â€19 pandemic. Muscle and Nerve, 2020, 62, 41-45.	1.0	54
9	Utility of Repetitive Nerve Stimulation in Myopathies. Pediatric Neurology Briefs, 2020, 34, 4.	0.2	1
10	PIGQ glycosylphosphatidylinositolâ€anchored protein deficiency: Characterizing the phenotype. American Journal of Medical Genetics, Part A, 2019, 179, 1270-1275.	0.7	11
11	Spinal Muscular Atrophy Diagnosed by Newborn Screening. Pediatric Neurology Briefs, 2019, 33, 5.	0.2	2
12	Gene Therapy for Spinal Muscular Atrophy: An Emerging Treatment Option for a Devastating Disease. Journal of Managed Care & Specialty Pharmacy, 2018, 24, S3-S16.	0.5	34
13	Friedreichâ€™s Ataxia: Clinical Presentation of a Compound Heterozygote Child with a Rare Nonsense Mutation and Comparison with Previously Published Cases. Case Reports in Neurological Medicine, 2018, 2018, 1-5.	0.3	2
14	Nusinersen versus Sham Control in Infantile-Onset Spinal Muscular Atrophy. New England Journal of Medicine, 2017, 377, 1723-1732.	13.9	1,533
15	EPG5-related Vici syndrome: a paradigm of neurodevelopmental disorders with defective autophagy. Brain, 2016, 139, 765-781.	3.7	99
16	Guidelines for Corticosteroid use in Treatment of DMD. Pediatric Neurology Briefs, 2016, 30, 21.	0.2	6
17	Delay in Diagnosis of Duchenne Muscular Dystrophy. Pediatric Neurology Briefs, 2015, 29, 5.	0.2	4
18	Orofacial EMG in Congenital Multiple Cranial Neuropathies. Pediatric Neurology Briefs, 2015, 29, 68.	0.2	0

#	ARTICLE	IF	CITATIONS
19	Symptomatic Cerebral Vasospasm Following Resection of a Medulloblastoma in a Child. Neurocritical Care, 2013, 18, 84-88.	1.2	12
20	Mouse Survival Motor Neuron Alleles That Mimic SMN2 Splicing and Are Inducible Rescue Embryonic Lethality Early in Development but Not Late. PLoS ONE, 2010, 5, e15887.	1.1	71
21	Comparative Proteomes of the Proliferating C2C12 Myoblasts and Fully Differentiated Myotubes Reveal the Complexity of the Skeletal Muscle Differentiation Program. Molecular and Cellular Proteomics, 2004, 3, 1065-1082.	2.5	75
22	Paraffin-wax-coated plates as matrix-assisted laser desorption/ionization sample support for high-throughput identification of proteins by peptide mass fingerprinting. Analytical Biochemistry, 2004, 327, 222-232.	1.1	61