

Gayathri Narayanappa

List of Publications by Year in descending order

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Version: 2024-02-01

10
papers

135
citations

1478505

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h-index

1372567

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all docs

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docs citations

10
times ranked

262
citing authors

#	ARTICLE	IF	CITATIONS
1	A Dominant C150Y Mutation in FHL1 Induces Structural Alterations in LIM2 Domain Causing Protein Aggregation In Human and Drosophila Indirect Flight Muscles. Journal of Molecular Neuroscience, 2021, 71, 2324-2335.	2.3	1
2	Infective myositis. Brain Pathology, 2021, 31, e12950.	4.1	25
3	A Novel L1 Linker Mutation in DES Resulted in Total Absence of Protein. Journal of Molecular Neuroscience, 2021, 71, 2468-2473.	2.3	6
4	Diagnosis of primary mitochondrial disorders -Emphasis on myopathological aspects. Mitochondrion, 2021, 61, 69-84.	3.4	4
5	Clinico-pathological and Molecular Spectrum of Mitochondrial Polymerase γ Mutations in a Cohort from India. Journal of Molecular Neuroscience, 2021, 71, 2219-2228.	2.3	6
6	Mitochondrial oxidative phosphorylation disorders in children: Phenotypic, genotypic and biochemical correlations in 85 patients from South India. Mitochondrion, 2017, 32, 42-49.	3.4	17
7	Muscle biopsies from human muscle diseases with myopathic pathology reveal common alterations in mitochondrial function. Journal of Neurochemistry, 2016, 138, 174-191.	3.9	33
8	Microsporidial polymyositis in human immunodeficiency virus infected patients, a rare life threatening opportunistic infection: Clinical suspicion, diagnosis, and management in resource limited settings. Muscle and Nerve, 2015, 51, 775-780.	2.2	7
9	Magnetic resonance imaging correlates of genetically characterized patients with mitochondrial disorders: A study from south India. Mitochondrion, 2015, 25, 6-16.	3.4	28
10	New mutation of the desmin gene identified in an extended Indian pedigree presenting with distal myopathy and cardiac disease. Neurology India, 2013, 61, 622.	0.4	8