

Sonia M Hasan

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

13
papers

159
citations

7
h-index

12
g-index

13
ext. papers

199
ext. citations

4.7
avg, IF

2.1
L-index

#	Paper	IF	Citations
13	Gain-of-function defects of astrocytic Kir4.1 channels in children with autism spectrum disorders and epilepsy. <i>Scientific Reports</i> , 2016 , 6, 34325	4.9	43
12	New insights into the pathogenesis and therapeutics of episodic ataxia type 1. <i>Frontiers in Cellular Neuroscience</i> , 2015 , 9, 317	6.1	40
11	deficiency leads to a syndrome of abnormal striatum, congenital cataract and intellectual disability. <i>Journal of Medical Genetics</i> , 2016 , 53, 786-792	5.8	14
10	Lethal digenic mutations in the K channels Kir4.1 () and SLACK () associated with severe-disabling seizures and neurodevelopmental delay. <i>Journal of Neurophysiology</i> , 2017 , 118, 2402-2411	3.2	12
9	A Calsequestrin-1 Mutation Associated with a Skeletal Muscle Disease Alters Sarcoplasmic Ca ²⁺ Release. <i>PLoS ONE</i> , 2016 , 11, e0155516	3.7	12
8	Hydrogen peroxide-induced reduction of delayed rectifier potassium current in hippocampal neurons involves oxidation of sulfhydryl groups. <i>Brain Research</i> , 2013 , 1520, 61-9	3.7	7
7	A channelopathy mutation in the voltage-sensor discloses contributions of a conserved phenylalanine to gating properties of Kv1.1 channels and ataxia. <i>Scientific Reports</i> , 2017 , 7, 4583	4.9	7
6	Oxidative stress alters physiological and morphological neuronal properties. <i>Neurochemical Research</i> , 2007 , 32, 1169-78	4.6	7
5	Identification of a New Mutation Underlying Regressive Episodic Ataxia Type I. <i>Frontiers in Neurology</i> , 2018 , 9, 587	4.1	6
4	Altered functional properties of a missense variant in the TRESK K channel (KCNK18) associated with migraine and intellectual disability. <i>Pflugers Archiv European Journal of Physiology</i> , 2020 , 472, 923-930	4.6	4
3	Commentary: A channelopathy mutation in the voltage-sensor discloses contributions of a conserved phenylalanine to gating properties of Kv1.1 channels and ataxia. <i>Frontiers in Cellular Neuroscience</i> , 2018 , 12, 174	6.1	3
2	Electromechanical coupling of the Kv1.1 voltage-gated K channel is fine-tuned by the simplest amino acid residue in the S4-S5 linker. <i>Pflugers Archiv European Journal of Physiology</i> , 2020 , 472, 899-909	4.6	2
1	(Kir5.1) Gene Ablation Causes Subfertility and Increases the Prevalence of Morphologically Abnormal Spermatozoa. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	2