

# Sonia M Hasan

## List of Publications by Year in Descending Order

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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	(Kir5.1) Gene Ablation Causes Subfertility and Increases the Prevalence of Morphologically Abnormal Spermatozoa. <i>International Journal of Molecular Sciences</i> , <b>2021</b> , 22,	6.3	2
12	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> , <b>2020</b> , 472, 923-930	4.6	4
11	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> , <b>2020</b> , 472, 899-909	4.6	2
10	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> , <b>2018</b> , 12, 174	6.1	3
9	Identification of a New Mutation Underlying Regressive Episodic Ataxia Type I. <i>Frontiers in Neurology</i> , <b>2018</b> , 9, 587	4.1	6
8	Lethal digenic mutations in the K channels Kir4.1 () and SLACK () associated with severe-disabling seizures and neurodevelopmental delay. <i>Journal of Neurophysiology</i> , <b>2017</b> , 118, 2402-2411	3.2	12
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> , <b>2017</b> , 7, 4583	4.9	7
6	Gain-of-function defects of astrocytic Kir4.1 channels in children with autism spectrum disorders and epilepsy. <i>Scientific Reports</i> , <b>2016</b> , 6, 34325	4.9	43
5	A Calsequestrin-1 Mutation Associated with a Skeletal Muscle Disease Alters Sarcoplasmic Ca <sup>2+</sup> Release. <i>PLoS ONE</i> , <b>2016</b> , 11, e0155516	3.7	12
4	deficiency leads to a syndrome of abnormal striatum, congenital cataract and intellectual disability. <i>Journal of Medical Genetics</i> , <b>2016</b> , 53, 786-792	5.8	14
3	New insights into the pathogenesis and therapeutics of episodic ataxia type 1. <i>Frontiers in Cellular Neuroscience</i> , <b>2015</b> , 9, 317	6.1	40
2	Hydrogen peroxide-induced reduction of delayed rectifier potassium current in hippocampal neurons involves oxidation of sulfhydryl groups. <i>Brain Research</i> , <b>2013</b> , 1520, 61-9	3.7	7
1	Oxidative stress alters physiological and morphological neuronal properties. <i>Neurochemical Research</i> , <b>2007</b> , 32, 1169-78	4.6	7