

# Nyamkhishig Sambuughin

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5247598/publications.pdf>

Version: 2024-02-01

8  
papers

241  
citations

1478505

6  
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1720034

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8  
docs citations

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times ranked

481  
citing authors

#	ARTICLE	IF	CITATIONS
1	Multifactorial Origin of Exertional Rhabdomyolysis, Recurrent Hematuria, and Episodic Pain in a Service Member with Sickle Cell Trait. <i>Case Reports in Genetics</i> , 2018, 2018, 1-6.	0.2	4
2	Response to Finsterer and Zarrouk-Mahjoub. <i>Molecular Genetics and Metabolism Reports</i> , 2018, 17, 2.	1.1	0
3	Pathogenic and rare deleterious variants in multiple genes suggest oligogenic inheritance in recurrent exertional rhabdomyolysis. <i>Molecular Genetics and Metabolism Reports</i> , 2018, 16, 76-81.	1.1	15
4	Round Table on Malignant Hyperthermia in Physically Active Populations: Meeting Proceedings. <i>Journal of Athletic Training</i> , 2017, 52, 377-383.	1.8	16
5	Exome analysis identifies <scp>B</scp>rody myopathy in a family diagnosed with malignant hyperthermia susceptibility. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2014, 2, 472-483.	1.2	20
6	Exome sequencing reveals SCO2 mutations in a family presented with fatal infantile hyperthermia. <i>Journal of Human Genetics</i> , 2013, 58, 226-228.	2.3	9
7	KBTBD13 interacts with Cullin 3 to form a functional ubiquitin ligase. <i>Biochemical and Biophysical Research Communications</i> , 2012, 421, 743-749.	2.1	34
8	Dominant Mutations in KBTBD13, a Member of the BTB/Kelch Family, Cause Nemaline Myopathy with Cores. <i>American Journal of Human Genetics</i> , 2010, 87, 842-847.	6.2	143