

# 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

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citations

1478505

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1720034

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times ranked

481  
citing authors

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