

# Sadia

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/9707071/publications.pdf>

Version: 2024-02-01

7  
papers

58  
citations

2682572  
2  
h-index

1872680  
6  
g-index

7  
all docs

7  
docs citations

7  
times ranked

28  
citing authors

#	ARTICLE	IF	CITATIONS
1	A homozygous missense variant in the <i>MLC1</i> gene underlies megalencephalic leukoencephalopathy with subcortical cysts in large kindred: Heterozygous carriers show seizure and mild motor function deterioration. <i>American Journal of Medical Genetics, Part A</i> , 2022, 188, 1075-1082.	1.2	1
2	Identification of a Novel Homozygous Missense (c.443A>T;p.N148I) Mutation in BBS2 in a Kashmiri Family with Bardet-Biedl Syndrome. <i>BioMed Research International</i> , 2021, 2021, 1-9.	1.9	1
3	Minocycline-Derived Silver Nanoparticles for Assessment of Their Antidiabetic Potential against Alloxan-Induced Diabetic Mice. <i>Pharmaceutics</i> , 2021, 13, 1678.	4.5	6
4	Current status of beta-thalassemia and its treatment strategies. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2021, 9, e1788.	1.2	44
5	Sequencing and Characterization of Mitochondrial Protein-Coding Genes for <i>Schizothorax niger</i> (Cypriniformes: Cyprinidae) with Phylogenetic Consideration. <i>BioMed Research International</i> , 2020, 2020, 1-13.	1.9	2
6	Identification of a recurrent nonsense mutation in HR gene responsible for atrichia with papular lesions in two Kashmiri families. <i>Journal of Gene Medicine</i> , 2020, 22, e3167.	2.8	2
7	A recurrent missense mutation in the EDAR gene causes severe autosomal recessive hypohidrotic ectodermal dysplasia in two consanguineous Kashmiri families. <i>Journal of Gene Medicine</i> , 2019, 21, e3113.	2.8	2