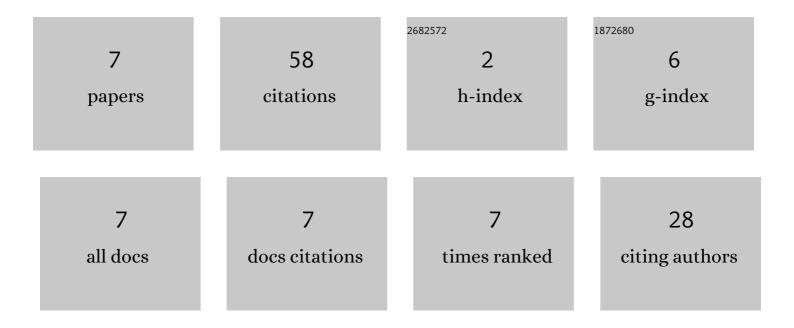


## List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/9707071/publications.pdf Version: 2024-02-01



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