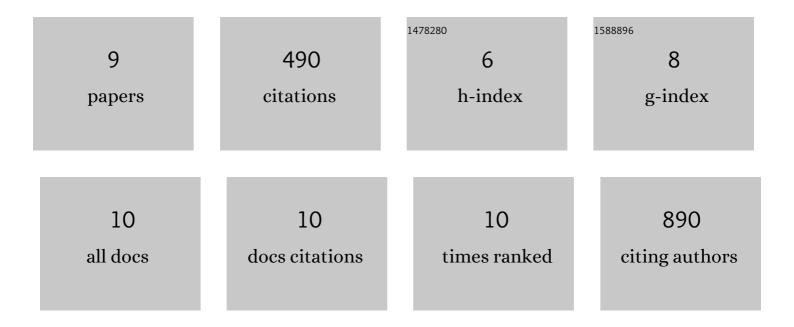
## Bhagyalaxmi Mohapatra

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/3181870/publications.pdf Version: 2024-02-01



#	Article	IF	CITATIONS
1	Dilated cardiomyopathy: a new insight into the rare but common cause of heart failure. Heart Failure Reviews, 2022, 27, 431-454.	1.7	12
2	A gain-of-function mutation in CITED2 is associated with congenital heart disease. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2021, 822, 111741.	0.4	7
3	Functional analysis of novel genetic variants of <i><scp>NKX2</scp>â€5</i> associated with nonsyndromic congenital heart disease. American Journal of Medical Genetics, Part A, 2021, 185, 3644-3663.	0.7	9
4	Implication of rare genetic variants of NODAL and ACVR1B in congenital heart disease patients from Indian population. Experimental Cell Research, 2021, 409, 112869.	1.2	1
5	Implication of GATA4 synonymous variants in congenital heart disease: A comprehensive in-silico approach. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2019, 813, 31-38.	0.4	9
6	Functionally significant, novel <i>GATA4</i> variants are frequently associated with Tetralogy of Fallot. Human Mutation, 2018, 39, 1957-1972.	1.1	9
7	Epidemiology of Congenital Heart Disease in India. Congenital Heart Disease, 2015, 10, 437-446.	0.0	54
8	Identification and functional characterization of NODAL rare variants in heterotaxy and isolated cardiovascular malformations. Human Molecular Genetics, 2009, 18, 861-871.	1.4	140
9	Mutations in the muscle LIM protein and α-actinin-2 genes in dilated cardiomyopathy and endocardial fibroelastosis. Molecular Genetics and Metabolism, 2003, 80, 207-215.	0.5	249