

Bhagyalaxmi Mohapatra

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/3181870/publications.pdf>

Version: 2024-02-01

9
papers

490
citations

1478280

6
h-index

1588896

8
g-index

10
all docs

10
docs citations

10
times ranked

890
citing authors

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