

Lili Hao

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5922360/publications.pdf>

Version: 2024-02-01

11
papers

119
citations

1306789

7
h-index

1281420

11
g-index

11
all docs

11
docs citations

11
times ranked

199
citing authors

#	ARTICLE	IF	CITATIONS
1	Multiple gene variations contributed to congenital heart disease via GATA family transcriptional regulation. <i>Journal of Translational Medicine</i> , 2017, 15, 69.	1.8	22
2	The roles of SMYD4 in epigenetic regulation of cardiac development in zebrafish. <i>PLoS Genetics</i> , 2018, 14, e1007578.	1.5	17
3	Early onset developmental delay and epilepsy in pediatric patients with WDR45 variants. <i>European Journal of Medical Genetics</i> , 2019, 62, 149-160.	0.7	16
4	Hypermethylation-mediated downregulation of lncRNA TBX5-AS1:2 in Tetralogy of Fallot inhibits cell proliferation by reducing TBX5 expression. <i>Journal of Cellular and Molecular Medicine</i> , 2020, 24, 6472-6484.	1.6	16
5	Hypomethylation and decreased expression of <i>BRG1</i> in the myocardium of patients with congenital heart disease. <i>Birth Defects Research</i> , 2017, 109, 1183-1195.	0.8	14
6	Exome sequencing identifies a mutation in TMC1 as a novel cause of autosomal recessive nonsyndromic hearing loss. <i>Journal of Translational Medicine</i> , 2016, 14, 29.	1.8	10
7	Two novel <i>ANK1</i> loss-of-function mutations in Chinese families with hereditary spherocytosis. <i>Journal of Cellular and Molecular Medicine</i> , 2019, 23, 4454-4463.	1.6	10
8	Long non-coding RNA SAP30-2:1 is downregulated in congenital heart disease and regulates cell proliferation by targeting HAND2. <i>Frontiers of Medicine</i> , 2021, 15, 91-100.	1.5	7
9	Expression Profiles of Exosomal MicroRNAs Derived from Cerebrospinal Fluid in Patients with Congenital Hydrocephalus Determined by MicroRNA Sequencing. <i>Disease Markers</i> , 2022, 2022, 1-16.	0.6	3
10	<i>WDR62</i> variants contribute to congenital heart disease by inhibiting cardiomyocyte proliferation. <i>Clinical and Translational Medicine</i> , 2022, 12, .	1.7	3
11	The Novel Pathogenic Mutation c.849dupT in <i>BRCA2</i> Contributes to the Nonsense-Mediated mRNA Decay of <i>BRCA2</i> in Familial Breast Cancer. <i>Journal of Breast Cancer</i> , 2018, 21, 330.	0.8	1