Hairui Sun

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/5204490/publications.pdf

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14	174	1478505 6	1125743 13
papers	citations	h-index	g-index

14	14	14	206
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Prognostic significance of IDH mutation in adult low-grade gliomas: a meta-analysis. Journal of Neuro-Oncology, 2013, 113, 277-284.	2.9	74
2	Increased frequency of FBN1 frameshift and nonsense mutations in Marfan syndrome patients with aortic dissection. Molecular Genetics & Genomic Medicine, 2020, 8, e1041.	1.2	21
3	Genetics and Clinical Features of Noncompaction Cardiomyopathy in the Fetal Population. Frontiers in Cardiovascular Medicine, 2020, 7, 617561.	2.4	18
4	Case Report: Characterization of a Novel NONO Intronic Mutation in a Fetus With X-Linked Syndromic Mental Retardation-34. Frontiers in Genetics, 2020, 11, 593688.	2.3	13
5	Characteristics of Cardiac Phenotype in Prenatal Familial Cases With NONO Mutations. Circulation Genomic and Precision Medicine, 2020, 13, e002847.	3.6	12
6	Expanding the phenotype associated with SMARCC2 variants: a fetus with tetralogy of Fallot. BMC Medical Genomics, 2022, 15, 40.	1.5	8
7	Genetic and Clinical Features of Heterotaxy in a Prenatal Cohort. Frontiers in Genetics, 2022, 13, 818241.	2.3	8
8	Expanding the phenotype of STRA6â€related disorder to include left ventricular nonâ€compaction. Molecular Genetics & Genomic Medicine, 2020, 8, e1377.	1.2	4
9	Diverse cardiac phenotypes among different carriers of the same MYH7 splicing variant allele (c.732+1G>A) from a family. BMC Medical Genomics, 2022, 15, 36.	1.5	4
10	A novel intron mutation in FBN-1 gene identified in a pregnant woman with Marfan syndrome. Hereditas, 2021, 158, 6.	1.4	3
11	Detection of TSC1/TSC2 mosaic variants in patients with cardiac rhabdomyoma and tuberous sclerosis complex by hybridâ€capture nextâ€generation sequencing. Molecular Genetics & Genomic Medicine, 2021, 9, e1802.	1.2	3
12	Case Report: Biventricular Noncompaction Cardiomyopathy With Pulmonary Stenosis and Bradycardia in a Fetus With KCNH2 Mutation. Frontiers in Genetics, 2022, 13, 821226.	2.3	3
13	Generation of a NONO homozygous knockout human induced pluripotent stem cell line by CRISPR/Cas9 editing. Stem Cell Research, 2020, 47, 101893.	0.7	2
14	Unique dual indexing PCR reduces chimeric contamination and improves mutation detection in cell-free DNA of pregnant women. Talanta, 2020, 217, 121035.	5.5	1