

Agnese Giovannetti

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/99429/publications.pdf>

Version: 2024-02-01

8
papers

69
citations

1937685

4
h-index

1720034

7
g-index

8
all docs

8
docs citations

8
times ranked

104
citing authors

#	ARTICLE	IF	CITATIONS
1	Landscapes of cellular phenotypic diversity in breast cancer xenografts and their impact on drug response. <i>Nature Communications</i> , 2021, 12, 1998.	12.8	37
2	Whole exome sequencing in an Italian family with isolated maxillary canine agenesis and canine eruption anomalies. <i>Archives of Oral Biology</i> , 2018, 91, 96-102.	1.8	8
3	GDF5 mutation case report and a systematic review of molecular and clinical spectrum: Expanding current knowledge on genotype-phenotype correlations. <i>Bone</i> , 2021, 144, 115803.	2.9	7
4	Prenatal whole exome sequencing detects a new homozygous fukutin (FKTN) mutation in a fetus with an ultrasound suspicion of familial Dandy-Walker malformation. <i>Molecular Genetics & Genomic Medicine</i> , 2020, 8, e1054.	1.2	6
5	TLR4 T399I Polymorphism and Endometriosis in a Cohort of Italian Women. <i>Diagnostics</i> , 2020, 10, 255.	2.6	6
6	Heterozygous nonsense <i>ARX</i> mutation in a family highlights the complexity of clinical and molecular diagnosis in case of chromosomal and single gene disorder coinheritance. <i>Molecular Genetics & Genomic Medicine</i> , 2020, 8, e1336.	1.2	4
7	MiRLog and dbmiR: Prioritization and functional annotation tools to study human microRNA sequence variants. <i>Human Mutation</i> , 2022, , .	2.5	1
8	Correlating Neuroimaging and CNVs Data: 7 Years of Cytogenomic Microarray Analysis on Patients Affected by Neurodevelopmental Disorders. <i>Journal of Pediatric Genetics</i> , 2021, 10, 292-299.	0.7	0