Jing

List of Publications by Year
in descending order

10 papers	79 citations	1683354 5 h-index	1473754 9 g-index
10 all docs	10 docs citations	10 times ranked	96 citing authors

#	Article	IF	Citations
1	STARD3NL inhibits the osteogenic differentiation by inactivating the Wnt/Î²â€catenin pathway via binding to Annexin A2 in osteoporosis. Journal of Cellular and Molecular Medicine, 2022, , .	1.6	1
2	Singleâ€cell transcriptomics reveals pathogenic dysregulation of previously unrecognised chondral stem/progenitor cells in children with microtia. Clinical and Translational Medicine, 2022, 12, e702.	1.7	1
3	Expression Profiles of Exosomal MicroRNAs Derived from Cerebrospinal Fluid in Patients with Congenital Hydrocephalus Determined by MicroRNA Sequencing. Disease Markers, 2022, 2022, 1-16.	0.6	3
4	Downregulation of MEG3 and upregulation of EZH2 cooperatively promote neuroblastoma progression. Journal of Cellular and Molecular Medicine, 2022, 26, 2377-2391.	1.6	18
5	WDR62 variants contribute to congenital heart disease by inhibiting cardiomyocyte proliferation. Clinical and Translational Medicine, 2022, 12, .	1.7	3
6	Hypermethylationâ€mediated downâ€regulation of lncRNA TBX5â€AS1:2 in Tetralogy of Fallot inhibits cell proliferation by reducingTBX5expression. Journal of Cellular and Molecular Medicine, 2020, 24, 6472-6484.	1.6	16
7	MAST1 modulates neuronal differentiation and cell cycle exit via P27 in neuroblastoma cells. FEBS Open Bio, 2020, 10, 1104-1114.	1.0	3
8	Mutations in <scp>TOP</scp>2B cause autosomalâ€dominant hereditary hearing loss via inhibition of the <scp>PI</scp>3Kâ€Akt signalling pathway. FEBS Letters, 2019, 593, 2008-2018.	1.3	13
9	Novel TRRAP mutation causes autosomal dominant nonâ€syndromic hearing loss. Clinical Genetics, 2019, 96, 300-308.	1.0	11
10	Two novel ANK1 lossâ€ofâ€function mutations in Chinese families with hereditary spherocytosis. Journal of Cellular and Molecular Medicine, 2019, 23, 4454-4463.	1.6	10