Harikiran Nistala

List of Publications by Year in descending order

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1040056 1372567 11 456 9 10 citations h-index g-index papers 12 12 12 844 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Fibrillin-1 and -2 differentially modulate endogenous TGF- \hat{l}^2 and BMP bioavailability during bone formation. Journal of Cell Biology, 2010, 190, 1107-1121.	5.2	173
2	Differential effects of alendronate and losartan therapy on osteopenia and aortic aneurysm in mice with severe Marfan syndrome. Human Molecular Genetics, 2010, 19, 4790-4798.	2.9	58
3	Extracellular Microfibrils Control Osteoblast-supported Osteoclastogenesis by Restricting TGFβ Stimulation of RANKL Production. Journal of Biological Chemistry, 2010, 285, 34126-34133.	3.4	49
4	Caffey disease: New perspectives on old questions. Bone, 2014, 60, 246-251.	2.9	45
5	Fibrillin-Rich Microfibrils—Structural and Instructive Determinants of Mammalian Development and Physiology. Connective Tissue Research, 2008, 49, 1-6.	2.3	35
6	Functional biology of the Steel syndrome founder allele and evidence for clan genomics derivation of COL27A1 pathogenic alleles worldwide. European Journal of Human Genetics, 2020, 28, 1243-1264.	2.8	27
7	Extracellular regulation of transforming growth factor \hat{l}^2 and bone morphogenetic protein signaling in bone. Annals of the New York Academy of Sciences, 2010, 1192, 253-256.	3.8	25
8	Generation of $\langle i \rangle$ Fbn1 $\langle i \rangle$ conditional null mice implicates the extracellular microfibrils in osteoprogenitor recruitment. Genesis, 2012, 50, 635-641.	1.6	19
9	NMIHBA results from hypomorphic <i>PRUNE1</i> variants that lack short-chain exopolyphosphatase activity. Human Molecular Genetics, 2021, 29, 3516-3531.	2.9	16
10	Drugâ€Based Therapies for Vascular Disease in Marfan Syndrome: From Mouse Models to Human Patients. Mount Sinai Journal of Medicine, 2010, 77, 366-373.	1.9	9
11	Differential effects of alendronate and losartan therapy on osteopenia and aortic aneurysm in mice with severe Marfan syndrome. Human Molecular Genetics, 2014, 23, 6137-6137.	2.9	0