

Megan Ealy

List of Publications by Year in descending order

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Version: 2024-02-01

16
papers

659
citations

623734

14
h-index

996975

15
g-index

16
all docs

16
docs citations

16
times ranked

754
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular characterization and prospective isolation of human fetal cochlear hair cell progenitors. <i>Nature Communications</i> , 2018, 9, 4027.	12.8	70
2	Single-cell analysis delineates a trajectory toward the human early otic lineage. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, 8508-8513.	7.1	60
3	Mutations in apoptosis-inducing factor cause X-linked recessive auditory neuropathy spectrum disorder. <i>Journal of Medical Genetics</i> , 2015, 52, 523-531.	3.2	92
4	Rare Variants in BMP2 and BMP4 Found in Otosclerosis Patients Reduce Smad Signaling. <i>Otology and Neurotology</i> , 2014, 35, 395-400.	1.3	10
5	<i>COL1A1</i> association and otosclerosis: A meta-analysis. <i>American Journal of Medical Genetics, Part A</i> , 2012, 158A, 1066-1070.	1.2	20
6	Otosclerosis. <i>Advances in Oto-Rhino-Laryngology</i> , 2011, 70, 122-129.	1.6	38
7	Commentary on "Otosclerosis: Thirty-Year Follow-Up After Surgery". <i>Annals of Otology, Rhinology and Laryngology</i> , 2011, 120, 615-616.	1.1	0
8	The prevalence of mitochondrial mutations associated with aminoglycoside-induced sensorineural hearing loss in an NICU population. <i>Laryngoscope</i> , 2011, 121, 1184-1186.	2.0	26
9	Genetic variants in the RELN gene are associated with otosclerosis in multiple European populations. <i>Human Genetics</i> , 2010, 127, 155-162.	3.8	28
10	Genetic variants in <i>RELN</i> are associated with otosclerosis in a non-European population from Tunisia. <i>Annals of Human Genetics</i> , 2010, 74, 399-405.	0.8	18
11	The Genetics of otosclerosis. <i>Hearing Research</i> , 2010, 266, 70-74.	2.0	44
12	A Genome-wide Analysis Identifies Genetic Variants in the RELN Gene Associated with Otosclerosis. <i>American Journal of Human Genetics</i> , 2009, 84, 328-338.	6.2	66
13	No Evidence for Association Between the Renin-Angiotensin-Aldosterone System and Otosclerosis in a Large Belgian-Dutch Population. <i>Otology and Neurotology</i> , 2009, 30, 1079-1083.	1.3	26
14	Association of Bone Morphogenetic Proteins With Otosclerosis. <i>Journal of Bone and Mineral Research</i> , 2008, 23, 507-516.	2.8	58
15	Gene expression analysis of human otosclerotic stapedial footplates. <i>Hearing Research</i> , 2008, 240, 80-86.	2.0	28
16	The coding polymorphism T263I in TGF- β 1 is associated with otosclerosis in two independent populations. <i>Human Molecular Genetics</i> , 2007, 16, 2021-2030.	2.9	75