Megan Ealy

List of Publications by Year
in descending order

16	659	623734 14	996975 15
papers	citations	h-index	g-index

16	16	16	754
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Mutations in apoptosis-inducing factor cause X-linked recessive auditory neuropathy spectrum disorder. Journal of Medical Genetics, 2015, 52, 523-531.	3.2	92
2	The coding polymorphism T263I in TGF-Î²1 is associated with otosclerosis in two independent populations. Human Molecular Genetics, 2007, 16, 2021-2030.	2.9	75
3	Molecular characterization and prospective isolation of human fetal cochlear hair cell progenitors. Nature Communications, 2018, 9, 4027.	12.8	70
4	A Genome-wide Analysis Identifies Genetic Variants in the RELN Gene Associated with Otosclerosis. American Journal of Human Genetics, 2009, 84, 328-338.	6.2	66
5	Single-cell analysis delineates a trajectory toward the human early otic lineage. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 8508-8513.	7.1	60
6	Association of Bone Morphogenetic Proteins With Otosclerosis. Journal of Bone and Mineral Research, 2008, 23, 507-516.	2.8	58
7	The Genetics of otosclerosis. Hearing Research, 2010, 266, 70-74.	2.0	44
8	Otosclerosis. Advances in Oto-Rhino-Laryngology, 2011, 70, 122-129.	1.6	38
9	Gene expression analysis of human otosclerotic stapedial footplates. Hearing Research, 2008, 240, 80-86.	2.0	28
10	Genetic variants in the RELN gene are associated with otosclerosis in multiple European populations. Human Genetics, 2010, 127, 155-162.	3.8	28
11	No Evidence for Association Between the Renin-Angiotensin-Aldosterone System and Otosclerosis in a Large Belgian-Dutch Population. Otology and Neurotology, 2009, 30, 1079-1083.	1.3	26
12	The prevalence of mitochondrial mutations associated with aminoglycosideâ€induced sensorineural hearing loss in an NICU population. Laryngoscope, 2011, 121, 1184-1186.	2.0	26
13	COL1A1 association and otosclerosis: A metaâ€analysis. American Journal of Medical Genetics, Part A, 2012, 158A, 1066-1070.	1.2	20
14	Genetic variants in RELN are associated with otosclerosis in a nonâ€European population from Tunisia. Annals of Human Genetics, 2010, 74, 399-405.	0.8	18
15	Rare Variants in BMP2 and BMP4 Found in Otosclerosis Patients Reduce Smad Signaling. Otology and Neurotology, 2014, 35, 395-400.	1.3	10
16	Commentary on â€œOtosclerosis: Thirty-Year Follow-Up After Surgeryâ€. Annals of Otology, Rhinology and Laryngology, 2011, 120, 615-616.	1.1	0