

Manou Sommen

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/981838/publications.pdf>

Version: 2024-02-01

8
papers

266
citations

1478505

6
h-index

1588992

8
g-index

8
all docs

8
docs citations

8
times ranked

521
citing authors

#	ARTICLE	IF	CITATIONS
1	A wide range of protective and predisposing variants in aggrecan influence the susceptibility for otosclerosis. <i>Human Genetics</i> , 2022, 141, 951-963.	3.8	6
2	Osmotic stress inhibits leaf growth of <i>Arabidopsis thaliana</i> by enhancing ARF-mediated auxin responses. <i>New Phytologist</i> , 2020, 226, 1766-1780.	7.3	31
3	Insufficient evidence for a role of SERPINF1 in otosclerosis. <i>Molecular Genetics and Genomics</i> , 2019, 294, 1001-1006.	2.1	11
4	Variants affecting diverse domains of MEPE are associated with two distinct bone disorders, a craniofacial bone defect and otosclerosis. <i>Genetics in Medicine</i> , 2019, 21, 1199-1208.	2.4	17
5	DNA Diagnostics of Hereditary Hearing Loss: A Targeted Resequencing Approach Combined with a Mutation Classification System. <i>Human Mutation</i> , 2016, 37, 812-819.	2.5	76
6	Heimler Syndrome Is Caused by Hypomorphic Mutations in the Peroxisome-Biogenesis Genes PEX1 and PEX6. <i>American Journal of Human Genetics</i> , 2015, 97, 535-545.	6.2	103
7	Genetic Association Analysis in a Clinically and Histologically Confirmed Otosclerosis Population Confirms Association With the TGFBI Gene but Suggests an Association of the RELN Gene With a Clinically Indistinguishable Otosclerosis-Like Phenotype. <i>Otology and Neurotology</i> , 2014, 35, 1058-1064.	1.3	17
8	Genetic and clinical diagnosis in non-syndromic hearing loss. <i>Hearing, Balance and Communication</i> , 2013, 11, 138-145.	0.4	5