Manou Sommen

List of Publications by Year
in descending order

8	266	1478505 6	1588992 8
papers	citations	h-index	g-index

8	8	8	521
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Heimler Syndrome Is Caused by Hypomorphic Mutations in the Peroxisome-Biogenesis Genes PEX1 and PEX6. American Journal of Human Genetics, 2015, 97, 535-545.	6.2	103
2	DNA Diagnostics of Hereditary Hearing Loss: A Targeted Resequencing Approach Combined with a Mutation Classification System. Human Mutation, 2016, 37, 812-819.	2.5	76
3	Osmotic stress inhibits leaf growth of Arabidopsis thaliana by enhancing ARFâ€mediated auxin responses. New Phytologist, 2020, 226, 1766-1780.	7.3	31
4	Genetic Association Analysis in a Clinically and Histologically Confirmed Otosclerosis Population Confirms Association With the TGFB1 Gene but Suggests an Association of the RELN Gene With a Clinically Indistinguishable Otosclerosis-Like Phenotype. Otology and Neurotology, 2014, 35, 1058-1064.	1.3	17
5	Variants affecting diverse domains of MEPE are associated with two distinct bone disorders, a craniofacial bone defect and otosclerosis. Genetics in Medicine, 2019, 21, 1199-1208.	2.4	17
6	Insufficient evidence for a role of SERPINF1 in otosclerosis. Molecular Genetics and Genomics, 2019, 294, 1001-1006.	2.1	11
7	A wide range of protective and predisposing variants in aggrecan influence the susceptibility for otosclerosis. Human Genetics, 2022, 141, 951-963.	3.8	6
8	Genetic and clinical diagnosis in non-syndromic hearing loss. Hearing, Balance and Communication, 2013, 11, 138-145.	0.4	5