Jeroen Jules Smits

List of Publications by Year
in descending order

9	95	1684188 5	1474206 9
papers	citations	h-index	g-index

12	12	12	189
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	De novo and inherited loss-of-function variants of ATP2B2 are associated with rapidly progressive hearing impairment. Human Genetics, 2019, 138, 61-72.	3.8	27
2	Cochlear supporting cells require GAS2 for cytoskeletal architecture and hearing. Developmental Cell, 2021, 56, 1526-1540.e7.	7.0	18
3	A RIPOR2 in-frame deletion is a frequent and highly penetrant cause of adult-onset hearing loss. Journal of Medical Genetics, 2021, 58, 96-104.	3.2	14
4	A Mid-scala Cochlear Implant Electrode Design Achieves a Stable Post-surgical Position in the Cochlea of Patients Over Timeâ€"A Prospective Observational Study. Otology and Neurotology, 2018, 39, e231-e239.	1.3	9
5	Revisiting Place-Pitch Match in CI Recipients Using 3D Imaging Analysis. Annals of Otology, Rhinology and Laryngology, 2016, 125, 378-384.	1.1	7
6	Variants in USP48 encoding ubiquitin hydrolase are associated with autosomal dominant non-syndromic hereditary hearing loss. Human Molecular Genetics, 2021, 30, 1785-1796.	2.9	6
7	Frequency and origin of the c.2090T>G p.(Leu697Trp) MYO3A variant associated with autosomal dominant hearing loss. European Journal of Human Genetics, 2022, 30, 13-21.	2.8	6
8	Genotype-Phenotype Correlations of Pathogenic COCH Variants in DFNA9: A HuGE Systematic Review and Audiometric Meta-Analysis. Biomolecules, 2022, 12, 220.	4.0	5
9	Exploring the missing heritability in subjects with hearing loss, enlarged vestibular aqueducts, and a single or no pathogenic SLC26A4 variant. Human Genetics, 2022, 141, 465-484.	3.8	3