# 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 | 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 |
| 2 | 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 |
| 3 | Genotype-Phenotype Correlations of Pathogenic COCH Variants in DFNA9: A HuGE Systematic Review and Audiometric Meta-Analysis. Biomolecules, 2022, 12, 220. | 4.0 | 5 |
| 4 | 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 |
| 5 | Cochlear supporting cells require GAS2 for cytoskeletal architecture and hearing. Developmental Cell, 2021, 56, 1526-1540.e7. | 7.0 | 18 |
| 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 | 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 |
| 8 | 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 |
| 9 | Revisiting Place-Pitch Match in CI Recipients Using 3D Imaging Analysis. Annals of Otology, Rhinology and Laryngology, 2016, 125, 378-384. | 1.1 | 7 |