Sawako Masuda

List of Publications by Year
in descending order

7 papers	111 citations	1684188 5 h-index	1720034 7 g-index
7 all docs	7 docs citations	7 times ranked	231 citing authors

#	Article	IF	Citations
1	Variants encoding a restricted carboxy-terminal domain of SLC12A2 cause hereditary hearing loss in humans. PLoS Genetics, 2020, 16, e1008643.	3.5	36
2	High prevalence of CDH23 mutations in patients with congenital high-frequency sporadic or recessively inherited hearing loss. Orphanet Journal of Rare Diseases, 2015, 10, 60.	2.7	34
3	Prevalence of TECTA mutation in patients with mid-frequency sensorineural hearing loss. Orphanet Journal of Rare Diseases, 2017, 12, 157.	2.7	19
4	Comparison of the prevalence and features of inner ear malformations in congenital unilateral and bilateral hearing loss. International Journal of Pediatric Otorhinolaryngology, 2019, 125, 92-97.	1.0	12
5	Phenotypeâ€“genotype correlation in patients with typical and atypical branchio-oto-renal syndrome. Scientific Reports, 2022, 12, 969.	3.3	6
6	Differences in hearing levels between siblings with hearing loss caused by GJB2 mutations. Auris Nasus Larynx, 2020, 47, 938-942.	1.2	2
7	Investigation of the hearing levels of siblings affected by a single GJB2 variant: Possibility of genetic modifiers. International Journal of Pediatric Otorhinolaryngology, 2021, 149, 110840.	1.0	2