Zheng-min Xu

List of Publications by Year
in descending order

14	152	1478505 6	1199594 12
papers	citations	h-index	g-index

16	16	16	212
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Systematic elucidation of the mechanism of geraniol via network pharmacology . Drug Design, Development and Therapy, 2019, Volume 13, 1069-1075.	4.3	56
2	The outcomes of endoscopic management in young children with subglottic stenosis. International Journal of Pediatric Otorhinolaryngology, 2017, 99, 141-145.	1.0	18
3	Performance of two hearing screening protocols in NICU in Shanghai. International Journal of Pediatric Otorhinolaryngology, 2011, 75, 1225-1229.	1.0	17
4	Prediction of frequency-specific hearing threshold using chirp auditory brainstem response in infants with hearing losses. International Journal of Pediatric Otorhinolaryngology, 2014, 78, 812-816.	1.0	15
5	Geraniol suppresses proinflammatory mediators in phorbol 12-myristate 13-acetate with A23187-induced HMC-1 cells. Drug Design, Development and Therapy, 2018, Volume 12, 2897-2903.	4.3	13
6	Identification of key genes in allergic rhinitis by bioinformatics analysis. Journal of International Medical Research, 2021, 49, 030006052110295.	1.0	10
7	The homozygote p.V27I/p.E114G variant of GJB2 is a putative indicator of nonsyndromic hearing loss in Chinese infants. International Journal of Pediatric Otorhinolaryngology, 2016, 84, 48-51.	1.0	6
8	Measurement of Thresholds Using Auditory Steady-State Response and Cochlear Microphonics in Children with Auditory Neuropathy. Journal of the American Academy of Audiology, 2019, 30, 672-676.	0.7	4
9	Auditory and Speech Outcomes of Cochlear Implantation in Children With Auditory Neuropathy Spectrum Disorder: A Systematic Review and Meta-Analysis. Annals of Otology, Rhinology and Laryngology, 2023, 132, 371-380.	1.1	4
10	Novel heterozygous GATA3 and SLC34A3 variants in a 6â€yearâ€old boy with Barakat syndrome and hypercalciuria. Molecular Genetics & Genomic Medicine, 2020, 8, e1222.	1.2	3
11	Apparent homozygosity for a novel splicing variant in EPS8 causes congenital profound hearing loss. European Journal of Medical Genetics, 2021, 64, 104362.	1.3	3
12	Prevalence of p.V37I variant of GJB2 among Chinese infants with mild or moderate hearing loss. International Journal of Clinical and Experimental Medicine, 2015, 8, 21674-8.	1.3	2
13	Recessive LOXHD1 variants cause a prelingual down-sloping hearing loss: genotype-phenotype correlation and three additional children with novel variants. International Journal of Pediatric Otorhinolaryngology, 2021, 145, 110715.	1.0	1
14	Imaging and audiological features of children with cochlear nerve deficiency. Ear, Nose and Throat Journal, 2022, , 014556132210966.	0.8	0