Cong Tian

List of Publications by Year
in descending order

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15 papers	358 citations	759055 12 h-index	996849 15 g-index
15 all docs	15 docs citations	15 times ranked	733 citing authors

#	Article	IF	Citations
1	Loss of the E3 ubiquitin ligase LRSAM1 sensitizes peripheral axons to degeneration in a mouse model of Charcot-Marie-Tooth disease. DMM Disease Models and Mechanisms, 2013, 6, 780-92.	1.2	44
2	Role for Toll-Like Receptor 2 in the Immune Response toStreptococcus pneumoniaeInfection in Mouse Otitis Media. Infection and Immunity, 2009, 77, 3100-3108.	1.0	38
3	A new mouse mutant of the Cdh23 gene with early-onset hearing loss facilitates evaluation of otoprotection drugs. Pharmacogenomics Journal, 2012, 12, 30-44.	0.9	37
4	The roles of USH1 proteins and PDZ domain-containing USH proteins in USH2 complex integrity in cochlear hair cells. Human Molecular Genetics, 2017, 26, ddw421.	1.4	35
5	Otitis media in a mouse model for Down syndrome. International Journal of Experimental Pathology, 2009, 90, 480-488.	0.6	28
6	Sh3pxd2b Mice Are a Model for Craniofacial Dysmorphology and Otitis Media. PLoS ONE, 2011, 6, e22622.	1.1	26
7	Ush1c gene expression levels in the ear and eye suggest different roles for Ush1c in neurosensory organs in a new Ush1c knockout mouse. Brain Research, 2010, 1328, 57-70.	1.1	24
8	Otitis Media in a New Mouse Model for CHARGE Syndrome with a Deletion in the Chd7 Gene. PLoS ONE, 2012, 7, e34944.	1.1	23
9	Pathological Features in the Lmna Mutant Mouse Provide a Novel Model of Human Otitis Media and Laminopathies. American Journal of Pathology, 2012, 181, 761-774.	1.9	20
10	A novel long intergenic non-coding RNA, Nostrill, regulates iNOS gene transcription and neurotoxicity in microglia. Journal of Neuroinflammation, 2021, 18, 16.	3.1	18
11	Ectopic Mineralization and Conductive Hearing Loss in Enpp1asj Mutant Mice, a New Model for Otitis Media and Tympanosclerosis. PLoS ONE, 2016, 11, e0168159.	1.1	18
12	Hearing loss without overt metabolic acidosis in ATP6V1B1 deficient MRL mice, a new genetic model for non-syndromic deafness with enlarged vestibular aqueducts. Human Molecular Genetics, 2017, 26, 3722-3735.	1.4	17
13	Detecting Novel Ototoxins and Potentiation of Ototoxicity by Disease Settings. Frontiers in Neurology, 2021, 12, 725566.	1.1	14
14	Mutation in Phex Gene Predisposes BALB/c-PhexHyp-Duk/Y Mice to Otitis Media. PLoS ONE, 2012, 7, e43010.	1.1	13
15	CACHD1-deficient mice exhibit hearing and balance deficits associated with a disruption of calcium homeostasis in the inner ear. Hearing Research, 2021, 409, 108327.	0.9	3