

Zheng-Yi Chen

List of Publications by Year in descending order

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Version: 2024-02-01

34
papers

3,226
citations

331670

21
h-index

377865

34
g-index

36
all docs

36
docs citations

36
times ranked

4837
citing authors

#	ARTICLE	IF	CITATIONS
1	Gene editing in a Myo6 semi-dominant mouse model rescues auditory function. <i>Molecular Therapy</i> , 2022, 30, 105-118.	8.2	31
2	Preventing autosomal-dominant hearing loss in Bth mice with CRISPR/CasRx-based RNA editing. <i>Signal Transduction and Targeted Therapy</i> , 2022, 7, 79.	17.1	22
3	Stem Cells and Gene Therapy in Progressive Hearing Loss: the State of the Art. <i>JARO - Journal of the Association for Research in Otolaryngology</i> , 2021, 22, 95-105.	1.8	15
4	Generation and characterization of a <i>P2rx2</i> V60L mouse model for DFNA41. <i>Human Molecular Genetics</i> , 2021, 30, 985-995.	2.9	4
5	The NIH Somatic Cell Genome Editing program. <i>Nature</i> , 2021, 592, 195-204.	27.8	84
6	Generation and Genetic Correction of USH2A c.2299delG Mutation in Patient-Derived Induced Pluripotent Stem Cells. <i>Genes</i> , 2021, 12, 805.	2.4	16
7	A Novel in vitro Model Delineating Hair Cell Regeneration and Neural Reinnervation in Adult Mouse Cochlea. <i>Frontiers in Molecular Neuroscience</i> , 2021, 14, 757831.	2.9	1
8	Direct Functional Protein Delivery with a Peptide into Neonatal and Adult Mammalian Inner Ear In Vivo. <i>Molecular Therapy - Methods and Clinical Development</i> , 2020, 18, 511-519.	4.1	5
9	Characterization of UMI028-A-1 stem cell line that contains a CRISPR/Cas9 induced hearing loss-associated variant (V60L (c.178G>A)) in the P2RX2 gene. <i>Stem Cell Research</i> , 2020, 49, 102017.	0.7	1
10	Ex vivo cell-based CRISPR/Cas9 genome editing for therapeutic applications. <i>Biomaterials</i> , 2020, 234, 119711.	11.4	58
11	Genome and base editing for genetic hearing loss. <i>Hearing Research</i> , 2020, 394, 107958.	2.0	18
12	Renewed proliferation in adult mouse cochlea and regeneration of hair cells. <i>Nature Communications</i> , 2019, 10, 5530.	12.8	71
13	In Vivo Assessment of Potential Therapeutic Approaches for USH2A-Associated Diseases. <i>Advances in Experimental Medicine and Biology</i> , 2019, 1185, 91-96.	1.6	26
14	Treatment of autosomal dominant hearing loss by in vivo delivery of genome editing agents. <i>Nature</i> , 2018, 553, 217-221.	27.8	412
15	Delivery of Adeno-Associated Virus Vectors in Adult Mammalian Inner-Ear Cell Subtypes Without Auditory Dysfunction. <i>Human Gene Therapy</i> , 2018, 29, 492-506.	2.7	64
16	The Key Transcription Factor Expression in the Developing Vestibular and Auditory Sensory Organs: A Comprehensive Comparison of Spatial and Temporal Patterns. <i>Neural Plasticity</i> , 2018, 2018, 1-9.	2.2	8
17	Creation of miniature pig model of human Waardenburg syndrome type 2A by ENU mutagenesis. <i>Human Genetics</i> , 2017, 136, 1463-1475.	3.8	28
18	Adenovirus Vectors Target Several Cell Subtypes of Mammalian Inner Ear In Vivo. <i>Neural Plasticity</i> , 2016, 2016, 1-8.	2.2	26

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19	Identification of Adeno-Associated Viral Vectors That Target Neonatal and Adult Mammalian Inner Ear Cell Subtypes. <i>Human Gene Therapy</i> , 2016, 27, 687-699.	2.7	79
20	Biomedical applications of gene editing. <i>Human Genetics</i> , 2016, 135, 967-969.	3.8	1
21	Myc and Fgf Are Required for Zebrafish Neuromast Hair Cell Regeneration. <i>PLoS ONE</i> , 2016, 11, e0157768.	2.5	22
22	The application of genome editing in studying hearing loss. <i>Hearing Research</i> , 2015, 327, 102-108.	2.0	46
23	Notch inhibition induces mitotically generated hair cells in mammalian cochleae via activating the Wnt pathway. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2015, 112, 166-171.	7.1	182
24	Gene Expression by Mouse Inner Ear Hair Cells during Development. <i>Journal of Neuroscience</i> , 2015, 35, 6366-6380.	3.6	308
25	Discovery and Characterization of a Peptide That Enhances Endosomal Escape of Delivered Proteins in Vitro and in Vivo. <i>Journal of the American Chemical Society</i> , 2015, 137, 14084-14093.	13.7	109
26	XIRP2, an Actin-Binding Protein Essential for Inner Ear Hair-Cell Stereocilia. <i>Cell Reports</i> , 2015, 10, 1811-1818.	6.4	32
27	Cationic lipid-mediated delivery of proteins enables efficient protein-based genome editing in vitro and in vivo. <i>Nature Biotechnology</i> , 2015, 33, 73-80.	17.5	1,180
28	Hair Cell Overexpression of Islet1 Reduces Age-Related and Noise-Induced Hearing Loss. <i>Journal of Neuroscience</i> , 2013, 33, 15086-15094.	3.6	41
29	Disrupting the Interaction between Retinoblastoma Protein and Raf-1 Leads to Defects in Progenitor Cell Proliferation and Survival during Early Inner Ear Development. <i>PLoS ONE</i> , 2013, 8, e83726.	2.5	4
30	Overlapping and distinct pRb pathways in the mammalian auditory and vestibular organs. <i>Cell Cycle</i> , 2011, 10, 337-351.	2.6	29
31	Diverse expression patterns of LIM-homeodomain transcription factors (LIM-HDs) in mammalian inner ear development. <i>Developmental Dynamics</i> , 2008, 237, 3305-3312.	1.8	33
32	The $\alpha 1$ subunit of nicotinic acetylcholine receptors in the inner ear: transcriptional regulation by ATOH1 and co-expression with the $\beta 3$ subunit in hair cells. <i>Journal of Neurochemistry</i> , 2007, 103, 2651-2664.	3.9	27
33	Proliferation of Functional Hair Cells in Vivo in the Absence of the Retinoblastoma Protein. <i>Science</i> , 2005, 307, 1114-1118.	12.6	240
34	Applications of genomics in the inner ear. <i>Pharmacogenomics</i> , 2003, 4, 735-745.	1.3	3