Matthew R Avenarius

List of Publications by Year
in descending order

25 papers	1,489 citations	516710 16 h-index	580821 25 g-index
28 all docs	28 docs citations	28 times ranked	2029 citing authors

#	Article	IF	Citations
1	<scp>EGFR</scp> internal tandem duplications in fusionâ€negative congenital and neonatal spindle cell tumors. Genes Chromosomes and Cancer, 2023, 62, 17-26.	2.8	3
2	The clinical utility of a riskâ€modifying <scp>SNP</scp> to detect carriers for spinal muscular atrophy with increased sensitivity. Molecular Genetics & Genomic Medicine, 2022, 10, e1897.	1.2	2
3	Loss of Baiap2l2 destabilizes the transducing stereocilia of cochlear hair cells and leads to deafness. Journal of Physiology, 2021, 599, 1173-1198.	2.9	28
4	Activation of the RAS pathway through uncommon BRAF mutations in mucinous pancreatic cysts without KRAS mutation. Modern Pathology, 2021, 34, 438-444.	5.5	19
5	Somatic PIK3R1 variation as a cause of vascular malformations and overgrowth. Genetics in Medicine, 2021, 23, 1882-1888.	2.4	26
6	Normal FISH CLL Represents a Heterogeneous Subgroup Where Prognosis Can be Refined with IGHV Mutational Status. Blood, 2021, 138, 1563-1563.	1.4	0
7	Genetic Characterization of Pediatric Sarcomas by Targeted RNA Sequencing. Journal of Molecular Diagnostics, 2020, 22, 1238-1245.	2.8	9
8	Complete sequencing of the SMN2 gene in SMA patients detects SMN gene deletion junctions and variants in SMN2 that modify the SMA phenotype. Human Genetics, 2019, 138, 241-256.	3.8	57
9	Genome sequencing identifies somatic BRAF duplication c.1794_1796dupTAC;p.Thr599dup in pediatric patient with low-grade ganglioglioma. Journal of Physical Education and Sports Management, 2018, 4, a002618.	1.2	7
10	Grxcr2 is required for stereocilia morphogenesis in the cochlea. PLoS ONE, 2018, 13, e0201713.	2.5	11
11	TRPV6, TRPM6 and TRPM7 Do Not Contribute to Hair-Cell Mechanotransduction. Frontiers in Cellular Neuroscience, 2018, 12, 41.	3.7	6
12	Heterodimeric capping protein is required for stereocilia length and width regulation. Journal of Cell Biology, 2017, 216, 3861-3881.	5.2	48
13	Stereocilia-staircase spacing is influenced by myosin III motors and their cargos espin-1 and espin-like. Nature Communications, 2016, 7, 10833.	12.8	72
14	Correlation of Actin Crosslinker and Capper Expression Levels with Stereocilia Growth Phases. Molecular and Cellular Proteomics, 2014, 13, 606-620.	3.8	26
15	Improved Biolistic Transfection of Hair Cells. PLoS ONE, 2012, 7, e46765.	2.5	14
16	Two Iranian families with a novel mutation in GJB2 causing autosomal dominant nonsyndromic hearing loss. American Journal of Medical Genetics, Part A, 2011, 155, 1202-1211.	1.2	9
17	Mutations in Grxcr1 Are The Basis for Inner Ear Dysfunction in the Pirouette Mouse. American Journal of Human Genetics, 2010, 86, 148-160.	6.2	49
18	Genetic male infertility and mutation of CATSPER ion channels. European Journal of Human Genetics, 2010, 18, 1178-1184.	2.8	139

#	Article	IF	Citations
19	Human Male Infertility Caused by Mutations in the CATSPER1 Channel Protein. American Journal of Human Genetics, 2009, 84, 505-510.	6.2	206
20	A Forward Genetics Screen in Mice Identifies Recessive Deafness Traits and Reveals That Pejvakin Is Essential for Outer Hair Cell Function. Journal of Neuroscience, 2007, 27, 2163-2175.	3.6	159
21	Evidence for a direct role of the disease modifier SCNM1 in splicing. Human Molecular Genetics, 2007, 16, 2506-2516.	2.9	41
22	The Coxsackievirus and Adenovirus Receptor: A new adhesion protein in cochlear development. Hearing Research, 2006, 215, 1-9.	2.0	28
23	GJB2 mutations: Passage through Iran. American Journal of Medical Genetics, Part A, 2005, 133A, 132-137.	1.2	77
24	A novel deletion involving the connexin-30 gene, del(GJB6-d13s1854), found in trans with mutations in the GJB2 gene (connexin-26) in subjects with DFNB1 non-syndromic hearing impairment. Journal of Medical Genetics, 2005, 42, 588-594.	3.2	282
25	OTOF mutations revealed by genetic analysis of hearing loss families including a potential temperature sensitive auditory neuropathy allele. Journal of Medical Genetics, 2005, 43, 576-581.	3.2	139