Steve D M Brown

List of Publications by Year
in descending order

64 papers	4,554 citations	172457 29 h-index	138484 58 g-index
65 all docs	65 docs citations	65 times ranked	8553 citing authors

#	Article	IF	Citations
1	Identifying genetic determinants of inflammatory pain in mice using a large-scale gene-targeted screen. Pain, 2022, 163, 1139-1157.	4.2	4
2	Neuroplastin genetically interacts with Cadherin 23 and the encoded isoform Np55 is sufficient for cochlear hair cell function and hearing. PLoS Genetics, 2022, 18, e1009937.	3.5	4
3	Introduction to Mammalian Genome Special Issue: Mammalian Genetic Resources. Mammalian Genome, 2022, 33, 1-3.	2.2	0
4	Extensive identification of genes involved in congenital and structural heart disorders and cardiomyopathy. , 2022, 1, 157-173.		22
5	Biophysical and morphological changes in inner hair cells and their efferent innervation in the ageing mouse cochlea. Journal of Physiology, 2021, 599, 269-287.	2.9	25
6	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
7	A resource of targeted mutant mouse lines for 5,061 genes. Nature Genetics, 2021, 53, 416-419.	21.4	60
8	Advances in mouse genetics for the study of human disease. Human Molecular Genetics, 2021, 30, R274-R284.	2.9	26
9	INFRAFRONTIER quality principles in systemic phenotyping. Mammalian Genome, 2021, , 1.	2.2	3
10	A holistic view of mouse enhancer architectures reveals analogous pleiotropic effects and correlation with human disease. BMC Genomics, 2020, 21, 754.	2.8	3
11	Pathophysiological changes in inner hair cell ribbon synapses in the ageing mammalian cochlea. Journal of Physiology, 2020, 598, 4339-4355.	2.9	23
12	The Jeff Mouse Mutant Model for Chronic Otitis Media Manifests Gain-of-Function as Well as Loss-of-Function Effects. Frontiers in Genetics, 2020, 11, 498.	2.3	2
13	Mutation in Fbxo11 Leads to Altered Immune Cell Content in Jeff Mouse Model of Otitis Media. Frontiers in Genetics, 2020, 11, 50.	2.3	6
14	Ageâ€related changes in the biophysical and morphological characteristics of mouse cochlear outer hair cells. Journal of Physiology, 2020, 598, 3891-3910.	2.9	29
15	High-throughput discovery of genetic determinants of circadian misalignment. PLoS Genetics, 2020, 16, e1008577.	3.5	10
16	Mouse mutant phenotyping at scale reveals novel genes controlling bone mineral density. PLoS Genetics, 2020, 16, e1009190.	3.5	19
17	High-throughput discovery of genetic determinants of circadian misalignment. , 2020, 16, e1008577.		0
18	High-throughput discovery of genetic determinants of circadian misalignment. , 2020, 16, e1008577.		0

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19	High-throughput discovery of genetic determinants of circadian misalignment. , 2020, 16, e1008577.		0
20	High-throughput discovery of genetic determinants of circadian misalignment. , 2020, 16, e1008577.		0
21	Modification of an aggressive model of Alport Syndrome reveals early differences in disease pathogenesis due to genetic background. Scientific Reports, 2019, 9, 20398.	3.3	11
22	Cellular content plays a crucial role in Nonâ€typeableHaemophilus influenzaeinfection of preinflamedJunbomouse middle ear. Cellular Microbiology, 2019, 21, e12960.	2.1	13
23	Transcript Analysis Reveals a Hypoxic Inflammatory Environment in Human Chronic Otitis Media With Effusion. Frontiers in Genetics, 2019, 10, 1327.	2.3	12
24	Cellular Immune Response against Nontypeable Haemophilus influenzae Infecting the Preinflamed Middle Ear of the Junbo Mouse. Infection and Immunity, 2019, 87, .	2.2	9
25	A genetic modifier suggests that endurance exercise exacerbates Huntington's disease. Human Molecular Genetics, 2018, 27, 1723-1731.	2.9	17
26	High-throughput mouse phenomics for characterizing mammalian gene function. Nature Reviews Genetics, 2018, 19, 357-370.	16.3	78
27	Helios is a key transcriptional regulator of outer hair cell maturation. Nature, 2018, 563, 696-700.	27.8	90
28	A Wars2 Mutant Mouse Model Displays OXPHOS Deficiencies and Activation of Tissue-Specific Stress Response Pathways. Cell Reports, 2018, 25, 3315-3328.e6.	6.4	35
29	A Requirement for Zic2 in the Regulation of Nodal Expression Underlies the Establishment of Left-Sided Identity. Scientific Reports, 2018, 8, 10439.	3.3	6
30	Genetic landscape of auditory dysfunction. Human Molecular Genetics, 2018, 27, R130-R135.	2.9	20
31	A large scale hearing loss screen reveals an extensive unexplored genetic landscape for auditory dysfunction. Nature Communications, 2017, 8, 886.	12.8	116
32	Prevalence of sexual dimorphism in mammalian phenotypic traits. Nature Communications, 2017, 8, 15475.	12.8	200
33	Disease model discovery from 3,328 gene knockouts by The International Mouse Phenotyping Consortium. Nature Genetics, 2017, 49, 1231-1238.	21.4	216
34	CIB2 interacts with TMC1 and TMC2 and is essential for mechanotransduction in auditory hair cells. Nature Communications, 2017, 8, 43.	12.8	121
35	A mutation in Nischarin causes otitis media via LIMK1 and NF-ÎºB pathways. PLoS Genetics, 2017, 13, e1006969.	3.5	36
36	Genome-wide association of multiple complex traits in outbred mice by ultra-low-coverage sequencing. Nature Genetics, 2016, 48, 912-918.	21.4	124

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37	Novel gene function revealed by mouse mutagenesis screens for models of age-related disease. Nature Communications, 2016, 7, 12444.	12.8	79
38	Mouse Models of NMNAT1-Leber Congenital Amaurosis (LCA9) Recapitulate Key Features of the Human Disease. American Journal of Pathology, 2016, 186, 1925-1938.	3.8	61
39	Absence of Neuroplastin-65 Affects Synaptogenesis in Mouse Inner Hair Cells and Causes Profound Hearing Loss. Journal of Neuroscience, 2016, 36, 222-234.	3.6	30
40	High-throughput discovery of novel developmental phenotypes. Nature, 2016, 537, 508-514.	27.8	1,001
41	Correction of the auditory phenotype in C57BL/6N mice via CRISPR/Cas9-mediated homology directed repair. Genome Medicine, 2016, 8, 16.	8.2	113
42	Mice with an N-Ethyl-N-Nitrosourea (ENU) Induced Tyr209Asn Mutation in Natriuretic Peptide Receptor 3 (NPR3) Provide a Model for Kyphosis Associated with Activation of the MAPK Signaling Pathway. PLoS ONE, 2016, 11, e0167916.	2.5	11
43	Interactions between the otitis media gene, Fbxo11, and p53 in the mouse embryonic lung. DMM Disease Models and Mechanisms, 2015, 8, 1531-42.	2.4	19
44	Comparative visualization of genotype-phenotype relationships. Nature Methods, 2015, 12, 698-699.	19.0	2
45	Analysis of mammalian gene function through broad-based phenotypic screens across a consortium of mouse clinics. Nature Genetics, 2015, 47, 969-978.	21.4	137
46	Applying the ARRIVE Guidelines to an In Vivo Database. PLoS Biology, 2015, 13, e1002151.	5.6	75
47	N-ethyl-N-Nitrosourea (ENU) Induced Mutations within the Klotho Gene Lead to Ectopic Calcification and Reduced Lifespan in Mouse Models. PLoS ONE, 2015, 10, e0122650.	2.5	16
48	The Actin-Binding Proteins Eps8 and Gelsolin Have Complementary Roles in Regulating the Growth and Stability of Mechanosensory Hair Bundles of Mammalian Cochlear Outer Hair Cells. PLoS ONE, 2014, 9, e87331.	2.5	15
49	A comparative phenotypic and genomic analysis of C57BL/6J and C57BL/6N mouse strains. Genome Biology, 2013, 14, R82.	9.6	403
50	Otitis media in the Tgif knockout mouse implicates TGFÎ² signalling in chronic middle ear inflammatory disease. Human Molecular Genetics, 2013, 22, 2553-2565.	2.9	50
51	EVI1 Acts as an Inducible Negative-Feedback Regulator of NF-ÎºB by Inhibiting p65 Acetylation. Journal of Immunology, 2012, 188, 6371-6380.	0.8	33
52	Unraveling the genetics of otitis media: from mouse to human and back again. Mammalian Genome, 2011, 22, 66-82.	2.2	58
53	Introduction. Mammalian Genome, 2011, 22, 361-361.	2.2	1
54	HIFâ€“VEGF Pathways Are Critical for Chronic Otitis Media in Junbo and Jeff Mouse Mutants. PLoS Genetics, 2011, 7, e1002336.	3.5	54

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55	Genome-Wide Association Study Using Extreme Truncate Selection Identifies Novel Genes Affecting Bone Mineral Density and Fracture Risk. PLoS Genetics, 2011, 7, e1001372.	3.5	233
56	A hearing and vestibular phenotyping pipeline to identify mouse mutants with hearing impairment. Nature Protocols, 2010, 5, 177-190.	12.0	93
57	Gelsolin Plays a Role in the Actin Polymerization Complex of Hair Cell Stereocilia. PLoS ONE, 2010, 5, e11627.	2.5	38
58	The Functional Annotation of Mammalian Genomes: The Challenge of Phenotyping. Annual Review of Genetics, 2009, 43, 305-333.	7.6	60
59	Mutation at the Evi1 Locus in Junbo Mice Causes Susceptibility to Otitis Media. PLoS Genetics, 2006, 2, e149.	3.5	88
60	Association of the FBXO11 Gene With Chronic Otitis Media With Effusion and Recurrent Otitis Media. JAMA Otolaryngology, 2006, 132, 729.	1.2	65
61	A mutation in the F-box gene, Fbxo11, causes otitis media in the Jeff mouse. Human Molecular Genetics, 2006, 15, 3273-3279.	2.9	98
62	A gene-driven ENU-based approach to generating an allelic series in any gene. Mammalian Genome, 2004, 15, 585-591.	2.2	148
63	The Deaf Mouse Mutant Jeff (Jf) is a Single Gene Model of Otitis Media. JARO - Journal of the Association for Research in Otolaryngology, 2003, 4, 130-138.	1.8	79
64	Novel ENU-induced eye mutations in the mouse: models for human eye disease. Human Molecular Genetics, 2002, 11, 755-767.	2.9	126