

Colin McKerlie

List of Publications by Year in descending order

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Version: 2024-02-01

33
papers

2,846
citations

471509

17
h-index

434195

31
g-index

40
all docs

40
docs citations

40
times ranked

6084
citing authors

#	ARTICLE	IF	CITATIONS
1	Identifying genetic determinants of inflammatory pain in mice using a large-scale gene-targeted screen. <i>Pain</i> , 2022, 163, 1139-1157.	4.2	4
2	Extensive identification of genes involved in congenital and structural heart disorders and cardiomyopathy. , 2022, 1, 157-173.		22
3	Process and Workflow for Preparation of Disparate Mouse Tissues for Proteomic Analysis. <i>Journal of Proteome Research</i> , 2021, 20, 305-316.	3.7	3
4	A resource of targeted mutant mouse lines for 5,061 genes. <i>Nature Genetics</i> , 2021, 53, 416-419.	21.4	60
5	Proteotyping of knockout mouse strains reveals sex- and strain-specific signatures in blood plasma. <i>Npj Systems Biology and Applications</i> , 2021, 7, 25.	3.0	2
6	INFRAFRONTIER quality principles in systemic phenotyping. <i>Mammalian Genome</i> , 2021, , 1.	2.2	3
7	Soft windowing application to improve analysis of high-throughput phenotyping data. <i>Bioinformatics</i> , 2020, 36, 1492-1500.	4.1	9
8	Variant in NHLRC2 leads to increased hnRNP C2 in developing neurons and the hippocampus of a mouse model of FINCA disease. <i>Molecular Medicine</i> , 2020, 26, 123.	4.4	5
9	PATHBIO: an international training program for precision mouse phenotyping. <i>Mammalian Genome</i> , 2020, 31, 49-53.	2.2	2
10	The Deep Genome Project. <i>Genome Biology</i> , 2020, 21, 18.	8.8	30
11	Human and mouse essentiality screens as a resource for disease gene discovery. <i>Nature Communications</i> , 2020, 11, 655.	12.8	64
12	Mouse mutant phenotyping at scale reveals novel genes controlling bone mineral density. <i>PLoS Genetics</i> , 2020, 16, e1009190.	3.5	19
13	Genome-wide screening of mouse knockouts reveals novel genes required for normal integumentary and oculocutaneous structure and function. <i>Scientific Reports</i> , 2019, 9, 11211.	3.3	6
14	A Comprehensive Plasma Metabolomics Dataset for a Cohort of Mouse Knockouts within the International Mouse Phenotyping Consortium. <i>Metabolites</i> , 2019, 9, 101.	2.9	40
15	Identification of genetic elements in metabolism by high-throughput mouse phenotyping. <i>Nature Communications</i> , 2018, 9, 288.	12.8	59
16	Identification of genes required for eye development by high-throughput screening of mouse knockouts. <i>Communications Biology</i> , 2018, 1, 236.	4.4	37
17	A Review of Current Standards and the Evolution of Histopathology Nomenclature for Laboratory Animals. <i>ILAR Journal</i> , 2018, 59, 29-39.	1.8	15
18	The International Mouse Phenotyping Consortium (IMPC): a functional catalogue of the mammalian genome that informs conservation. <i>Conservation Genetics</i> , 2018, 19, 995-1005.	1.5	82

#	ARTICLE	IF	CITATIONS
19	A large scale hearing loss screen reveals an extensive unexplored genetic landscape for auditory dysfunction. <i>Nature Communications</i> , 2017, 8, 886.	12.8	116
20	Prevalence of sexual dimorphism in mammalian phenotypic traits. <i>Nature Communications</i> , 2017, 8, 15475.	12.8	200
21	Disease model discovery from 3,328 gene knockouts by The International Mouse Phenotyping Consortium. <i>Nature Genetics</i> , 2017, 49, 1231-1238.	21.4	216
22	High-throughput discovery of novel developmental phenotypes. <i>Nature</i> , 2016, 537, 508-514.	27.8	1,001
23	Human Fetal Testicular Tissue Xenotransplantation: A Platform to Study the Effect of Gonadotropins on Human Germ Cell Development In Utero. <i>Journal of Urology</i> , 2015, 194, 585-591.	0.4	11
24	Analysis of mammalian gene function through broad-based phenotypic screens across a consortium of mouse clinics. <i>Nature Genetics</i> , 2015, 47, 969-978.	21.4	137
25	The Bulk of Autotaxin Activity Is Dispensable for Adult Mouse Life. <i>PLoS ONE</i> , 2015, 10, e0143083.	2.5	55
26	Histopathology reveals correlative and unique phenotypes in a high throughput mouse phenotyping screen. <i>DMM Disease Models and Mechanisms</i> , 2014, 7, 515-24.	2.4	44
27	Analysis of Phenotype. , 2014, , 431-487.		5
28	The mouse pathology ontology, MPATH; structure and applications. <i>Journal of Biomedical Semantics</i> , 2013, 4, 18.	1.6	32
29	Phenotype and genotype comparison of C57BL/6N substrains contributing to the International Mouse Phenotyping Consortium (IMPC). <i>FASEB Journal</i> , 2013, 27, 1b195.	0.5	0
30	The mammalian gene function resource: the international knockout mouse consortium. <i>Mammalian Genome</i> , 2012, 23, 580-586.	2.2	292
31	Pathology of the Laboratory Mouse. <i>Toxicologic Pathology</i> , 2011, 39, 559-562.	1.8	17
32	Two mouse mutations mapped to chromosome 11 with differing morphologies but similar progressive inflammatory alopecia. <i>Experimental Dermatology</i> , 2005, 14, 373-379.	2.9	8
33	Inactivation of <i>Fac</i> in mice produces inducible chromosomal instability and reduced fertility reminiscent of Fanconi anaemia. <i>Nature Genetics</i> , 1996, 12, 448-451.	21.4	241