

Fuyi Xu

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/5244774/publications.pdf>

Version: 2024-02-01

31
papers

237
citations

1039406

9
h-index

1125271

13
g-index

32
all docs

32
docs citations

32
times ranked

380
citing authors

#	ARTICLE	IF	CITATIONS
1	Systems genetics analysis defines importance of TMEM43<i>LUMA</i> for cardiac- and metabolic-related pathways. <i>Physiological Genomics</i> , 2022, 54, 22-35.	1.0	10
2	A systems genetics approach delineates the role of Bcl2 in leukemia pathogenesis. <i>Leukemia Research</i> , 2022, 114, 106804.	0.4	2
3	Analysis of cardiac function and morphology in genetic reference population of BXD strains reveals associated eQTLs and candidate causal and modifier genes. <i>FASEB Journal</i> , 2022, 36, .	0.2	0
4	Characterizing modifier genes of cardiac fibrosis phenotype in hypertrophic cardiomyopathy. <i>International Journal of Cardiology</i> , 2021, 330, 135-141.	0.8	6
5	Deficiency in nebulin repeats of sarcomeric nebulin is detrimental for cardiomyocyte tolerance to exercise and biomechanical stress. <i>American Journal of Physiology - Heart and Circulatory Physiology</i> , 2021, 320, H2130-H2146.	1.5	3
6	Combining whole exome sequencing with in silico analysis and clinical data to identify candidate variants in pediatric left ventricular noncompaction. <i>International Journal of Cardiology</i> , 2021, 347, 29-37.	0.8	2
7	Ace2 and Tmprss2 Expressions Are Regulated by Dhx32 and Influence the Gastrointestinal Symptoms Caused by SARS-CoV-2. <i>Journal of Personalized Medicine</i> , 2021, 11, 1212.	1.1	5
8	Hippocampal Transcriptome-Wide Association Study Reveals Correlations Between Impaired Glutamatergic Synapse Pathway and Age-Related Hearing Loss in BXD-Recombinant Inbred Mice. <i>Frontiers in Neuroscience</i> , 2021, 15, 745668.	1.4	4
9	Humanized GPRC6AKGKY is a gain-of-function polymorphism in mice. <i>Scientific Reports</i> , 2020, 10, 11143.	1.6	11
10	Genomic Characteristics and Selection Signatures in Indigenous Chongming White Goat (Capra) Tj ETQq0 0 0 rgBT/Overlock 10 Tf 50 3	1.1	7
11	Exploring the Role of Chemokine Receptor 6 (Ccr6) in the BXD Mouse Model of Gulf War Illness. <i>Frontiers in Neuroscience</i> , 2020, 14, 818.	1.4	4
12	The Genetic Dissection of Ace2 Expression Variation in the Heart of Murine Genetic Reference Population. <i>Frontiers in Cardiovascular Medicine</i> , 2020, 7, 582949.	1.1	13
13	Role of GPRC6A in Regulating Hepatic Energy Metabolism in Mice. <i>Scientific Reports</i> , 2020, 10, 7216.	1.6	15
14	Identifying modifier genes for hypertrophic cardiomyopathy. <i>Journal of Molecular and Cellular Cardiology</i> , 2020, 144, 119-126.	0.9	12
15	Genome-wide transcriptome architecture in a mouse model of Gulf War Illness. <i>Brain, Behavior, and Immunity</i> , 2020, 89, 209-223.	2.0	13
16	Electrical Stimulation Induces Retinal Müller Cell Proliferation and Their Progenitor Cell Potential. <i>Cells</i> , 2020, 9, 781.	1.8	24
17	Modeling the Genetic Basis of Individual Differences in Susceptibility to Gulf War Illness. <i>Brain Sciences</i> , 2020, 10, 143.	1.1	11
18	An Age-Related Hearing Protection Locus on Chromosome 16 of BXD Strain Mice. <i>Neural Plasticity</i> , 2020, 2020, 1-11.	1.0	4

#	ARTICLE	IF	CITATIONS
19	Candidate Regulators of Dyslipidemia in Chromosome 1 Substitution Lines Using Liver Co-Expression Profiling Analysis. <i>Frontiers in Genetics</i> , 2020, 10, 1258.	1.1	1
20	Genetic Dissection of Hypertrophic Cardiomyopathy with Myocardial RNA-Seq. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3040.	1.8	26
21	Genome-Wide Analysis of MicroRNA-related Single Nucleotide Polymorphisms (SNPs) in Mouse Genome. <i>Scientific Reports</i> , 2020, 10, 5789.	1.6	11
22	Genetic Dissection of the Regulatory Mechanisms of Ace2 in the Infected Mouse Lung. <i>Frontiers in Immunology</i> , 2020, 11, 607314.	2.2	14
23	A systems genetics approach to revealing the molecular network of the retina. <i>Molecular Vision</i> , 2020, 26, 459-471.	1.1	0
24	Familial Left Ventricular Non-Compaction Is Associated With a Rare p.V407I Variant in Bone Morphogenetic Protein 10. <i>Circulation Journal</i> , 2019, 83, 1737-1746.	0.7	12
25	A multiplex SNP genotyping by allele-specific PCR based on stem-loop and universal fluorescent primers of Chr1 dixin mice. <i>Electrophoresis</i> , 2019, 40, 1600-1605.	1.3	2
26	Exploring the involvement of Tac2 in the mouse hippocampal stress response through gene networking. <i>Gene</i> , 2019, 696, 176-185.	1.0	4
27	Genetic Dissection of Femoral and Tibial Microarchitecture. <i>JBMR Plus</i> , 2019, 3, e10241.	1.3	6
28	Investigating a downstream gene of using the systems genetics method. <i>Molecular Vision</i> , 2019, 25, 222-236.	1.1	2
29	Sequence analysis of chromosome 1 revealed different selection patterns between Chinese wild mice and laboratory strains. <i>Molecular Genetics and Genomics</i> , 2017, 292, 1111-1121.	1.0	2
30	Chromosome 1 Sequence Analysis of C57BL/6J-Chr1KM Mouse Strain. <i>International Journal of Genomics</i> , 2017, 2017, 1-9.	0.8	3
31	Genome Sequencing of Chromosome 1 Substitution Lines Derived from Chinese Wild Mice Revealed a Unique Resource for Genetic Studies of Complex Traits. <i>G3: Genes, Genomes, Genetics</i> , 2016, 6, 3571-3580.	0.8	8