

Sibylle Sabrautzki

List of Publications by Year in descending order

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Version: 2024-02-01

16
papers

260
citations

1040056

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996975

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17
all docs

17
docs citations

17
times ranked

629
citing authors

#	ARTICLE	IF	CITATIONS
1	Welfare Assessment of Adult Laboratory Zebrafish: A Practical Guide. <i>Zebrafish</i> , 2021, 18, 282-292.	1.1	2
2	A point mutation in the <i>Pdia6</i> gene results in loss of pancreatic β -cell identity causing overt diabetes. <i>Molecular Metabolism</i> , 2021, 54, 101334.	6.5	3
3	Systemic Jak1 activation provokes hepatic inflammation and imbalanced FGF23 production and cleavage. <i>FASEB Journal</i> , 2021, 35, e21302.	0.5	13
4	Combining fish and environmental PCR for diagnostics of diseased laboratory zebrafish in recirculating systems. <i>PLoS ONE</i> , 2019, 14, e0222360.	2.5	16
5	Mutation in the mouse histone gene <i>Hist2h3c1</i> leads to degeneration of the lens vesicle and severe microphthalmia. <i>Experimental Eye Research</i> , 2019, 188, 107632.	2.6	4
6	The elevation of circulating fibroblast growth factor 23 without kidney disease does not increase cardiovascular disease risk. <i>Kidney International</i> , 2018, 94, 49-59.	5.2	62
7	Point mutation of <i>Ffar1</i> abrogates fatty acid-dependent insulin secretion, but protects against HFD-induced glucose intolerance. <i>Molecular Metabolism</i> , 2017, 6, 1304-1312.	6.5	19
8	Standardized, systemic phenotypic analysis reveals kidney dysfunction as main alteration of <i>Kctd1 l27N</i> mutant mice. <i>Journal of Biomedical Science</i> , 2017, 24, 57.	7.0	8
9	Viable <i>Ednra Y129F</i> mice feature human mandibulofacial dysostosis with alopecia (MFDA) syndrome due to the homologue mutation. <i>Mammalian Genome</i> , 2016, 27, 587-598.	2.2	5
10	Exome sequencing identifies a nonsense mutation in <i>Fam46a</i> associated with bone abnormalities in a new mouse model for skeletal dysplasia. <i>Mammalian Genome</i> , 2016, 27, 111-121.	2.2	27
11	Generation and Standardized, Systemic Phenotypic Analysis of <i>Pou3f3L423P</i> Mutant Mice. <i>PLoS ONE</i> , 2016, 11, e0150472.	2.5	14
12	The First <i>Scube3</i> Mutant Mouse Line with Pleiotropic Phenotypic Alterations. <i>G3: Genes, Genomes, Genetics</i> , 2016, 6, 4035-4046.	1.8	9
13	New Mutation in the Mouse <i>Xpd/Ercc2</i> Gene Leads to Recessive Cataracts. <i>PLoS ONE</i> , 2015, 10, e0125304.	2.5	24
14	Genomic characterization of mutant laboratory mouse strains by exome sequencing and annotation lift-over. <i>BMC Genomics</i> , 2015, 16, 351.	2.8	0
15	An ENU Mutagenesis-Derived Mouse Model with a Dominant <i>Jak1</i> Mutation Resembling Phenotypes of Systemic Autoimmune Disease. <i>American Journal of Pathology</i> , 2013, 183, 352-368.	3.8	24
16	New mouse models for metabolic bone diseases generated by genome-wide ENU mutagenesis. <i>Mammalian Genome</i> , 2012, 23, 416-430.	2.2	30