

M Elizabeth Ross

List of Publications by Year in descending order

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Version: 2024-02-01

21
papers

872
citations

687220

13
h-index

677027

22
g-index

23
all docs

23
docs citations

23
times ranked

1581
citing authors

#	ARTICLE	IF	CITATIONS
1	Single-nuclei isoform RNA sequencing unlocks barcoded exon connectivity in frozen brain tissue. <i>Nature Biotechnology</i> , 2022, 40, 1082-1092.	9.4	52
2	Astrocytes derived from ASD individuals alter behavior and destabilize neuronal activity through aberrant Ca ²⁺ signaling. <i>Molecular Psychiatry</i> , 2022, 27, 2470-2484.	4.1	26
3	A spatially resolved brain region- and cell type-specific isoform atlas of the postnatal mouse brain. <i>Nature Communications</i> , 2021, 12, 463.	5.8	109
4	Genome-wide investigation identifies a rare copy-number variant burden associated with human spina bifida. <i>Genetics in Medicine</i> , 2021, 23, 1211-1218.	1.1	10
5	Unraveling the complex genetics of neural tube defects: From biological models to human genomics and back. <i>Genesis</i> , 2021, 59, e23459.	0.8	19
6	Systems biology analysis of human genomes points to key pathways conferring spina bifida risk. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2021, 118, .	3.3	11
7	Loss of <i>RAD9B</i> impairs early neural development and contributes to the risk for human spina bifida. <i>Human Mutation</i> , 2020, 41, 786-799.	1.1	14
8	Unlocking the genetic complexity of congenital hydrocephalus. <i>Nature Medicine</i> , 2020, 26, 1682-1683.	15.2	5
9	Postzygotic inactivating mutations of RHOA cause a mosaic neuroectodermal syndrome. <i>Nature Genetics</i> , 2019, 51, 1438-1441.	9.4	25
10	SLC35A2 ^Δ CDG: Functional characterization, expanded molecular, clinical, and biochemical phenotypes of 30 unreported Individuals. <i>Human Mutation</i> , 2019, 40, 908-925.	1.1	39
11	Tight junction protein occludin regulates progenitor Self-Renewal and survival in developing cortex. <i>ELife</i> , 2019, 8, .	2.8	27
12	Mature Hippocampal Neurons Require LIS1 for Synaptic Integrity: Implications for Cognition. <i>Biological Psychiatry</i> , 2018, 83, 518-529.	0.7	11
13	Single-cell isoform RNA sequencing characterizes isoforms in thousands of cerebellar cells. <i>Nature Biotechnology</i> , 2018, 36, 1197-1202.	9.4	253
14	Wiring the Human Brain: A User's Handbook. <i>Neuron</i> , 2017, 95, 482-485.	3.8	11
15	Mutation in noncoding <i>RNA RNU12</i> causes early onset cerebellar ataxia. <i>Annals of Neurology</i> , 2017, 81, 68-78.	2.8	68
16	Interneuron precursor transplants in adult hippocampus reverse psychosis-relevant features in a mouse model of hippocampal disinhibition. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 7450-7455.	3.3	81
17	Cell cycle regulation and interneuron production. <i>Developmental Neurobiology</i> , 2011, 71, 2-9.	1.5	13
18	Genes and the long and winding road to cortical construction and cognition. <i>Neurobiology of Disease</i> , 2010, 38, 145-147.	2.1	1

#	ARTICLE	IF	CITATIONS
19	Gene-environment interactions, folate metabolism and the embryonic nervous system. Wiley Interdisciplinary Reviews: Systems Biology and Medicine, 2010, 2, 471-480.	6.6	28
20	Functional interactions between the LRP6 WNT co-receptor and folate supplementation. Human Molecular Genetics, 2010, 19, 4560-4572.	1.4	41
21	Mechanistic insights into folate supplementation from Crooked tail and other NTD-prone mutant mice. Birth Defects Research Part A: Clinical and Molecular Teratology, 2009, 85, 314-321.	1.6	26