Ming Zhang

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/11784251/publications.pdf Version: 2024-02-01



Μίνο Ζηλνο

#	Article	IF	CITATIONS
1	Isoformâ€specific antibodies reveal distinct subcellular localizations of <scp>C</scp> 9orf72 in amyotrophic lateral sclerosis. Annals of Neurology, 2015, 78, 568-583.	5.3	123
2	The C9orf72 repeat expansion itself is methylated in ALS and FTLD patients. Acta Neuropathologica, 2015, 129, 715-727.	7.7	114
3	A Predictive Metabolic Signature for the Transition From Gestational Diabetes Mellitus to Type 2 Diabetes. Diabetes, 2016, 65, 2529-2539.	0.6	113
4	Drug Repositioning for Alzheimer's Disease Based on Systematic â€~omics' Data Mining. PLoS ONE, 2016, e0168812.	, 11. 2.5	95
5	Mutation analysis of <i>CHCHD10</i> in different neurodegenerative diseases. Brain, 2015, 138, e380-e380.	7.6	86
6	Drug Repositioning for Diabetes Based on 'Omics' Data Mining. PLoS ONE, 2015, 10, e0126082.	2.5	74
7	Characterization of Zinc Influx Transporters (ZIPs) in Pancreatic β Cells. Journal of Biological Chemistry, 2015, 290, 18757-18769.	3.4	58
8	Jump from Pre-mutation to Pathologic Expansion in C9orf72. American Journal of Human Genetics, 2015, 96, 962-970.	6.2	50
9	Progesterone Receptor Membrane Component 1 Is a Functional Part of the Glucagon-like Peptide-1 (GLP-1) Receptor Complex in Pancreatic β Cells. Molecular and Cellular Proteomics, 2014, 13, 3049-3062.	3.8	48
10	DNA methylation age-acceleration is associated with disease duration and age at onset in C9orf72 patients. Acta Neuropathologica, 2017, 134, 271-279.	7.7	46
11	A C6orf10/LOC101929163 locus is associated with age of onset in C9orf72 carriers. Brain, 2018, 141, 2895-2907.	7.6	39
12	Genetic and epigenetic study of ALS-discordant identical twins with double mutations in <i>SOD1</i> and <i>ARHGEF28</i> . Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 1268-1270.	1.9	35
13	Unaffected mosaic <i>C9orf72</i> case. Neurology, 2018, 90, e323-e331.	1.1	33
14	DNA methylation age acceleration is associated with ALS age of onset and survival. Acta Neuropathologica, 2020, 139, 943-946.	7.7	30
15	NMDA Receptor Hypofunction Induces Dysfunctions of Energy Metabolism And Semaphorin Signaling in Rats: A Synaptic Proteome Study. Schizophrenia Bulletin, 2012, 38, 579-591.	4.3	26
16	The ONDRISeq panel: custom-designed next-generation sequencing of genes related to neurodegeneration. Npj Genomic Medicine, 2016, 1, 16032.	3.8	26
17	Positive association between ALDH1A2 and schizophrenia in the Chinese population. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2009, 33, 1491-1495.	4.8	25
18	A Novel GLP1 Receptor Interacting Protein ATP6ap2 Regulates Insulin Secretion in Pancreatic Beta Cells. Journal of Biological Chemistry, 2015, 290, 25045-25061.	3.4	25

Ming Zhang

#	Article	IF	CITATIONS
19	Parkinsonism due to A53E αâ€synuclein gene mutation: Clinical, genetic, epigenetic, and biochemical features. Movement Disorders, 2018, 33, 1950-1955.	3.9	25
20	Metabolomic Analysis Reveals Metabolic Disturbance in the Cortex and Hippocampus of Subchronic MK-801 Treated Rats. PLoS ONE, 2013, 8, e60598.	2.5	24
21	The Identification of Novel Protein-Protein Interactions in Liver that Affect Glucagon Receptor Activity. PLoS ONE, 2015, 10, e0129226.	2.5	19
22	Marked Differences in C9orf72 Methylation Status and Isoform Expression between C9/ALS Human Embryonic and Induced Pluripotent Stem Cells. Stem Cell Reports, 2016, 7, 927-940.	4.8	19
23	Targeted Next-generation Sequencing and Bioinformatics Pipeline to Evaluate Genetic Determinants of Constitutional Disease. Journal of Visualized Experiments, 2018, , .	0.3	17
24	Mutation analysis of CHCHD2 in Canadian patients with familial Parkinson's disease. Neurobiology of Aging, 2016, 38, 217.e7-217.e8.	3.1	16
25	<i>C9orf72</i> and <i>ATXN2</i> repeat expansions coexist in a family with ataxia, dementia, and parkinsonism. Movement Disorders, 2017, 32, 158-162.	3.9	15
26	Mutation analysis of C9orf72 in patients with corticobasal syndrome. Neurobiology of Aging, 2015, 36, 2905.e1-2905.e5.	3.1	13
27	Neuropathologic description of <i>CHCHD10</i> mutated amyotrophic lateral sclerosis. Neurology: Genetics, 2020, 6, e394.	1.9	13
28	Proteome alterations of cortex and hippocampus tissues in mice subjected to vitamin A depletion. Journal of Nutritional Biochemistry, 2011, 22, 1003-1008.	4.2	12
29	Genetic and epigenetic study of an Alzheimer's disease family with monozygotic triplets. Brain, 2019, 142, 3375-3381.	7.6	11
30	Genetic analysis of CHCHD2 and CHCHD10 in Italian patients with Parkinson's disease. Neurobiology of Aging, 2017, 53, 193.e7-193.e8.	3.1	8
31	Mutation analysis of CHCHD2 and CHCHD10 in Italian patients with mitochondrial myopathy. Neurobiology of Aging, 2018, 66, 181.e1-181.e2.	3.1	8
32	Combined epigenetic/genetic study identified an ALS age of onset modifier. Acta Neuropathologica Communications, 2021, 9, 75.	5.2	7
33	Proteome alteration of U251 human astrocytoma cell after inhibiting retinoic acid synthesis. Molecular and Cellular Biochemistry, 2009, 323, 185-193.	3.1	5
34	Vitamin A depletion alters sensitivity of motor behavior to MK-801 in C57BL/6J mice. Behavioral and Brain Functions, 2010, 6, 7.	3.3	5
35	DNA methylation age acceleration is associated with age of onset in Chinese spinocerebellar ataxia type 3 patients. Neurobiology of Aging, 2022, 113, 1-6.	3.1	3
36	Response to a letter to the editor. Neurobiology of Aging, 2019, 78, 195-196.	3.1	0

#	Article	IF	CITATIONS
37	A study of the mechanism of yinzhihuang"Equation missing" No EquationSource Format="TEX", only image injection in the treatment of infantile hepatitis syndromeinjection in the treatment of infantile hepatitis syndromeinjection in the treatment of infantile hepatitis syndromeinjection in the treatment of infantile hepatitis syndrome.		0