Ming Zhang

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/11784251/publications.pdf

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37 papers	1,271 citations	361413 20 h-index	377865 34 g-index
40 all docs	40 docs citations	40 times ranked	2687 citing authors

#	Article	IF	Citations
1	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
2	Combined epigenetic/genetic study identified an ALS age of onset modifier. Acta Neuropathologica Communications, 2021, 9, 75.	5.2	7
3	DNA methylation age acceleration is associated with ALS age of onset and survival. Acta Neuropathologica, 2020, 139, 943-946.	7.7	30
4	Neuropathologic description of CHCHD10 mutated amyotrophic lateral sclerosis. Neurology: Genetics, 2020, 6, e394.	1.9	13
5	Genetic and epigenetic study of an Alzheimerâ€™s disease family with monozygotic triplets. Brain, 2019, 142, 3375-3381.	7.6	11
6	Response to a letter to the editor. Neurobiology of Aging, 2019, 78, 195-196.	3.1	0
7	Targeted Next-generation Sequencing and Bioinformatics Pipeline to Evaluate Genetic Determinants of Constitutional Disease. Journal of Visualized Experiments, 2018, , .	0.3	17
8	Unaffected mosaic C9orf72 case. Neurology, 2018, 90, e323-e331.	1.1	33
9	Mutation analysis of CHCHD2 and CHCHD10 in Italian patients with mitochondrial myopathy. Neurobiology of Aging, 2018, 66, 181.e1-181.e2.	3.1	8
10	Parkinsonism due to A53E Î±â€synuclein gene mutation: Clinical, genetic, epigenetic, and biochemical features. Movement Disorders, 2018, 33, 1950-1955.	3.9	25
11	A C6orf10/LOC101929163 locus is associated with age of onset in C9orf72 carriers. Brain, 2018, 141, 2895-2907.	7.6	39
12	C9orf72 and ATXN2 repeat expansions coexist in a family with ataxia, dementia, and parkinsonism. Movement Disorders, 2017, 32, 158-162.	3.9	15
13	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
14	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
15	Genetic and epigenetic study of ALS-discordant identical twins with double mutations inSOD1andARHGEF28. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 1268-1270.	1.9	35
16	The ONDRISeq panel: custom-designed next-generation sequencing of genes related to neurodegeneration. Npj Genomic Medicine, 2016, 1, 16032.	3.8	26
17	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
18	A Predictive Metabolic Signature for the Transition From Gestational Diabetes Mellitus to Type 2 Diabetes. Diabetes, 2016, 65, 2529-2539.	0.6	113

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19	Mutation analysis of CHCHD2 in Canadian patients with familial Parkinson's disease. Neurobiology of Aging, 2016, 38, 217.e7-217.e8.	3.1	16
20	Drug Repositioning for Alzheimerâ€™s Disease Based on Systematic â€˜omicsâ€™ Data Mining. PLoS ONE, 2016, 11, e0168812.	2.3	95
21	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
22	The Identification of Novel Protein-Protein Interactions in Liver that Affect Glucagon Receptor Activity. PLoS ONE, 2015, 10, e0129226.	2.5	19
23	Jump from Pre-mutation to Pathologic Expansion in C9orf72. American Journal of Human Genetics, 2015, 96, 962-970.	6.2	50
24	Characterization of Zinc Influx Transporters (ZIPs) in Pancreatic Î² Cells. Journal of Biological Chemistry, 2015, 290, 18757-18769.	3.4	58
25	The C9orf72 repeat expansion itself is methylated in ALS and FTLD patients. Acta Neuropathologica, 2015, 129, 715-727.	7.7	114
26	Drug Repositioning for Diabetes Based on 'Omics' Data Mining. PLoS ONE, 2015, 10, e0126082.	2.5	74
27	Mutation analysis of C9orf72 in patients with corticobasal syndrome. Neurobiology of Aging, 2015, 36, 2905.e1-2905.e5.	3.1	13
28	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
29	Mutation analysis ofCHCHD10in different neurodegenerative diseases. Brain, 2015, 138, e380-e380.	7.6	86
30	Progesterone Receptor Membrane Component 1 Is a Functional Part of the Glucagon-like Peptide-1 (GLP-1) Receptor Complex in Pancreatic Î² Cells. Molecular and Cellular Proteomics, 2014, 13, 3049-3062.	3.8	48
31	Metabolomic Analysis Reveals Metabolic Disturbance in the Cortex and Hippocampus of Subchronic MK-801 Treated Rats. PLoS ONE, 2013, 8, e60598.	2.5	24
32	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
33	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
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	Proteome alteration of U251 human astrocytoma cell after inhibiting retinoic acid synthesis. Molecular and Cellular Biochemistry, 2009, 323, 185-193.	3.1	5
36	Positive association between ALDH1A2 and schizophrenia in the Chinese population. Progress in Neuro-Psychopharmacology and Biological Psychiatry, 2009, 33, 1491-1495.	4.8	25

#	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 syndrome. , 1995, 1, 122-124.		0