

Adam Keith Walker

List of Publications by Year in descending order

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Version: 2024-02-01

51
papers

2,305
citations

279778

23
h-index

223791

46
g-index

58
all docs

58
docs citations

58
times ranked

3149
citing authors

#	ARTICLE	IF	CITATIONS
1	Functional recovery in new mouse models of ALS/FTLD after clearance of pathological cytoplasmic TDP-43. <i>Acta Neuropathologica</i> , 2015, 130, 643-660.	7.7	215
2	Protein disulphide isomerase protects against protein aggregation and is S-nitrosylated in amyotrophic lateral sclerosis. <i>Brain</i> , 2010, 133, 105-116.	7.6	156
3	ALS-Associated TDP-43 Induces Endoplasmic Reticulum Stress, Which Drives Cytoplasmic TDP-43 Accumulation and Stress Granule Formation. <i>PLoS ONE</i> , 2013, 8, e81170.	2.5	141
4	The Pathobiology of TDP-43 C-Terminal Fragments in ALS and FTLD. <i>Frontiers in Neuroscience</i> , 2019, 13, 335.	2.8	135
5	Redefining the Role of Metallothionein within the Injured Brain. <i>Journal of Biological Chemistry</i> , 2008, 283, 15349-15358.	3.4	130
6	Mutant FUS induces endoplasmic reticulum stress in amyotrophic lateral sclerosis and interacts with protein disulfide-isomerase. <i>Neurobiology of Aging</i> , 2012, 33, 2855-2868.	3.1	88
7	Defects in optineurin- and myosin VI-mediated cellular trafficking in amyotrophic lateral sclerosis. <i>Human Molecular Genetics</i> , 2015, 24, 3830-3846.	2.9	71
8	Astrocytic TDP-43 Pathology in Alexander Disease. <i>Journal of Neuroscience</i> , 2014, 34, 6448-6458.	3.6	64
9	Mutant SOD1 inhibits ER-Golgi transport in amyotrophic lateral sclerosis. <i>Journal of Neurochemistry</i> , 2014, 129, 190-204.	3.9	61
10	Extracellular wildtype and mutant SOD1 induces ER-Golgi pathology characteristic of amyotrophic lateral sclerosis in neuronal cells. <i>Cellular and Molecular Life Sciences</i> , 2013, 70, 4181-4195.	5.4	59
11	Stress signaling from the endoplasmic reticulum: A central player in the pathogenesis of amyotrophic lateral sclerosis. <i>IUBMB Life</i> , 2011, 63, n/a-n/a.	3.4	58
12	Protein Quality Control and the Amyotrophic Lateral Sclerosis/Frontotemporal Dementia Continuum. <i>Frontiers in Molecular Neuroscience</i> , 2017, 10, 119.	2.9	58
13	Proteomics Approaches for Biomarker and Drug Target Discovery in ALS and FTD. <i>Frontiers in Neuroscience</i> , 2019, 13, 548.	2.8	57
14	Impaired NHEJ repair in amyotrophic lateral sclerosis is associated with TDP-43 mutations. <i>Molecular Neurodegeneration</i> , 2020, 15, 51.	10.8	54
15	Neuroinflammation in schizophrenia: the role of nuclear factor kappa B. <i>Translational Psychiatry</i> , 2021, 11, 528.	4.8	54
16	Pathogenic mutation in the ALS/FTD gene, CCNF, causes elevated Lys48-linked ubiquitylation and defective autophagy. <i>Cellular and Molecular Life Sciences</i> , 2018, 75, 335-354.	5.4	44
17	An insoluble frontotemporal lobar degeneration-associated TDP-43 C-terminal fragment causes neurodegeneration and hippocampus pathology in transgenic mice. <i>Human Molecular Genetics</i> , 2015, 24, 7241-7254.	2.9	39
18	Mislocalisation of TDP-43 to the cytoplasm causes cortical hyperexcitability and reduced excitatory neurotransmission in the motor cortex. <i>Journal of Neurochemistry</i> , 2021, 157, 1300-1315.	3.9	36

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19	Nuclear factor kappa B activation appears weaker in schizophrenia patients with high brain cytokines than in non-schizophrenic controls with high brain cytokines. <i>Journal of Neuroinflammation</i> , 2020, 17, 215.	7.2	33
20	Bim Links ER Stress and Apoptosis in Cells Expressing Mutant SOD1 Associated with Amyotrophic Lateral Sclerosis. <i>PLoS ONE</i> , 2012, 7, e35413.	2.5	31
21	TDP-43 pathology: From noxious assembly to therapeutic removal. <i>Progress in Neurobiology</i> , 2022, 211, 102229.	5.7	30
22	Casein kinase II phosphorylation of cyclin F at serine 621 regulates the Lys48-ubiquitylation E3 ligase activity of the SCF (cyclin F) complex. <i>Open Biology</i> , 2017, 7, 170058.	3.6	29
23	Circulating epinephrine is not required for chronic stress to enhance metastasis. <i>Psychoneuroendocrinology</i> , 2019, 99, 191-195.	2.7	26
24	Impaired glymphatic function in the early stages of disease in a TDP-43 mouse model of amyotrophic lateral sclerosis. <i>Translational Neurodegeneration</i> , 2022, 11, 17.	8.0	26
25	Novel monoclonal antibodies to normal and pathologically altered human TDP-43 proteins. <i>Acta Neuropathologica Communications</i> , 2014, 2, 33.	5.2	25
26	ERp57 is protective against mutant SOD1-induced cellular pathology in amyotrophic lateral sclerosis. <i>Human Molecular Genetics</i> , 2018, 27, 1311-1331.	2.9	24
27	Regional, cellular and species difference of two key neuroinflammatory genes implicated in schizophrenia. <i>Brain, Behavior, and Immunity</i> , 2020, 88, 826-839.	4.1	23
28	Label-Free Fluorescent Poly(amidoamine) Dendrimer for Traceable and Controlled Drug Delivery. <i>Biomacromolecules</i> , 2019, 20, 2148-2158.	5.4	19
29	<sc>Nâ€“</sc>linked glycosylation modulates dimerization of protein disulfide isomerase familyA<sc>A</sc> memberA2 (<sc>PDIA</sc>2). <i>FEBS Journal</i> , 2013, 280, 233-243.	4.7	18
30	Protein Disulfide Isomerase and the Endoplasmic Reticulum in Amyotrophic Lateral Sclerosis. <i>Journal of Neuroscience</i> , 2010, 30, 3865-3867.	3.6	15
31	Mechanisms of Neuroprotection by Protein Disulphide Isomerase in Amyotrophic Lateral Sclerosis. <i>Neurology Research International</i> , 2011, 2011, 1-7.	1.3	15
32	Riluzole does not ameliorate disease caused by cytoplasmic TDPâ€“43 in a mouse model of amyotrophic lateral sclerosis. <i>European Journal of Neuroscience</i> , 2021, 54, 6237-6255.	2.6	15
33	Stilbenes from <i>Veratrum maackii</i> Regel Protect against Ethanol-Induced DNA Damage in Mouse Cerebellum and Cerebral Cortex. <i>ACS Chemical Neuroscience</i> , 2018, 9, 1616-1624.	3.5	14
34	Workflow for Rapidly Extracting Biological Insights from Complex, Multicondition Proteomics Experiments with WGCNA and PloGO2. <i>Journal of Proteome Research</i> , 2020, 19, 2898-2906.	3.7	13
35	Trajectory of change in brain complement factors from neonatal to young adult humans. <i>Journal of Neurochemistry</i> , 2021, 157, 479-493.	3.9	12
36	Unbiased Label-Free Quantitative Proteomics of Cells Expressing Amyotrophic Lateral Sclerosis (ALS) Mutations in CENF Reveals Activation of the Apoptosis Pathway: A Workflow to Screen Pathogenic Gene Mutations. <i>Frontiers in Molecular Neuroscience</i> , 2021, 14, 627740.	2.9	12

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37	Peripheral NF- κ B dysregulation in people with schizophrenia drives inflammation: putative anti-inflammatory functions of NF- κ B kinases. <i>Translational Psychiatry</i> , 2022, 12, 21.	4.8	12
38	Metallothionein expression by NG2 glial cells following CNS injury. <i>Cellular and Molecular Life Sciences</i> , 2007, 64, 2716-2722.	5.4	11
39	Disrupting circadian rhythms promotes cancer-induced inflammation in mice. <i>Brain, Behavior, & Immunity - Health</i> , 2022, 21, 100428.	2.5	9
40	Genetic and immunopathological analysis of CHCHD10 in Australian amyotrophic lateral sclerosis and frontotemporal dementia and transgenic TDP-43 mice. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020, 91, 162-171.	1.9	8
41	The Cysteine (Cys) Residues Cys-6 and Cys-111 in Mutant Superoxide Dismutase 1 (SOD1) A4V Are Required for Induction of Endoplasmic Reticulum Stress in Amyotrophic Lateral Sclerosis. <i>Journal of Molecular Neuroscience</i> , 2020, 70, 1357-1368.	2.3	8
42	Longitudinal exploration of cancer-related cognitive impairment in patients with newly diagnosed aggressive lymphoma: protocol for a feasibility study. <i>BMJ Open</i> , 2020, 10, e038312.	1.9	6
43	Neurodegenerative disease-associated protein aggregates are poor inducers of the heat shock response in neuronal cells. <i>Journal of Cell Science</i> , 2020, 133, .	2.0	6
44	Cancer-related cognitive impairment in patients with newly diagnosed aggressive lymphoma undergoing standard chemotherapy: a longitudinal feasibility study. <i>Supportive Care in Cancer</i> , 0, , .	2.2	6
45	Cryptic inclusions UNCover losses driving neurodegeneration. <i>Trends in Genetics</i> , 2022, 38, 889-891.	6.7	4
46	Where There's Smoke, There's Fire" But Who Is Lighting the Match? Bolstering Transcriptional Evidence for the Role of Nuclear Factor- κ B in Neuroimmune Activation in Schizophrenia. <i>Biological Psychiatry</i> , 2019, 85, 5-7.	1.3	2
47	miR-23a suppression accelerates functional decline in the rNLS8 mouse model of TDP-43 proteinopathy. <i>Neurobiology of Disease</i> , 2022, 162, 105559.	4.4	2
48	Clocking onto chemotherapy to enhance cancer treatment. <i>Brain, Behavior, and Immunity</i> , 2022, 100, 172-173.	4.1	1
49	Early and progressive dysfunction revealed by in vivo neurite imaging in the rNLS8 TDP-43 mouse model of ALS. <i>NeuroImage: Clinical</i> , 2022, 34, 103016.	2.7	1
50	Note in reference to "Mutant FUS induces endoplasmic reticulum stress in amyotrophic lateral sclerosis and interacts with protein disulfide-isomerase" [Neurobiol. Aging 33(12) (2012) 2855-2868]. <i>Neurobiology of Aging</i> , 2017, 60, 205.	3.1	0
51	Involvement of endoplasmic reticulum stress in TDP-43-linked neurodegenerative disease. <i>Postdoc Journal</i> , 2014, 2, .	0.4	0