

Mel B Feany

List of Publications by Year in descending order

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77
papers

13,671
citations

50276
46
h-index

74163
75
g-index

84
all docs

84
docs citations

84
times ranked

13946
citing authors

#	ARTICLE	IF	CITATIONS
1	Case Study 1: A 55-Year-Old Woman With Progressive Cognitive, Perceptual, and Motor Impairments. <i>Journal of Neuropsychiatry and Clinical Neurosciences</i> , 2022, 34, 8-15.	1.8	2
2	Anastasis Drives Senescence and Non-Cell Autonomous Neurodegeneration in the Astroglial Pathology of Alexander Disease. <i>Journal of Neuroscience</i> , 2022, 42, 2584-2597.	3.6	2
3	Î±-synuclein impairs autophagosome maturation through abnormal actin stabilization. <i>PLoS Genetics</i> , 2021, 17, e1009359.	3.5	49
4	Oligomerization of Lrrk controls actin severing and Î±-synuclein neurotoxicity in vivo. <i>Molecular Neurodegeneration</i> , 2021, 16, 33.	10.8	6
5	Precision Medicine on the Fly: Using <i>Drosophila</i> to Decipher Gene-Environment Interactions in Parkinson's Disease. <i>Toxicological Sciences</i> , 2021, 182, 159-167.	3.1	8
6	Elevated Oxidative Stress and DNA Damage in Cortical Neurons of Chemotherapy Patients. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 705-712.	1.7	9
7	Parkinson's disease risk genes act in glia to control neuronal Î±-synuclein toxicity. <i>Neurobiology of Disease</i> , 2021, 159, 105482.	4.4	19
8	Antisense therapy in a rat model of Alexander disease reverses GFAP pathology, white matter deficits, and motor impairment. <i>Science Translational Medicine</i> , 2021, 13, eabg4711.	12.4	21
9	Iatrogenic Neuropathology of Systemic Therapies. <i>Surgical Pathology Clinics</i> , 2020, 13, 331-342.	1.7	4
10	Comparative proteomic analysis highlights metabolic dysfunction in Î±-synucleinopathy. <i>Npj Parkinson's Disease</i> , 2020, 6, 40.	5.3	16
11	Biotin rescues mitochondrial dysfunction and neurotoxicity in a tauopathy model. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 33608-33618.	7.1	20
12	New-Onset Delusions Herald an Underlying Neurodegenerative Condition. <i>Journal of Clinical Psychiatry</i> , 2020, 81, .	2.2	1
13	Glial Î±-synuclein promotes neurodegeneration characterized by a distinct transcriptional program in vivo. <i>Glia</i> , 2019, 67, 1933-1957.	4.9	27
14	PARP Inhibitors and Parkinson's Disease. <i>New England Journal of Medicine</i> , 2019, 380, 492-494.	27.0	31
15	Development of gene-environment interaction model in <i>Drosophila</i> for neurodegenerative disease: A step towards personalized medicine. <i>FASEB Journal</i> , 2019, 33, 813.14.	0.5	0
16	Î±-synuclein Induces Mitochondrial Dysfunction through Spectrin and the Actin Cytoskeleton. <i>Neuron</i> , 2018, 97, 108-124.e6.	8.1	181
17	A Conserved Cytoskeletal Signaling Cascade Mediates Neurotoxicity of FTDP-17 Tau Mutations <i>In Vivo</i> . <i>Journal of Neuroscience</i> , 2018, 38, 108-119.	3.6	35
18	Lrrk promotes tau neurotoxicity through dysregulation of actin and mitochondrial dynamics. <i>PLoS Biology</i> , 2018, 16, e2006265.	5.6	44

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19	Tissue and cellular rigidity and mechanosensitive signaling activation in Alexander disease. <i>Nature Communications</i> , 2018, 9, 1899.	12.8	43
20	Nortriptyline inhibits aggregation and neurotoxicity of alpha-synuclein by enhancing reconfiguration of the monomeric form. <i>Neurobiology of Disease</i> , 2017, 106, 191-204.	4.4	28
21	Aging-related tau astrogliopathy (ARTAG): harmonized evaluation strategy. <i>Acta Neuropathologica</i> , 2016, 131, 87-102.	7.7	380
22	Defective Phagocytic Corpse Processing Results in Neurodegeneration and Can Be Rescued by TORC1 Activation. <i>Journal of Neuroscience</i> , 2016, 36, 3170-3183.	3.6	50
23	An <i>In Vivo</i> Pharmacological Screen Identifies Cholinergic Signaling as a Therapeutic Target in Glial-Based Nervous System Disease. <i>Journal of Neuroscience</i> , 2016, 36, 1445-1455.	3.6	34
24	Lamin Dysfunction Mediates Neurodegeneration in Tauopathies. <i>Current Biology</i> , 2016, 26, 129-136.	3.9	184
25	Nitric oxide mediates glial-induced neurodegeneration in Alexander disease. <i>Nature Communications</i> , 2015, 6, 8966.	12.8	44
26	Connecting the dots between tau dysfunction and neurodegeneration. <i>Trends in Cell Biology</i> , 2015, 25, 46-53.	7.9	108
27	p53 prevents neurodegeneration by regulating synaptic genes. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 18055-18060.	7.1	65
28	Tau promotes neurodegeneration through global chromatin relaxation. <i>Nature Neuroscience</i> , 2014, 17, 357-366.	14.8	370
29	Glial cells are critical for the neuropathology of complex I deficiency in <i>Drosophila</i> . <i>Human Molecular Genetics</i> , 2014, 23, 4686-4692.	2.9	34
30	Functional screening in <i>Drosophila</i> identifies Alzheimer's disease susceptibility genes and implicates Tau-mediated mechanisms. <i>Human Molecular Genetics</i> , 2014, 23, 870-877.	2.9	147
31	Why size matters – balancing mitochondrial dynamics in Alzheimer's disease. <i>Trends in Neurosciences</i> , 2013, 36, 325-335.	8.6	150
32	Alexander Disease. <i>Journal of Neuroscience</i> , 2012, 32, 5017-5023.	3.6	210
33	A neuroprotective role for the DNA damage checkpoint in tauopathy. <i>Aging Cell</i> , 2012, 11, 360-362.	6.7	47
34	Tau Promotes Neurodegeneration via DRP1 Mislocalization <i>In Vivo</i> . <i>Neuron</i> , 2012, 75, 618-632.	8.1	331
35	Parkinson's Disease: Genetics and Pathogenesis. <i>Annual Review of Pathology: Mechanisms of Disease</i> , 2011, 6, 193-222.	22.4	654
36	Functional Screening of Alzheimer Pathology Genome-wide Association Signals in <i>Drosophila</i> . <i>American Journal of Human Genetics</i> , 2011, 88, 232-238.	6.2	81

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37	Protein Misfolding and Oxidative Stress Promote Glial-Mediated Neurodegeneration in an Alexander Disease Model. <i>Journal of Neuroscience</i> , 2011, 31, 2868-2877.	3.6	67
38	Glial Fibrillary Tangles and JAK/STAT-Mediated Glial and Neuronal Cell Death in a <i>Drosophila</i> Model of Glial Tauopathy. <i>Journal of Neuroscience</i> , 2010, 30, 16102-16113.	3.6	64
39	Lysosomal Dysfunction Promotes Cleavage and Neurotoxicity of Tau In Vivo. <i>PLoS Genetics</i> , 2010, 6, e1001026.	3.5	132
40	Parkinson's disease: Insights from non-traditional model organisms. <i>Progress in Neurobiology</i> , 2010, 92, 558-571.	5.7	60
41	New Approaches to the Pathology and Genetics of Neurodegeneration. <i>American Journal of Pathology</i> , 2010, 176, 2058-2066.	3.8	15
42	The Unfolded Protein Response Protects from Tau Neurotoxicity In Vivo. <i>PLoS ONE</i> , 2010, 5, e13084.	2.5	80
43	Tyrosine and serine phosphorylation of τ -synuclein have opposing effects on neurotoxicity and soluble oligomer formation. <i>Journal of Clinical Investigation</i> , 2009, 119, 3257-65.	8.2	158
44	Inactivation of <i>Drosophila</i> Huntingtin affects long-term adult functioning and the pathogenesis of a Huntington's disease model. <i>DMM Disease Models and Mechanisms</i> , 2009, 2, 247-266.	2.4	80
45	Cathepsin D expression level affects alpha-synuclein processing, aggregation, and toxicity in vivo. <i>Molecular Brain</i> , 2009, 2, 5.	2.6	232
46	τ -Synuclein S129 Phosphorylation Mutants Do Not Alter Nigrostriatal Toxicity in a Rat Model of Parkinson Disease. <i>Journal of Neuropathology and Experimental Neurology</i> , 2009, 68, 515-524.	1.7	111
47	Tau Phosphorylation Sites Work in Concert to Promote Neurotoxicity In Vivo. <i>Molecular Biology of the Cell</i> , 2007, 18, 5060-5068.	2.1	178
48	Connecting cell-cycle activation to neurodegeneration in <i>Drosophila</i> . <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2007, 1772, 446-456.	3.8	24
49	Calpain-Cleavage of τ -Synuclein. <i>American Journal of Pathology</i> , 2007, 170, 1725-1738.	3.8	213
50	Sirtuin 2 Inhibitors Rescue τ -Synuclein-Mediated Toxicity in Models of Parkinson's Disease. <i>Science</i> , 2007, 317, 516-519.	12.6	995
51	Aggregated τ -Synuclein Mediates Dopaminergic Neurotoxicity In Vivo. <i>Journal of Neuroscience</i> , 2007, 27, 3338-3346.	3.6	271
52	S/P and T/P phosphorylation is critical for tau neurotoxicity in <i>Drosophila</i> . <i>Journal of Neuroscience Research</i> , 2007, 85, 1271-1278.	2.9	108
53	Abnormal bundling and accumulation of F-actin mediates tau-induced neuronal degeneration in vivo. <i>Nature Cell Biology</i> , 2007, 9, 139-148.	10.3	399
54	Oxidative stress mediates tau-induced neurodegeneration in <i>Drosophila</i> . <i>Journal of Clinical Investigation</i> , 2007, 117, 236-245.	8.2	262

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55	Î±-synuclein acts in the nucleus to inhibit histone acetylation and promote neurotoxicity. Human Molecular Genetics, 2006, 15, 3012-3023.	2.9	486
56	TOR-Mediated Cell-Cycle Activation Causes Neurodegeneration in a Drosophila Tauopathy Model. Current Biology, 2006, 16, 230-241.	3.9	251
57	Accelerated Accumulation of Misfolded Prion Protein and Spongiform Degeneration in a Drosophila Model of Gerstmann-Straussler-Scheinker Syndrome. Journal of Neuroscience, 2006, 26, 12408-12414.	3.6	53
58	Î±-Synuclein phosphorylation controls neurotoxicity and inclusion formation in a Drosophila model of Parkinson disease. Nature Neuroscience, 2005, 8, 657-663.	14.8	575
59	Proliferative Potential of Human Astrocytes. Journal of Neuropathology and Experimental Neurology, 2005, 64, 163-169.	1.7	51
60	Cathepsin D-deficient Drosophila recapitulate the key features of neuronal ceroid lipofuscinoses. Neurobiology of Disease, 2005, 19, 194-199.	4.4	68
61	Disease-related phenotypes in a Drosophila model of hereditary spastic paraplegia are ameliorated by treatment with vinblastine. Journal of Clinical Investigation, 2005, 115, 3026-3034.	8.2	99
62	Comparison of pathways controlling toxicity in the eye and brain in Drosophila models of human neurodegenerative diseases. Human Molecular Genetics, 2004, 13, 2011-2018.	2.9	99
63	Yeast genetics targets lipids in Parkinson's disease. Trends in Genetics, 2004, 20, 273-277.	6.7	29
64	Post-transcriptional suppression of pathogenic prion protein expression in Drosophila neurons. Journal of Neurochemistry, 2003, 85, 1614-1623.	3.9	23
65	Parkin. Neuron, 2003, 38, 13-16.	8.1	108
66	Polyglutamines Stop Traffic. Neuron, 2003, 40, 1-2.	8.1	39
67	Gene expression changes presage neurodegeneration in a Drosophila model of Parkinson's disease. Human Molecular Genetics, 2003, 12, 2457-2466.	2.9	111
68	Mitochondrial pathology and apoptotic muscle degeneration in <i>Drosophila parkin</i> mutants. Proceedings of the National Academy of Sciences of the United States of America, 2003, 100, 4078-4083.	7.1	1,117
69	Title is missing!. Current Opinion in Neurology, 2003, 16, 443-449.	3.6	20
70	From fruit fly to bedside. Current Opinion in Neurology, 2003, 16, 443-449.	3.6	83
71	Genetic Modifiers of Tauopathy in Drosophila. Genetics, 2003, 165, 1233-1242.	2.9	237
72	Modelling neurodegenerative diseases in Drosophila: a fruitful approach?. Nature Reviews Neuroscience, 2002, 3, 237-243.	10.2	144

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73	Tauopathy in <i>Drosophila</i> : Neurodegeneration Without Neurofibrillary Tangles. Science, 2001, 293, 711-714.	12.6	868
74	Studying Human Neurodegenerative Diseases in Flies and Worms. Journal of Neuropathology and Experimental Neurology, 2000, 59, 847-856.	1.7	34
75	A <i>Drosophila</i> model of Parkinson's disease. Nature, 2000, 404, 394-398.	27.8	1,927
76	Neurodegenerative disorders with extensive tau pathology: A comparative study and review. Annals of Neurology, 1996, 40, 139-148.	5.3	301
77	The synaptic vesicle protein synaptotagmin promotes formation of filopodia in fibroblasts. Nature, 1993, 364, 537-540.	27.8	63