

Michael W Baughn

List of Publications by Year in descending order

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11
papers

3,171
citations

840776

11
h-index

1281871

11
g-index

11
all docs

11
docs citations

11
times ranked

4453
citing authors

#	ARTICLE	IF	CITATIONS
1	Divergent roles of ALS-linked proteins FUS/TLS and TDP-43 intersect in processing long pre-mRNAs. <i>Nature Neuroscience</i> , 2012, 15, 1488-1497.	14.8	628
2	Targeting RNA Foci in iPSC-Derived Motor Neurons from ALS Patients with a <i>C9ORF72</i> Repeat Expansion. <i>Science Translational Medicine</i> , 2013, 5, 208ra149.	12.4	586
3	Targeted degradation of sense and antisense <i>C9orf72</i> RNA foci as therapy for ALS and frontotemporal degeneration. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, E4530-9.	7.1	508
4	Gain of Toxicity from ALS/FTD-Linked Repeat Expansions in C9ORF72 Is Alleviated by Antisense Oligonucleotides Targeting GGGGCC-Containing RNAs. <i>Neuron</i> , 2016, 90, 535-550.	8.1	437
5	Premature polyadenylation-mediated loss of stathmin-2 is a hallmark of TDP-43-dependent neurodegeneration. <i>Nature Neuroscience</i> , 2019, 22, 180-190.	14.8	345
6	ALS/FTD-Linked Mutation in FUS Suppresses Intra-axonal Protein Synthesis and Drives Disease Without Nuclear Loss-of-Function of FUS. <i>Neuron</i> , 2018, 100, 816-830.e7.	8.1	185
7	Sense-encoded poly-GR dipeptide repeat proteins correlate to neurodegeneration and uniquely co-localize with TDP-43 in dendrites of repeat-expanded C9orf72 amyotrophic lateral sclerosis. <i>Acta Neuropathologica</i> , 2018, 135, 459-474.	7.7	152
8	Sporadic ALS has compartment-specific aberrant exon splicing and altered cellâ€™matrix adhesion biology. <i>Human Molecular Genetics</i> , 2010, 19, 313-328.	2.9	114
9	Frequency and Complexity of De Novo Structural Mutation in Autism. <i>American Journal of Human Genetics</i> , 2016, 98, 667-679.	6.2	88
10	Transcriptomeâ€™pathology correlation identifies interplay between TDP-43 and the expression of its kinase CK1E in sporadic ALS. <i>Acta Neuropathologica</i> , 2018, 136, 405-423.	7.7	69
11	Lack of C9ORF72 coding mutations supports a gain of function for repeat expansions in amyotrophic lateral sclerosis. <i>Neurobiology of Aging</i> , 2013, 34, 2234.e13-2234.e19.	3.1	59