Jacqueline S De Belleroche

List of Publications by Year in Descending Order

33 papers	6,041 citations	22 h-index	33 g-index
33 ext. papers	7,081 ext. citations	9.5 avg, IF	4.74 L-index

#	Paper	IF	Citations
33	Tissue-selective regulation of protein homeostasis and unfolded protein response signalling in sporadic ALS. *Journal of Cellular and Molecular Medicine*, **2020**, 24, 6055-6069	5.6	6
32	Endoplasmic reticulum stress signalling - from basic mechanisms to clinical applications. *FEBS Journal*, **2019**, 286, 241-278	5.7	309
31	Genome-wide Analyses Identify KIF5A as a Novel ALS Gene. *Neuron*, **2018**, 97, 1268-1283.e6	13.9	296
30	Focus on the Role of D-serine and D-amino Acid Oxidase in Amyotrophic Lateral Sclerosis/Motor Neuron Disease (ALS). *Frontiers in Molecular Biosciences*, **2018**, 5, 8	5.6	17
29	Amyotrophic lateral sclerosis (ALS) and Alzheimer's disease (AD) are characterised by differential activation of ER stress pathways: focus on UPR target genes. *Cell Stress and Chaperones*, **2018**, 23, 897-912	4	38
28	ALS-associated missense and nonsense TBK1 mutations can both cause loss of kinase function. *Neurobiology of Aging*, **2018**, 71, 266.e1-266.e10	5.6	44
27	Mutations in the vesicular trafficking protein annexin A11 are associated with amyotrophic lateral sclerosis. *Science Translational Medicine*, **2017**, 9,	17.5	74
26	Characterisation of the pathogenic effects of the in vivo expression of an ALS-linked mutation in D-amino acid oxidase: Phenotype and loss of spinal cord motor neurons. *PLoS ONE*, **2017**, 12, e0188912	3.7	6
25	NEK1 variants confer susceptibility to amyotrophic lateral sclerosis. *Nature Genetics*, **2016**, 48, 1037-42	36.3	149
24	CCNF mutations in amyotrophic lateral sclerosis and frontotemporal dementia. *Nature Communications*, **2016**, 7, 11253	17.4	126
23	Common variants in the chromosome 2p23 region containing the SLC30A3 (ZnT3) gene are associated with schizophrenia in female but not male individuals in a large collection of European samples. *Psychiatry Research*, **2016**, 246, 335-340	9.9	14
22	Experimental approaches for elucidating co-agonist regulation of NMDA receptor in motor neurons: Therapeutic implications for amyotrophic lateral sclerosis (ALS). *Journal of Pharmaceutical and Biomedical Analysis*, **2015**, 116, 2-6	3.5	10
21	The CHCHD10 P34S variant is not associated with ALS in a UK cohort of familial and sporadic patients. *Neurobiology of Aging*, **2015**, 36, 2908.e17-8	5.6	19
20	VCP mutations are not a major cause of familial amyotrophic lateral sclerosis in the UK. *Journal of the Neurological Sciences*, **2015**, 349, 209-13	3.2	9
19	Exome-wide rare variant analysis identifies TUBA4A mutations associated with familial ALS. *Neuron*, **2014**, 84, 324-31	13.9	229
18	Pathogenic effects of amyotrophic lateral sclerosis-linked mutation in D-amino acid oxidase are mediated by D-serine. *Neurobiology of Aging*, **2014**, 35, 876-85	5.6	26
17	Allelic variants in the zinc transporter-3 gene, SLC30A3, a candidate gene identified from gene expression studies, show gender-specific association with schizophrenia. *European Psychiatry*, **2014**, 29, 172-8	6	15

16	The role of D-serine and glycine as co-agonists of NMDA receptors in motor neuron degeneration and amyotrophic lateral sclerosis (ALS). *Frontiers in Synaptic Neuroscience*, **2014**, 6, 10	3.5	38
15	Sequestosome-1 (SQSTM1) sequence variants in ALS cases in the UK: prevalence and coexistence of SQSTM1 mutations in ALS kindred with PDB. *European Journal of Human Genetics*, **2014**, 22, 492-6	5.3	41
14	The C9ORF72 expansion mutation is a common cause of ALS+/-FTD in Europe and has a single founder. *European Journal of Human Genetics*, **2013**, 21, 102-8	5.3	159
13	The role of D-amino acids in amyotrophic lateral sclerosis pathogenesis: a review. *Amino Acids*, **2012** , 43, 1823-31	3.5	38
12	Transcription and pathway analysis of the superior temporal cortex and anterior prefrontal cortex in schizophrenia. *Journal of Neuroscience Research*, **2011**, 89, 1218-27	4.4	55
11	Reply to Millecamps et al.: Elucidating the role of D amino acid oxidase in familial amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences of the United States of America*, **2010**, 107, E108-E108	11.5	78
10	Familial amyotrophic lateral sclerosis is associated with a mutation in D-amino acid oxidase. *Proceedings of the National Academy of Sciences of the United States of America*, **2010**, 107, 7556-61	11.5	191
9	Vesicle associated membrane protein B (VAPB) is decreased in ALS spinal cord. *Neurobiology of Aging*, **2010**, 31, 969-85	5.6	99
8	Thioredoxin reductase 1 haplotypes modify familial amyotrophic lateral sclerosis onset. *Free Radical Biology and Medicine*, **2009**, 46, 202-11	7.8	17
7	Mutations in FUS, an RNA processing protein, cause familial amyotrophic lateral sclerosis type 6. *Science*, **2009**, 323, 1208-1211	33.3	1890
6	Protective effects of heat shock protein 27 in a model of ALS occur in the early stages of disease progression. *Neurobiology of Disease*, **2008**, 30, 42-55	7.5	89
5	TDP-43 mutations in familial and sporadic amyotrophic lateral sclerosis. *Science*, **2008**, 319, 1668-72	33.3	1877
4	Identification of genetic heterogeneity in Refsum□s disease. *European Journal of Human Genetics*, **2000**, 8, 649-51	5.3	33
3	FC09.06 Cholecystokinin CCKb Receptor mRNA Isoforms: Expression in Postmortem Monkey and Human Brain □Alterations Following Schizophrenia. *European Psychiatry*, **2000**, 15, 284s-285s	6	
2	Hypothalamic Hypertensive Factor. *Hypertension*, **1997**, 30, 1493-1498	8.5	10
1	Elevated levels of amino acids in the CSF of motor neuron disease patients. *Neurochemical Pathology*, **1984**, 2, 1-6		39