

Kurt J De Vos

List of Publications by Year in descending order

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45
papers

8,746
citations

117571

34
h-index

243529

44
g-index

49
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49
docs citations

49
times ranked

11403
citing authors

#	ARTICLE	IF	CITATIONS
1	<i>C9ORF72</i> -derived poly-GA DPRs undergo endocytic uptake in astrocytes and spread to motor neurons. <i>Life Science Alliance</i> , 2022, 5, e202101276.	1.3	6
2	Astrocyte adenosine deaminase loss increases motor neuron toxicity in amyotrophic lateral sclerosis. <i>Brain</i> , 2019, 142, 586-605.	3.7	84
3	The role of mitochondria in amyotrophic lateral sclerosis. <i>Neuroscience Letters</i> , 2019, 710, 132933.	1.0	356
4	<i>C9orf72</i> plays a central role in Rab GTPase-dependent regulation of autophagy. <i>Small GTPases</i> , 2018, 9, 399-408.	0.7	45
5	Neurobiology of axonal transport defects in motor neuron diseases: Opportunities for translational research?. <i>Neurobiology of Disease</i> , 2017, 105, 283-299.	2.1	173
6	<i>C9orf72</i> expansion disrupts ATM-mediated chromosomal break repair. <i>Nature Neuroscience</i> , 2017, 20, 1225-1235.	7.1	138
7	SRSF1-dependent nuclear export inhibition of <i>C9ORF72</i> repeat transcripts prevents neurodegeneration and associated motor deficits. <i>Nature Communications</i> , 2017, 8, 16063.	5.8	106
8	Amyotrophic lateral sclerosis-associated mutant SOD1 inhibits anterograde axonal transport of mitochondria by reducing Miro1 levels. <i>Human Molecular Genetics</i> , 2017, 26, 4668-4679.	1.4	83
9	Protein Homeostasis in Amyotrophic Lateral Sclerosis: Therapeutic Opportunities?. <i>Frontiers in Molecular Neuroscience</i> , 2017, 10, 123.	1.4	62
10	<i>C9ORF72</i> hexanucleotide repeat exerts toxicity in a stable, inducible motor neuronal cell model, which is rescued by partial depletion of Pten. <i>Human Molecular Genetics</i> , 2017, 26, 1133-1145.	1.4	23
11	The <i>C9orf72</i> protein interacts with Rab1a and the ULK1 complex to regulate initiation of autophagy. <i>EMBO Journal</i> , 2016, 35, 1656-1676.	3.5	327
12	ALS/FTD-associated FUS activates GSK-3 β to disrupt the VAPB-PTIP51 interaction and ER-mitochondria associations. <i>EMBO Reports</i> , 2016, 17, 1326-1342.	2.0	201
13	Reduced number of axonal mitochondria and tau hypophosphorylation in mouse P301L tau knockin neurons. <i>Neurobiology of Disease</i> , 2016, 85, 1-10.	2.1	57
14	ER-mitochondria associations are regulated by the VAPB-PTIP51 interaction and are disrupted by ALS/FTD-associated TDP-43. <i>Nature Communications</i> , 2014, 5, 3996.	5.8	463
15	Increasing microtubule acetylation rescues axonal transport and locomotor deficits caused by LRRK2 Roc-COR domain mutations. <i>Nature Communications</i> , 2014, 5, 5245.	5.8	229
16	Axonal Transport Defects in a Mitofusin 2 Loss of Function Model of Charcot-Marie-Tooth Disease in Zebrafish. <i>PLoS ONE</i> , 2013, 8, e67276.	1.1	55
17	Amyotrophic lateral sclerosis-associated mutant VAPBP56S perturbs calcium homeostasis to disrupt axonal transport of mitochondria. <i>Human Molecular Genetics</i> , 2012, 21, 1979-1988.	1.4	112
18	VAPB interacts with the mitochondrial protein PTIP51 to regulate calcium homeostasis. <i>Human Molecular Genetics</i> , 2012, 21, 1299-1311.	1.4	423

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19	Lemur tyrosine kinase-2 signalling regulates kinesin-1 light chain-2 phosphorylation and binding of Smad2 cargo. <i>Oncogene</i> , 2012, 31, 2773-2782.	2.6	44
20	A comparison of in vitro properties of resting SOD1 transgenic microglia reveals evidence of reduced neuroprotective function. <i>BMC Neuroscience</i> , 2011, 12, 91.	0.8	19
21	Phosphorylation of kinesin light chain 1 at serine 460 modulates binding and trafficking of calyntenin-1. <i>Journal of Cell Science</i> , 2011, 124, 1032-1042.	1.2	55
22	A14...Fast axonal transport of mitochondria is altered in Huntington's disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2010, 81, A5.1-A5.	0.9	1
23	Deficiency of the Copper Chaperone for Superoxide Dismutase Increases Amyloid- β^2 Production. <i>Journal of Alzheimer's Disease</i> , 2010, 21, 1101-1105.	1.2	23
24	Amyotrophic lateral sclerosis mutant vesicle-associated membrane protein-associated protein-B transgenic mice develop TAR-DNA-binding protein-43 pathology. <i>Neuroscience</i> , 2010, 167, 774-785.	1.1	69
25	Neurofilament subunit (NFL) head domain phosphorylation regulates axonal transport of neurofilaments. <i>European Journal of Cell Biology</i> , 2009, 88, 193-202.	1.6	46
26	Direct evidence for axonal transport defects in a novel mouse model of mutant spastin-induced hereditary spastic paraplegia (HSP) and human HSP patients. <i>Journal of Neurochemistry</i> , 2009, 110, 34-44.	2.1	135
27	Riluzole protects against glutamate-induced slowing of neurofilament axonal transport. <i>Neuroscience Letters</i> , 2009, 454, 161-164.	1.0	34
28	Mutations in FUS, an RNA Processing Protein, Cause Familial Amyotrophic Lateral Sclerosis Type 6. <i>Science</i> , 2009, 323, 1208-1211.	6.0	2,295
29	Role of Axonal Transport in Neurodegenerative Diseases. <i>Annual Review of Neuroscience</i> , 2008, 31, 151-173.	5.0	638
30	Familial amyotrophic lateral sclerosis-linked SOD1 mutants perturb fast axonal transport to reduce axonal mitochondria content. <i>Human Molecular Genetics</i> , 2007, 16, 2720-2728.	1.4	365
31	Cell-Free Assays for Mitochondria-Cytoskeleton Interactions. <i>Methods in Cell Biology</i> , 2007, 80, 683-706.	0.5	2
32	Visualization and Quantification of Mitochondrial Dynamics in Living Animal Cells. <i>Methods in Cell Biology</i> , 2007, 80, 627-682.	0.5	79
33	Therapeutic activity of C5a receptor antagonists in a rat model of neurodegeneration. <i>FASEB Journal</i> , 2006, 20, 1407-1417.	0.2	129
34	Mitochondrial Function and Actin Regulate Dynamin-Related Protein 1-Dependent Mitochondrial Fission. <i>Current Biology</i> , 2005, 15, 678-683.	1.8	320
35	Expression of Phosphatidylinositol (4,5) Bisphosphate-specific Pleckstrin Homology Domains Alters Direction But Not the Level of Axonal Transport of Mitochondria. <i>Molecular Biology of the Cell</i> , 2003, 14, 3636-3649.	0.9	69
36	RPTP- β acts as a transducer of mechanical force on β 3-integrin-cytoskeleton linkages. <i>Journal of Cell Biology</i> , 2003, 161, 143-153.	2.3	194

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37	Tumor Necrosis Factor Induces Hyperphosphorylation of Kinesin Light Chain and Inhibits Kinesin-Mediated Transport of Mitochondria. <i>Journal of Cell Biology</i> , 2000, 149, 1207-1214.	2.3	103
38	Role of Reactive Oxygen Species in Tumor Necrosis Factor Toxicity. , 2000, , 245-264.		0
39	Redox regulation of TNF signaling. <i>BioFactors</i> , 1999, 10, 145-156.	2.6	126
40	Significance of Host Cell Kinesin in the Development of <i>Chlamydia psittaci</i> . <i>Infection and Immunity</i> , 1999, 67, 5441-5446.	1.0	12
41	Atractyloside-induced release of cathepsin B, a protease with caspase-processing activity. <i>FEBS Letters</i> , 1998, 438, 150-158.	1.3	275
42	The 55-kDa Tumor Necrosis Factor Receptor Induces Clustering of Mitochondria through Its Membrane-proximal Region. <i>Journal of Biological Chemistry</i> , 1998, 273, 9673-9680.	1.6	150
43	A Caspase-activated Factor (CAF) Induces Mitochondrial Membrane Depolarization and Cytochrome c Release by a Nonproteolytic Mechanism. <i>Journal of Experimental Medicine</i> , 1998, 188, 2193-2198.	4.2	34
44	Induction of Unresponsiveness to Tumor Necrosis Factor (TNF) after Autocrine TNF Expression Requires TNF Membrane Retention. <i>Journal of Biological Chemistry</i> , 1998, 273, 3271-3277.	1.6	12
45	Direct evidence for tumor necrosis factor-induced mitochondrial reactive oxygen intermediates and their involvement in cytotoxicity.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 1995, 92, 8115-8119.	3.3	566