

Lyle W Ostrow

List of Publications by Year in descending order

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Version: 2024-02-01

38
papers

4,861
citations

279701

23
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37
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all docs

42
docs citations

42
times ranked

6882
citing authors

#	ARTICLE	IF	CITATIONS
1	The C9orf72 repeat expansion disrupts nucleocytoplasmic transport. <i>Nature</i> , 2015, 525, 56-61.	13.7	835
2	RAN proteins and RNA foci from antisense transcripts in <i>C9ORF72</i> ALS and frontotemporal dementia. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, E4968-77.	3.3	681
3	An antisense oligonucleotide against SOD1 delivered intrathecally for patients with SOD1 familial amyotrophic lateral sclerosis: a phase 1, randomised, first-in-man study. <i>Lancet Neurology</i> , The, 2013, 12, 435-442.	4.9	534
4	Genome-wide Analyses Identify KIF5A as a Novel ALS Gene. <i>Neuron</i> , 2018, 97, 1268-1283.e6.	3.8	517
5	Degeneration and impaired regeneration of gray matter oligodendrocytes in amyotrophic lateral sclerosis. <i>Nature Neuroscience</i> , 2013, 16, 571-579.	7.1	485
6	Human endogenous retrovirus-K contributes to motor neuron disease. <i>Science Translational Medicine</i> , 2015, 7, 307ra153.	5.8	369
7	TDP-43 loss and ALS-risk SNPs drive mis-splicing and depletion of UNC13A. <i>Nature</i> , 2022, 603, 131-137.	13.7	188
8	Postmortem Cortex Samples Identify Distinct Molecular Subtypes of ALS: Retrotransposon Activation, Oxidative Stress, and Activated Glia. <i>Cell Reports</i> , 2019, 29, 1164-1177.e5.	2.9	184
9	Validation of serum neurofilaments as prognostic and potential pharmacodynamic biomarkers for ALS. <i>Neurology</i> , 2020, 95, e59-e69.	1.5	119
10	Shared polygenic risk and causal inferences in amyotrophic lateral sclerosis. <i>Annals of Neurology</i> , 2019, 85, 470-481.	2.8	118
11	Diseases of the nERVous system: retrotransposon activity in neurodegenerative disease. <i>Mobile DNA</i> , 2019, 10, 32.	1.3	91
12	Mechanosensation and endothelin in astrocytesâ€™ hypothetical roles in CNS pathophysiology. <i>Brain Research Reviews</i> , 2005, 48, 488-508.	9.1	87
13	Unexpected similarities between C9ORF72 and sporadic forms of ALS/FTD suggest a common disease mechanism. <i>ELife</i> , 2018, 7, .	2.8	53
14	Transcriptional profiling of HERV-K(HML-2) in amyotrophic lateral sclerosis and potential implications for expression of HML-2 proteins. <i>Molecular Neurodegeneration</i> , 2018, 13, 39.	4.4	47
15	Association of Variants in the <i>SPTLC1</i> Gene With Juvenile Amyotrophic Lateral Sclerosis. <i>JAMA Neurology</i> , 2021, 78, 1236.	4.5	46
16	Stretch induced endothelin-1 secretion by adult rat astrocytes involves calcium influx via stretch-activated ion channels (SACs). <i>Biochemical and Biophysical Research Communications</i> , 2011, 410, 81-86.	1.0	43
17	Distinct conformers of transmissible misfolded SOD1 distinguish human SOD1-FALS from other forms of familial and sporadic ALS. <i>Acta Neuropathologica</i> , 2016, 132, 827-840.	3.9	42
18	Nuclear export and translation of circular repeat-containing intronic RNA in C9ORF72-ALS/FTD. <i>Nature Communications</i> , 2021, 12, 4908.	5.8	41

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19	Mechanosensitive Ion Channels as Drug Targets. <i>CNS and Neurological Disorders</i> , 2004, 3, 287-295.	4.3	40
20	Properties of LINE-1 proteins and repeat element expression in the context of amyotrophic lateral sclerosis. <i>Mobile DNA</i> , 2018, 9, 35.	1.3	37
21	Clonally expanded CD8 T cells characterize amyotrophic lateral sclerosis-4. <i>Nature</i> , 2022, 606, 945-952.	13.7	35
22	An integrated multi-omic analysis of iPSC-derived motor neurons from C9ORF72 ALS patients. <i>IScience</i> , 2021, 24, 103221.	1.9	27
23	Loss of TDP-43 function and rimmed vacuoles persist after T cell depletion in a xenograft model of sporadic inclusion body myositis. <i>Science Translational Medicine</i> , 2022, 14, eabi9196.	5.8	27
24	Stretch-Induced Endothelin-1 Production by Astrocytes. <i>Journal of Cardiovascular Pharmacology</i> , 2000, 36, S274-S277.	0.8	26
25	Fatal <i>Exserohilum</i> Meningitis and Central Nervous System Vasculitis After Cervical Epidural Methylprednisolone Injection. <i>Annals of Internal Medicine</i> , 2012, 157, 835.	2.0	24
26	Editorial by concerned physicians: Unintended effect of the orphan drug act on the potential cost of 3,4-diaminopyridine. <i>Muscle and Nerve</i> , 2016, 53, 165-168.	1.0	24
27	Iatrogenic <i>Exserohilum</i> infection of the central nervous system: mycological identification and histopathological findings. <i>Modern Pathology</i> , 2013, 26, 166-170.	2.9	23
28	Loss of RAD-23 Protects Against Models of Motor Neuron Disease by Enhancing Mutant Protein Clearance. <i>Journal of Neuroscience</i> , 2015, 35, 14286-14306.	1.7	23
29	ALS Untangled No. 20: The Deanna Protocol. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2013, 14, 319-323.	1.1	19
30	Expanding the spectrum of monoclonal light chain deposition disease in muscle. <i>Muscle and Nerve</i> , 2012, 45, 755-761.	1.0	15
31	Eastchester clapping sign: A novel test of parietal neglect. <i>Annals of Neurology</i> , 2009, 66, 114-117.	2.8	14
32	Disseminated zoster with paresis in a multiple sclerosis patient treated with dimethyl fumarate. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2016, 3, e203.	3.1	13
33	S-acylation of SOD1, CCS, and a stable SOD1-CCS heterodimer in human spinal cords from ALS and non-ALS subjects. <i>Scientific Reports</i> , 2017, 7, 41141.	1.6	12
34	MG53 Preserves Neuromuscular Junction Integrity and Alleviates ALS Disease Progression. <i>Antioxidants</i> , 2021, 10, 1522.	2.2	6
35	Retromer dysfunction in amyotrophic lateral sclerosis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, .	3.3	5
36	Tonic status and electrodecremental paroxysms in an adult without epilepsy. <i>Epileptic Disorders</i> , 2011, 13, 99-101.	0.7	3

#	ARTICLE	IF	CITATIONS
37	ALSUntangled #63: ketogenic diets. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2023, 24, 159-163.	1.1	3
38	Teaching NeuroImages: Neurolymphomatosis. Neurology, 2019, 93, e1229-e1230.	1.5	0