Philip Van Damme

List of Publications by Year
in descending order

313 papers	21,205 citations	7551 77 h-index	13727 129 g-index
333 all docs	333 docs citations	333 times ranked	21460 citing authors

#	Article	IF	Citations
1	A modified Camel and Cactus Test detects presymptomatic semantic impairment in genetic frontotemporal dementia within the GENFI cohort. Applied Neuropsychology Adult, 2022, 29, 112-119.	0.7	18
2	RNF170 mutation causes autosomal dominant sensory ataxia with variable pyramidal involvement. European Journal of Neurology, 2022, 29, 345-349.	1.7	2
3	A data-driven disease progression model of fluid biomarkers in genetic frontotemporal dementia. Brain, 2022, 145, 1805-1817.	3.7	27
4	Stratifying the Presymptomatic Phase of Genetic Frontotemporal Dementia by Serum <scp>NfL</scp> and <scp>pNfH</scp>: A Longitudinal Multicentre Study. Annals of Neurology, 2022, 91, 33-47.	2.8	21
5	Genome-wide identification of the genetic basis of amyotrophic lateral sclerosis. Neuron, 2022, 110, 992-1008.e11.	3.8	51
6	Structural variation analysis of 6,500 whole genome sequences in amyotrophic lateral sclerosis. Npj Genomic Medicine, 2022, 7, 8.	1.7	23
7	The importance of offering early genetic testing in everyone with amyotrophic lateral sclerosis. Brain, 2022, 145, 1207-1210.	3.7	21
8	Electrodiagnosis of Guillain-Barre syndrome in the International GBS Outcome Study: Differences in methods and reference values. Clinical Neurophysiology, 2022, 138, 231-240.	0.7	7
9	Conceptual framework for the definition of preclinical and prodromal frontotemporal dementia. Alzheimer's and Dementia, 2022, 18, 1408-1423.	0.4	24
10	Structural brain splitting is a hallmark of Granulin-related frontotemporal dementia. Neurobiology of Aging, 2022, , .	1.5	1
11	Genome-wide study of DNA methylation shows alterations in metabolic, inflammatory, and cholesterol pathways in ALS. Science Translational Medicine, 2022, 14, eabj0264.	5.8	38
12	Genome-wide association analyses identify new Brugada syndrome risk loci and highlight a new mechanism of sodium channel regulation in disease susceptibility. Nature Genetics, 2022, 54, 232-239.	9.4	55
13	Clinical trials in pediatric ALS: a TRICALS feasibility study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, 23, 481-488.	1.1	3
14	Insights into the identification of a molecular signature for amyotrophic lateral sclerosis exploiting integrated microRNA profiling of iPSC-derived motor neurons and exosomes. Cellular and Molecular Life Sciences, 2022, 79, 189.	2.4	12
15	Frontotemporal Lobar Degeneration Case with an N-Terminal TUBA4A Mutation Exhibits Reduced TUBA4A Levels in the Brain and TDP-43 Pathology. Biomolecules, 2022, 12, 440.	1.8	5
16	Whole-genome sequencing reveals that variants in the Interleukin 18 Receptor Accessory Protein 3â€²UTR protect against ALS. Nature Neuroscience, 2022, 25, 433-445.	7.1	16
17	Semantic modelling of common data elements for rare disease registries, and a prototype workflow for their deployment over registry data. Journal of Biomedical Semantics, 2022, 13, 9.	0.9	11
18	Respiratory onset of amyotrophic lateral sclerosis in a pregnant woman with a novel SOD1 mutation. European Journal of Neurology, 2022, 29, 1279-1283.	1.7	2

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19	The role of inflammation in neurodegeneration: novel insights into the role of the immune system in C9orf72 HRE-mediated ALS/FTD. Molecular Neurodegeneration, 2022, 17, 22.	4.4	24
20	Prognostic relationship of neurofilaments, CHIT1, YKL-40 and MCP-1 in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 681-682.	0.9	7
21	HDAC3 Inhibition Stimulates Myelination in a CMT1A Mouse Model. Molecular Neurobiology, 2022, 59, 3414-3430.	1.9	7
22	Cellular Stress Induces Nucleocytoplasmic Transport Deficits Independent of Stress Granules. Biomedicines, 2022, 10, 1057.	1.4	5
23	Neuromuscular complications after COVID-19 vaccination: a series of eight patients. Acta Neurologica Belgica, 2022, 122, 753-761.	0.5	9
24	Long-term Safety and Efficacy of Avalglucosidase Alfa in Patients With Late-Onset Pompe Disease. Neurology, 2022, 99, .	1.5	16
25	Characterising ALS disease progression according to El Escorial and Gold Coast criteria. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 865-870.	0.9	10
26	Brain metabolic changes across Kingâ€™s stages in amyotrophic lateral sclerosis: a 18F-2-fluoro-2-deoxy-d-glucose-positron emission tomography study. European Journal of Nuclear Medicine and Molecular Imaging, 2021, 48, 1124-1133.	3.3	10
27	TDP-43 proteinopathies: a new wave of neurodegenerative diseases. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, 86-95.	0.9	174
28	PCYT2 mutations disrupting etherlipid biosynthesis: phenotypes converging on the CDP-ethanolamine pathway. Brain, 2021, 144, e17-e17.	3.7	6
29	Necrosomeâ€positive granulovacuolar degeneration is associated with TDPâ€43 pathological lesions in the hippocampus of ALS/FTLD cases. Neuropathology and Applied Neurobiology, 2021, 47, 328-345.	1.8	15
30	Papillary thyroid carcinoma presenting with severe Guillain-BarrÃ© syndrome. Acta Clinica Belgica, 2021, 76, 236-238.	0.5	0
31	The Effect of <scp>SMN</scp> Gene Dosage on <scp>ALS</scp> Risk and Disease Severity. Annals of Neurology, 2021, 89, 686-697.	2.8	10
32	Progression of Behavioral Disturbances and Neuropsychiatric Symptoms in Patients With Genetic Frontotemporal Dementia. JAMA Network Open, 2021, 4, e2030194.	2.8	42
33	Prognostic value of neurofilament light chain in chronic inflammatory demyelinating polyneuropathy. Brain Communications, 2021, 3, fcab018.	1.5	7
34	STING-Induced Inflammation â€" A Novel Therapeutic Target in ALS?. New England Journal of Medicine, 2021, 384, 765-767.	13.9	6
35	C9orf72 ALS-FTD: recent evidence for dysregulation of the autophagy-lysosome pathway at multiple levels. Autophagy, 2021, 17, 3306-3322.	4.3	52
36	Chitotriosidase as biomarker for early stage amyotrophic lateral sclerosis: a multicenter study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, 22, 276-286.	1.1	14

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37	HDAC6 inhibition restores TDPâ€43 pathology and axonal transport defects in human motor neurons with TARDBP mutations. EMBO Journal, 2021, 40, e106177.	3.5	51
38	AAV9-mediated gene delivery of MCT1 to oligodendrocytes does not provide a therapeutic benefit in a mouse model of ALS. Molecular Therapy - Methods and Clinical Development, 2021, 20, 508-519.	1.8	12
39	C9orf72 -derived arginine-containing dipeptide repeats associate with axonal transport machinery and impede microtubule-based motility. Science Advances, 2021, 7, .	4.7	57
40	Liquidâ€“Liquid Phase Separation Enhances TDP-43 LCD Aggregation but Delays Seeded Aggregation. Biomolecules, 2021, 11, 548.	1.8	18
41	Detection of multiple myositis-specific autoantibodies in unique patients with idiopathic inflammatory myopathy: A single centre-experience and literature review. Seminars in Arthritis and Rheumatism, 2021, 51, 486-494.	1.6	8
42	Altered perivascular fibroblast activity precedes ALS disease onset. Nature Medicine, 2021, 27, 640-646.	15.2	69
43	Distinguishing Primary Lateral Sclerosis from Parkinsonian Syndromes with the Help of Advanced Imaging. Journal of Nuclear Medicine, 2021, 62, 1318-1319.	2.8	1
44	Neuropathy of the phrenic nerve associated with antiganglioside antibodies. European Journal of Neurology, 2021, 28, 2138-2141.	1.7	1
45	The Revised Self-Monitoring Scale detects early impairment of social cognition in genetic frontotemporal dementia within the GENFI cohort. Alzheimer's Research and Therapy, 2021, 13, 127.	3.0	12
46	Tweaking Progranulin Expression: Therapeutic Avenues and Opportunities. Frontiers in Molecular Neuroscience, 2021, 14, 713031.	1.4	28
47	Innovating Clinical Trials for Amyotrophic Lateral Sclerosis. Neurology, 2021, 97, 528-536.	1.5	19
48	Association of Variants in the SPTLC1 Gene With Juvenile Amyotrophic Lateral Sclerosis. JAMA Neurology, 2021, 78, 1236.	4.5	46
49	Human motor units in microfluidic devices are impaired by FUS mutations and improved by HDAC6 inhibition. Stem Cell Reports, 2021, 16, 2213-2227.	2.3	47
50	Psychopathology in premanifest C9orf72 repeat expansion carriers. Journal of Neurology, Neurosurgery and Psychiatry, 2021, , jnnp-2021-327774.	0.9	1
51	Guillain-BarrÃ© syndrome after SARS-CoV-2 infection in an international prospective cohort study. Brain, 2021, 144, 3392-3404.	3.7	39
52	Generation of Human Motor Units with Functional Neuromuscular Junctions in Microfluidic Devices. Journal of Visualized Experiments, 2021, , .	0.2	4
53	Correlations between measures of ALS respiratory function: is there an alternative to FVC?. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2021, 22, 495-504.	1.1	2
54	Eculizumab in refractory generalized myasthenia gravis previously treated with rituximab: subgroup analysis of <scp>REGAIN</scp> and its extension study. Muscle and Nerve, 2021, 64, 662-669.	1.0	11

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55	Safety and efficacy of oral levosimendan in people with amyotrophic lateral sclerosis (the REFALS) Tj ETQq1 1 0.784314 rgBT /Overloc 821-831.	4.9	9
56	Differential early subcortical involvement in genetic FTD within the GENFI cohort. NeuroImage: Clinical, 2021, 30, 102646.	1.4	28
57	Histone Deacetylase Inhibition Regulates Lipid Homeostasis in a Mouse Model of Amyotrophic Lateral Sclerosis. International Journal of Molecular Sciences, 2021, 22, 11224.	1.8	27
58	A double-blind, placebo-controlled, randomized trial of PXT3003 for the treatment of Charcotâ€“Marieâ€“Tooth type 1A. Orphanet Journal of Rare Diseases, 2021, 16, 433.	1.2	23
59	SCFD1 expression quantitative trait loci in amyotrophic lateral sclerosis are differentially expressed. Brain Communications, 2021, 3, fcab236.	1.5	14
60	A panel of CSF proteins separates genetic frontotemporal dementia from presymptomatic mutation carriers: a GENFI study. Molecular Neurodegeneration, 2021, 16, 79.	4.4	9
61	FAIRification Efforts of Clinical Researchers: The Current State of Affairs. Studies in Health Technology and Informatics, 2021, 287, 35-39.	0.2	1
62	Common and rare variant association analyses in amyotrophic lateral sclerosis identify 15 risk loci with distinct genetic architectures and neuron-specific biology. Nature Genetics, 2021, 53, 1636-1648.	9.4	223
63	Investigating the Endo-Lysosomal System in Major Neurocognitive Disorders Due to Alzheimerâ€™s Disease, Frontotemporal Lobar Degeneration and Lewy Body Disease: Evidence for SORL1 as a Cross-Disease Gene. International Journal of Molecular Sciences, 2021, 22, 13633.	1.8	8
64	TSPO Versus P2X7 as a Target for Neuroinflammation: An In Vitro and In Vivo Study. Journal of Nuclear Medicine, 2020, 61, 604-607.	2.8	42
65	Age at symptom onset and death and disease duration in genetic frontotemporal dementia: an international retrospective cohort study. Lancet Neurology, The, 2020, 19, 145-156.	4.9	175
66	RNA-Sequencing Highlights Inflammation and Impaired Integrity of the Vascular Wall in Brain Arteriovenous Malformations. Stroke, 2020, 51, 268-274.	1.0	22
67	Non-invasive characterization of amyotrophic lateral sclerosis in a hTDP-43A315T mouse model: A PET-MR study. NeuroImage: Clinical, 2020, 27, 102327.	1.4	9
68	Placebo effect in chronic inflammatory demyelinating polyneuropathy: The <scp>PATH</scp> study and a systematic review. Journal of the Peripheral Nervous System, 2020, 25, 230-237.	1.4	15
69	TRICALS: creating a highway toward a cure. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 496-501.	1.1	20
70	Rare Variant Burden Analysis within Enhancers Identifies CAV1 as an ALS Risk Gene. Cell Reports, 2020, 33, 108456.	2.9	24
71	Diagnostic yield of testing for RFC1 repeat expansions in patients with unexplained adult-onset cerebellar ataxia. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1233-1234.	0.9	9
72	Consistent improvement with eculizumab across muscle groups in myasthenia gravis. Annals of Clinical and Translational Neurology, 2020, 7, 1327-1339.	1.7	16

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73	Dipeptide repeat protein and TDP-43 pathology along the hypothalamicâ€“pituitary axis in C9orf72 and non-C9orf72 ALS and FTLD-TDP cases. Acta Neuropathologica, 2020, 140, 777-781.	3.9	8
74	Use of Multimodal Imaging and Clinical Biomarkers in Presymptomatic Carriers of C9orf72 Repeat Expansion. JAMA Neurology, 2020, 77, 1008.	4.5	45
75	Amyotrophic lateral sclerosis: a clinical review. European Journal of Neurology, 2020, 27, 1918-1929.	1.7	451
76	A multi-center study of neurofilament assay reliability and inter-laboratory variability. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2020, 21, 452-458.	1.1	15
77	Moving Toward Multicenter Therapeutic Trials in Amyotrophic Lateral Sclerosis: Feasibility of Data Pooling Using Different Translocator Protein PET Radioligands. Journal of Nuclear Medicine, 2020, 61, 1621-1627.	2.8	22
78	Neurofilament light chain and C reactive protein explored as predictors of survival in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 436-437.	0.9	25
79	Phase 1â€“2 Trial of Antisense Oligonucleotide Tofersen for SOD1 ALS. New England Journal of Medicine, 2020, 383, 109-119.	13.9	354
80	Myositis as a neuromuscular complication of immune checkpoint inhibitors. Acta Neurologica Belgica, 2020, 120, 355-364.	0.5	17
81	Combined brain and spinal FDG PET allows differentiation between ALS and ALS mimics. European Journal of Nuclear Medicine and Molecular Imaging, 2020, 47, 2681-2690.	3.3	15
82	Late-onset Pompe disease (LOPD) in Belgium: clinical characteristics and outcome measures. Orphanet Journal of Rare Diseases, 2020, 15, 83.	1.2	26
83	ATXN1 repeat expansions confer risk for amyotrophic lateral sclerosis and contribute to TDP-43 mislocalization. Brain Communications, 2020, 2, fcaa064.	1.5	33
84	Intracerebroventricular delivery of vascular endothelial growth factor in patients with amyotrophic lateral sclerosis, a phase I study. Brain Communications, 2020, 2, fcaa160.	1.5	16
85	Is there a glucose metabolic signature of spreading TDP-43 pathology in amyotrophic lateral sclerosis?. Quarterly Journal of Nuclear Medicine and Molecular Imaging, 2020, 64, 96-104.	0.4	6
86	Knowledge, attitudes and behaviours towards vaccination: a survey of university students in Europe. European Journal of Public Health, 2020, 30, .	0.1	0
87	Quantitative Nucleocytoplasmic Transport Assays in Cellular Models of Neurodegeneration. Bio-protocol, 2020, 10, e3659.	0.2	2
88	Serum neurofilament light chain levels as a marker of upper motor neuron degeneration in patients with Amyotrophic Lateral Sclerosis. Neuropathology and Applied Neurobiology, 2019, 45, 291-304.	1.8	82
89	Long-term efficacy and safety of eculizumab in Japanese patients with generalized myasthenia gravis: A subgroup analysis of the REGAIN open-label extension study. Journal of the Neurological Sciences, 2019, 407, 116419.	0.3	18
90	Dystrophin deficiency leads to dysfunctional glutamate clearance in iPSC derived astrocytes. Translational Psychiatry, 2019, 9, 200.	2.4	18

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91	Reduction of ephrin-A5 aggravates disease progression in amyotrophic lateral sclerosis. Acta Neuropathologica Communications, 2019, 7, 114.	2.4	11
92	Restoration of histone acetylation ameliorates disease and metabolic abnormalities in a FUS mouse model. Acta Neuropathologica Communications, 2019, 7, 107.	2.4	61
93	P.69NEO1 and NEO-EXT studies: exploratory efficacy of repeat avalglucosidase alfa dosing for up to 5 years in participants with late-onset Pompe disease (LOPD). Neuromuscular Disorders, 2019, 29, S60-S61.	0.3	0
94	C9orf72-generated poly-GR and poly-PR do not directly interfere with nucleocytoplasmic transport. Scientific Reports, 2019, 9, 15728.	1.6	47
95	Phenotypes and malignancy risk of different FUS mutations in genetic amyotrophic lateral sclerosis. Annals of Clinical and Translational Neurology, 2019, 6, 2384-2394.	1.7	49
96	Serum neurofilament heavy chains as early marker of motor neuron degeneration. Annals of Clinical and Translational Neurology, 2019, 6, 1971-1979.	1.7	29
97	A Phase 3 Multicenter, Prospective, Open-Label Efficacy and Safety Study of Immune Globulin (Human) 10% Caprylate/Chromatography Purified in Patients with Myasthenia Gravis Exacerbations. European Neurology, 2019, 81, 223-230.	0.6	23
98	Long-term safety and efficacy of subcutaneous immunoglobulin IgPro20 in CIDP. Neurology: Neuroimmunology and NeuroInflammation, 2019, 6, e590.	3.1	37
99	Differentiation but not ALS mutations in FUS rewires motor neuron metabolism. Nature Communications, 2019, 10, 4147.	5.8	41
100	Reducing EphA4 before disease onset does not affect survival in a mouse model of Amyotrophic Lateral Sclerosis. Scientific Reports, 2019, 9, 14112.	1.6	10
101	Efficacy and safety of IVIG in CIDP: Combined data of the PRIMA and PATH studies. Journal of the Peripheral Nervous System, 2019, 24, 48-55.	1.4	17
102	Restabilization treatment after intravenous immunoglobulin withdrawal in chronic inflammatory demyelinating polyneuropathy: Results from the preâ€randomization phase of the Polyneuropathy And Treatment with Hizentra study. Journal of the Peripheral Nervous System, 2019, 24, 72-79.	1.4	13
103	Analytical performance of a CE-marked immunoassay to quantify phosphorylated neurofilament heavy chains. Clinical Chemistry and Laboratory Medicine, 2019, 57, e199-e202.	1.4	1
104	Randomized phase 2 study of FcRn antagonist efgartigimod in generalized myasthenia gravis. Neurology, 2019, 92, e2661-e2673.	1.5	169
105	Inflammatory markers in cerebrospinal fluid: independent prognostic biomarkers in amyotrophic lateral sclerosis?. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, jnnp-2018-319586.	0.9	42
106	A phase III trial of tirasemtiv as a potential treatment for amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 584-594.	1.1	29
107	Motor cortex metabolite alterations in amyotrophic lateral sclerosis assessed in vivo using edited and non-edited magnetic resonance spectroscopy. Brain Research, 2019, 1718, 22-31.	1.1	24
108	NEO1 and NEO-EXT studies: Long-term safety of repeat avalglucosidase alfa dosing for 4.5 years in late-onset Pompe disease patients. Molecular Genetics and Metabolism, 2019, 126, S115-S116.	0.5	0

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109	An ALS case with 38 (G4C2)-repeats in the C9orf72 gene shows TDP-43 and sparse dipeptide repeat protein pathology. Acta Neuropathologica, 2019, 137, 855-858.	3.9	12
110	Microglia lacking a peroxisomal Î²-oxidation enzyme chronically alter their inflammatory profile without evoking neuronal and behavioral deficits. Journal of Neuroinflammation, 2019, 16, 61.	3.1	20
111	Exome array analysis of rare and low frequency variants in amyotrophic lateral sclerosis. Scientific Reports, 2019, 9, 5931.	1.6	16
112	Shared polygenic risk and causal inferences in amyotrophic lateral sclerosis. Annals of Neurology, 2019, 85, 470-481.	2.8	118
113	Mutations in the Glycosyltransferase Domain of GLT8D1 Are Associated with Familial Amyotrophic Lateral Sclerosis. Cell Reports, 2019, 26, 2298-2306.e5.	2.9	57
114	Clinical spectrum of the anti-GQ1b antibody syndrome: a case series of eight patients. Acta Neurologica Belgica, 2019, 119, 29-36.	0.5	17
115	Longâ€term safety and efficacy of eculizumab in generalized myasthenia gravis. Muscle and Nerve, 2019, 60, 14-24.	1.0	162
116	AB0696â€…DETECTION OF COEXISTING MYOSITIS-SPECIFIC AUTOANTIBODIES WITH LINE AND DOT IMMUNOASSAYS IN PATIENTS WITH IDIOPATHIC INFLAMMATORY MYOPATHIES. , 2019, , .		0
117	Circadian sleep/wake-associated cells show dipeptide repeat protein aggregates in C9orf72-related ALS and FTLD cases. Acta Neuropathologica Communications, 2019, 7, 189.	2.4	22
118	White matter hyperintensities in progranulin-associated frontotemporal dementia: A longitudinal GENFI study. NeuroImage: Clinical, 2019, 24, 102077.	1.4	27
119	EphA4 loss improves social memory performance and alters dendritic spine morphology without changes in amyloid pathology in a mouse model of Alzheimerâ€™s disease. Alzheimer's Research and Therapy, 2019, 11, 102.	3.0	17
120	Lowering EphA4 Does Not Ameliorate Disease in a Mouse Model for Severe Spinal Muscular Atrophy. Frontiers in Neuroscience, 2019, 13, 1233.	1.4	2
121	Safety, tolerability, pharmacokinetics, pharmacodynamics, and exploratory efficacy of the novel enzyme replacement therapy avalglucosidase alfa (neoGAA) in treatment-naÃ¯ve and alglucosidase alfa-treated patients with late-onset Pompe disease: A phase 1, open-label, multicenter, multinational, ascending dose study. Neuromuscular Disorders, 2019, 29, 167-186.	0.3	59
122	Association of NIPA1 repeat expansions with amyotrophic lateral sclerosis in a large international cohort. Neurobiology of Aging, 2019, 74, 234.e9-234.e15.	1.5	26
123	Derivation of norms for the Dutch version of the Edinburgh cognitive and behavioral ALS screen. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 19-27.	1.1	17
124	Detection of myositis-specific antibodies. Annals of the Rheumatic Diseases, 2019, 78, e7-e7.	0.5	48
125	FUS (fused in sarcoma) is a component of the cellular response to topoisomerase Iâ€“induced DNA breakage and transcriptional stress. Life Science Alliance, 2019, 2, e201800222.	1.3	20
126	Anterior interosseous mononeuropathy associated with HEV infection. Neurology: Neuroimmunology and NeuroInflammation, 2018, 5, e429.	3.1	2

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127	Intravenous versus subcutaneous immunoglobulin â€“ Authors' reply. Lancet Neurology, The, 2018, 17, 393-394.	4.9	0
128	Conditional deletion of Id2 or Notch1 in oligodendrocyte progenitor cells does not ameliorate disease outcome in SOD1G93A mice. Neurobiology of Aging, 2018, 68, 1-4.	1.5	16
129	How much of the missing heritability of ALS is hidden in known ALS genes?. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 794-794.	0.9	6
130	HDAC6 is a therapeutic target in mutant GARS-induced Charcot-Marie-Tooth disease. Brain, 2018, 141, 673-687.	3.7	93
131	Elongator subunit 3 (ELP3) modifies ALS through tRNA modification. Human Molecular Genetics, 2018, 27, 1276-1289.	1.4	56
132	Inhibition of histone deacetylase 6 (HDAC6) protects against vincristine-induced peripheral neuropathies and inhibits tumor growth. Neurobiology of Disease, 2018, 111, 59-69.	2.1	52
133	A zebrafish model for C9orf72 ALS reveals RNA toxicity as a pathogenic mechanism. Acta Neuropathologica, 2018, 135, 427-443.	3.9	98
134	Prognosis for patients with amyotrophic lateral sclerosis: development and validation of a personalised prediction model. Lancet Neurology, The, 2018, 17, 423-433.	4.9	342
135	Genome-wide Analyses Identify KIF5A as a Novel ALS Gene. Neuron, 2018, 97, 1268-1283.e6.	3.8	517
136	Comparison of elevated phosphorylated neurofilament heavy chains in serum and cerebrospinal fluid of patients with amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 367-373.	0.9	86
137	NEK1 genetic variability in a Belgian cohort of ALS and ALS-FTD patients. Neurobiology of Aging, 2018, 61, 255.e1-255.e7.	1.5	32
138	Reconsidering the causality of TIA1 mutations in ALS. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2018, 19, 1-3.	1.1	22
139	Multicenter evaluation of neurofilaments in early symptom onset amyotrophic lateral sclerosis. Neurology, 2018, 90, e22-e30.	1.5	148
140	Subcutaneous immunoglobulin for maintenance treatment in chronic inflammatory demyelinating polyneuropathy (PATH): a randomised, double-blind, placebo-controlled, phase 3 trial. Lancet Neurology, The, 2018, 17, 35-46.	4.9	193
141	REGISTRIES AND CARE OF NEUROMUSCULAR DISORDERS. Neuromuscular Disorders, 2018, 28, S117.	0.3	0
142	Glucose metabolic brain patterns to discriminate amyotrophic lateral sclerosis from Parkinson plus syndromes. EJNMMI Research, 2018, 8, 110.	1.1	7
143	Non-invasive assessment of disease progression and neuroprotective effects of dietary coconut oil supplementation in the ALS SOD1G93A mouse model: A 1H-magnetic resonance spectroscopic study. NeuroImage: Clinical, 2018, 20, 1092-1105.	1.4	14
144	Regional variation of Guillain-BarrÃ© syndrome. Brain, 2018, 141, 2866-2877.	3.7	190

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145	Progranulin reduces insoluble TDP-43 levels, slows down axonal degeneration and prolongs survival in mutant TDP-43 mice. Molecular Neurodegeneration, 2018, 13, 55.	4.4	38
146	Low-Frequency and Rare-Coding Variation Contributes to Multiple Sclerosis Risk. Cell, 2018, 175, 1679-1687.e7.	13.5	115
147	Genetic Architecture of Adaptive Immune System Identifies Key Immune Regulators. Cell Reports, 2018, 25, 798-810.e6.	2.9	36
148	FUS-induced neurotoxicity in Drosophila is prevented by downregulating nucleocytoplasmic transport proteins. Human Molecular Genetics, 2018, 27, 4103-4116.	1.4	33
149	Topographical Guidance of PSC-Derived Cortical Neurons. Journal of Nanomaterials, 2018, 2018, 1-10.	1.5	3
150	From lexical regularities to axiomatic patterns for the quality assurance of biomedical terminologies and ontologies. Journal of Biomedical Informatics, 2018, 84, 59-74.	2.5	11
151	Project MinE: study design and pilot analyses of a large-scale whole-genome sequencing study in amyotrophic lateral sclerosis. European Journal of Human Genetics, 2018, 26, 1537-1546.	1.4	129
152	Astrocyte-derived Jagged-1 mitigates deleterious Notch signaling in amyotrophic lateral sclerosis. Neurobiology of Disease, 2018, 119, 26-40.	2.1	35
153	Generation of a human induced pluripotent stem cellâ€“based model for tauopathies combining three microtubuleâ€associated protein TAU mutations which displays several phenotypes linked to neurodegeneration. Alzheimer's and Dementia, 2018, 14, 1261-1280.	0.4	41
154	Oral fingolimod for chronic inflammatory demyelinating polyradiculoneuropathy (FORCIDP Trial): a double-blind, multicentre, randomised controlled trial. Lancet Neurology, The, 2018, 17, 689-698.	4.9	48
155	In Vivo Electrophysiological Measurement of Compound Muscle Action Potential from the Forelimbs in Mouse Models of Motor Neuron Degeneration. Journal of Visualized Experiments, 2018, , .	0.2	12
156	Molecular Dissection of FUS Points at Synergistic Effect of Low-Complexity Domains in Toxicity. Cell Reports, 2018, 24, 529-537.e4.	2.9	74
157	Multicenter validation of [¹⁸F]-FDG PET and support-vector machine discriminant analysis in automatically classifying patients with amyotrophic lateral sclerosis versus controls. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2018, 19, 570-577.	1.1	19
158	No supportive evidence for TIA1 gene mutations in a European cohort of ALS-FTD spectrum patients. Neurobiology of Aging, 2018, 69, 293.e9-293.e11.	1.5	15
159	Diagnostic and Prognostic Performance of Neurofilaments in ALS. Frontiers in Neurology, 2018, 9, 1167.	1.1	100
160	ATXN2 trinucleotide repeat length correlates with risk of ALS. Neurobiology of Aging, 2017, 51, 178.e1-178.e9.	1.5	86
161	Investigating the role of ALS genes CHCHD10 and TUBA4A in Belgian FTD-ALS spectrum patients. Neurobiology of Aging, 2017, 51, 177.e9-177.e16.	1.5	60
162	Safety and efficacy of ozanezumab in patients with amyotrophic lateral sclerosis: a randomised, double-blind, placebo-controlled, phase 2 trial. Lancet Neurology, The, 2017, 16, 208-216.	4.9	62

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