

Andreas Hermann

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

193
papers

4,758
citations

35
h-index

63
g-index

221
ext. papers

5,900
ext. citations

5.7
avg, IF

5.31
L-index

#	Paper	IF	Citations
193	Direct reprogramming of fibroblasts into neural stem cells by defined factors. <i>Cell Stem Cell</i> , 2012 , 10, 465-72	18	441
192	Efficient generation of neural stem cell-like cells from adult human bone marrow stromal cells. <i>Journal of Cell Science</i> , 2004 , 117, 4411-22	5.3	366
191	A star-PEG-heparin hydrogel platform to aid cell replacement therapies for neurodegenerative diseases. <i>Biomaterials</i> , 2009 , 30, 5049-60	15.6	254
190	Transcriptional profiles of CD133+ and CD133- glioblastoma-derived cancer stem cell lines suggest different cells of origin. <i>Cancer Research</i> , 2010 , 70, 2030-40	10.1	209
189	Impaired DNA damage response signaling by FUS-NLS mutations leads to neurodegeneration and FUS aggregate formation. <i>Nature Communications</i> , 2018 , 9, 335	17.4	139
188	A randomized, double blind, placebo-controlled trial of pioglitazone in combination with riluzole in amyotrophic lateral sclerosis. <i>PLoS ONE</i> , 2012 , 7, e37885	3.7	106
187	Comparative analysis of neuroectodermal differentiation capacity of human bone marrow stromal cells using various conversion protocols. <i>Journal of Neuroscience Research</i> , 2006 , 83, 1502-14	4.4	106
186	Nusinersen in adults with 5q spinal muscular atrophy: a non-interventional, multicentre, observational cohort study. <i>Lancet Neurology</i> , 2020 , 19, 317-325	24.1	98
185	Human iPSC models of neuronal ceroid lipofuscinosis capture distinct effects of TPP1 and CLN3 mutations on the endocytic pathway. <i>Human Molecular Genetics</i> , 2014 , 23, 2005-22	5.6	95
184	Intrathecal application of neuroectodermally converted stem cells into a mouse model of ALS: limited intraparenchymal migration and survival narrows therapeutic effects. <i>Journal of Neural Transmission</i> , 2007 , 114, 1395-406	4.3	93
183	C9ORF72 interaction with cofilin modulates actin dynamics in motor neurons. <i>Nature Neuroscience</i> , 2016 , 19, 1610-1618	25.5	87
182	Multipotent neural stem cells from the adult tegmentum with dopaminergic potential develop essential properties of functional neurons. <i>Stem Cells</i> , 2006 , 24, 949-64	5.8	71
181	4-Aminopyridine Induced Activity Rescues Hypoexcitable Motor Neurons from Amyotrophic Lateral Sclerosis Patient-Derived Induced Pluripotent Stem Cells. <i>Stem Cells</i> , 2016 , 34, 1563-75	5.8	65
180	Isogenic FUS-eGFP iPSC Reporter Lines Enable Quantification of FUS Stress Granule Pathology that Is Rescued by Drugs Inducing Autophagy. <i>Stem Cell Reports</i> , 2018 , 10, 375-389	8	64
179	Direct conversion of mouse fibroblasts into induced neural stem cells. <i>Nature Protocols</i> , 2014 , 9, 871-81	18.8	63
178	Chorein-sensitive polymerization of cortical actin and suicidal cell death in chorea-acanthocytosis. <i>FASEB Journal</i> , 2012 , 26, 1526-34	0.9	60
177	Mesodermal cell types induce neurogenesis from adult human hippocampal progenitor cells. <i>Journal of Neurochemistry</i> , 2006 , 98, 629-40	6	57

176	Alterations in the hypothalamic melanocortin pathway in amyotrophic lateral sclerosis. <i>Brain</i> , 2016 , 139, 1106-22	11.2	56
175	Diagnosis and treatment of chorea syndromes. <i>Current Neurology and Neuroscience Reports</i> , 2015 , 15, 514	6.6	56
174	Genotype-phenotype study in patients with valosin-containing protein mutations associated with multisystem proteinopathy. <i>Clinical Genetics</i> , 2018 , 93, 119-125	4	55
173	Restorative approaches in Parkinson's Disease: which cell type wins the race?. <i>Journal of the Neurological Sciences</i> , 2010 , 289, 93-103	3.2	51
172	Transcription profiling of adult and fetal human neuroprogenitors identifies divergent paths to maintain the neuroprogenitor cell state. <i>Stem Cells</i> , 2007 , 25, 1231-40	5.8	51
171	Age-dependent neuroectodermal differentiation capacity of human mesenchymal stromal cells: limitations for autologous cell replacement strategies. <i>Cytotherapy</i> , 2010 , 12, 17-30	4.8	49
170	Antibodies inhibit transmission and aggregation of poly-GA dipeptide repeat proteins. <i>EMBO Molecular Medicine</i> , 2017 , 9, 687-702	12	48
169	Isolation of neural crest derived chromaffin progenitors from adult adrenal medulla. <i>Stem Cells</i> , 2009 , 27, 2602-13	5.8	47
168	FUS Mislocalization and Vulnerability to DNA Damage in ALS Patients Derived hiPSCs and Aging Motoneurons. <i>Frontiers in Cellular Neuroscience</i> , 2016 , 10, 290	6.1	44
167	Directed growth of adult human white matter stem cell-derived neurons on aligned fibrillar collagen. <i>Tissue Engineering - Part A</i> , 2010 , 16, 1103-13	3.9	42
166	Effect of High-Caloric Nutrition on Survival in Amyotrophic Lateral Sclerosis. <i>Annals of Neurology</i> , 2020 , 87, 206-216	9.4	42
165	Nerve ultrasound in the differentiation of multifocal motor neuropathy (MMN) and amyotrophic lateral sclerosis with predominant lower motor neuron disease (ALS/LMND). <i>Journal of Neurology</i> , 2016 , 263, 35-44	5.5	41
164	Age-dependent neurodegeneration and organelle transport deficiencies in mutant TDP43 patient-derived neurons are independent of TDP43 aggregation. <i>Neurobiology of Disease</i> , 2018 , 115, 167-181	7.5	41
163	Stepwise acquirement of hallmark neuropathology in FUS-ALS iPSC models depends on mutation type and neuronal aging. <i>Neurobiology of Disease</i> , 2015 , 82, 420-429	7.5	40
162	Differentiation efficiency of induced pluripotent stem cells depends on the number of reprogramming factors. <i>Stem Cells</i> , 2012 , 30, 570-9	5.8	40
161	Chorein sensitivity of cytoskeletal organization and degranulation of platelets. <i>FASEB Journal</i> , 2013 , 27, 2799-806	0.9	38
160	Rostro-caudal gradual loss of cellular diversity within the periventricular regions of the ventricular system. <i>Stem Cells</i> , 2009 , 27, 928-41	5.8	37
159	A new molecular link between defective autophagy and erythroid abnormalities in chorea-acanthocytosis. <i>Blood</i> , 2016 , 128, 2976-2987	2.2	36

158	Cerebrospinal fluid promotes survival and astroglial differentiation of adult human neural progenitor cells but inhibits proliferation and neuronal differentiation. <i>BMC Neuroscience</i> , 2010 , 11, 48	3.2	35
157	Epigenetic conversion of human adult bone mesodermal stromal cells into neuroectodermal cell types for replacement therapy of neurodegenerative disorders. <i>Expert Opinion on Biological Therapy</i> , 2006 , 6, 653-70	5.4	35
156	Current state of knowledge in Chorea-Acanthocytosis as core Neuroacanthocytosis syndrome. <i>European Journal of Medical Genetics</i> , 2018 , 61, 699-705	2.6	34
155	Physical activity and environmental enrichment regulate the generation of neural precursors in the adult mouse substantia nigra in a dopamine-dependent manner. <i>BMC Neuroscience</i> , 2012 , 13, 132	3.2	33
154	Neuronal Dysfunction in iPSC-Derived Medium Spiny Neurons from Chorea-Acanthocytosis Patients Is Reversed by Src Kinase Inhibition and F-Actin Stabilization. <i>Journal of Neuroscience</i> , 2016 , 36, 12027-12043	6.6	32
153	Safety and efficacy of rasagiline as an add-on therapy to riluzole in patients with amyotrophic lateral sclerosis: a randomised, double-blind, parallel-group, placebo-controlled, phase 2 trial. <i>Lancet Neurology</i> , 2018 , 17, 681-688	24.1	30
152	The TGF- β System As a Potential Pathogenic Player in Disease Modulation of Amyotrophic Lateral Sclerosis. <i>Frontiers in Neurology</i> , 2017 , 8, 669	4.1	30
151	Small molecules for modulating protein driven liquid-liquid phase separation in treating neurodegenerative disease		30
150	Eye-tracking-based assessment suggests preserved well-being in locked-in patients. <i>Annals of Neurology</i> , 2017 , 81, 310-315	9.4	29
149	Knocking out C9ORF72 Exacerbates Axonal Trafficking Defects Associated with Hexanucleotide Repeat Expansion and Reduces Levels of Heat Shock Proteins. <i>Stem Cell Reports</i> , 2020 , 14, 390-405	8	29
148	Reduced hnRNPA3 increases C9orf72 repeat RNA levels and dipeptide-repeat protein deposition. <i>EMBO Reports</i> , 2016 , 17, 1314-25	6.5	29
147	Motor Unit Number Index (MUNIX) of hand muscles is a disease biomarker for adult spinal muscular atrophy. <i>Clinical Neurophysiology</i> , 2019 , 130, 315-319	4.3	29
146	Neurofilaments and tau in CSF in an infant with SMA type 1 treated with nusinersen. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019 , 90, 1068-1069	5.5	27
145	Prognostic factors in ALS: a comparison between Germany and China. <i>Journal of Neurology</i> , 2019 , 266, 1516-1525	5.5	27
144	Therapeutic decisions in ALS patients: cross-cultural differences and clinical implications. <i>Journal of Neurology</i> , 2018 , 265, 1600-1606	5.5	27
143	The concept and diagnostic criteria of primary lateral sclerosis. <i>Acta Neurologica Scandinavica</i> , 2017 , 136, 204-211	3.8	27
142	Neurorestoration in Parkinson's disease by cell replacement and endogenous regeneration. <i>Expert Opinion on Biological Therapy</i> , 2004 , 4, 131-43	5.4	27
141	Clinical features and differential diagnosis of flail arm syndrome. <i>Journal of Neurology</i> , 2016 , 263, 390-395	3.5	26

140	Phenotypes and malignancy risk of different FUS mutations in genetic amyotrophic lateral sclerosis. <i>Annals of Clinical and Translational Neurology</i> , 2019 , 6, 2384-2394	5.3	25
139	Non-Motor Symptoms in Patients Suffering from Motor Neuron Diseases. <i>Frontiers in Neurology</i> , 2016 , 7, 117	4.1	25
138	Poly-glycine-alanine exacerbates C9orf72 repeat expansion-mediated DNA damage via sequestration of phosphorylated ATM and loss of nuclear hnRNPA3. <i>Acta Neuropathologica</i> , 2020 , 139, 99-118	14.3	25
137	Neurochemical markers in CSF of adolescent and adult SMA patients undergoing nusinersen treatment. <i>Therapeutic Advances in Neurological Disorders</i> , 2019 , 12, 1756286419846058	6.6	24
136	Peripheral proinflammatory Th1/Th17 immune cell shift is linked to disease severity in amyotrophic lateral sclerosis. <i>Scientific Reports</i> , 2020 , 10, 5941	4.9	24
135	Reduced intraepidermal nerve fiber density in patients with REM sleep behavior disorder. <i>Parkinsonism and Related Disorders</i> , 2016 , 29, 10-6	3.6	24
134	Chorein Sensitive Arrangement of Cytoskeletal Architecture. <i>Cellular Physiology and Biochemistry</i> , 2015 , 37, 399-408	3.9	24
133	Oligodendrocytes enforce immune tolerance of the uninfected brain by purging the peripheral repertoire of autoreactive CD8+ T cells. <i>Immunity</i> , 2012 , 37, 134-46	32.3	24
132	Induced neural stem cells (iNSCs) in neurodegenerative diseases. <i>Journal of Neural Transmission</i> , 2013 , 120 Suppl 1, S19-25	4.3	22
131	Depending on the stress, histone deacetylase inhibitors act as heat shock protein co-inducers in motor neurons and potentiate arimoclomol, exerting neuroprotection through multiple mechanisms in ALS models. <i>Cell Stress and Chaperones</i> , 2020 , 25, 173-191	4	22
130	CHCHD10 mutations p.R15L and p.G66V cause motoneuron disease by haploinsufficiency. <i>Human Molecular Genetics</i> , 2018 , 27, 706-715	5.6	21
129	Altered calcium dynamics and glutamate receptor properties in iPSC-derived motor neurons from ALS patients with C9orf72, FUS, SOD1 or TDP43 mutations. <i>Human Molecular Genetics</i> , 2019 , 28, 2835-2850	5.6	20
128	Provision of assistive technology devices among people with ALS in Germany: a platform-case management approach. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2018 , 19, 342-350	3.6	20
127	Genome-wide expression profiling and functional network analysis upon neuroectodermal conversion of human mesenchymal stem cells suggest HIF-1 and miR-124a as important regulators. <i>Experimental Cell Research</i> , 2010 , 316, 2760-78	4.2	20
126	Communication Matters-Pitfalls and Promise of Hightech Communication Devices in Palliative Care of Severely Physically Disabled Patients With Amyotrophic Lateral Sclerosis. <i>Frontiers in Neurology</i> , 2018 , 9, 603	4.1	20
125	Human adult white matter progenitor cells are multipotent neuroprogenitors similar to adult hippocampal progenitors. <i>Stem Cells Translational Medicine</i> , 2014 , 3, 458-69	6.9	18
124	Parkinson's disease-like midbrain hyperechogenicity is frequent in amyotrophic lateral sclerosis. <i>Journal of Neurology</i> , 2013 , 260, 454-7	5.5	18
123	Proton MR spectroscopy of neural stem cells: does the proton-NMR peak at 1.28 ppm function as a biomarker for cell type or state?. <i>Rejuvenation Research</i> , 2011 , 14, 371-81	2.6	18

122	Differentiation of chromaffin progenitor cells to dopaminergic neurons. <i>Cell Transplantation</i> , 2012 , 21, 2471-86	4	18
121	Rapid and label-free classification of human glioma cells by infrared spectroscopic imaging. <i>Cytometry Part A: the Journal of the International Society for Analytical Cytology</i> , 2008 , 73A, 1158-64	4.6	18
120	Neurofilament light chain in serum of adolescent and adult SMA patients under treatment with nusinersen. <i>Journal of Neurology</i> , 2020 , 267, 36-44	5.5	18
119	Lithium Sensitivity of Store Operated Ca ²⁺ Entry and Survival of Fibroblasts Isolated from Chorea-Acanthocytosis Patients. <i>Cellular Physiology and Biochemistry</i> , 2017 , 42, 2066-2077	3.9	17
118	Human TDP-43 and FUS selectively affect motor neuron maturation and survival in a murine cell model of ALS by non-cell-autonomous mechanisms. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2015 , 16, 431-41	3.6	17
117	Perivascular Mesenchymal Stem Cells From the Adult Human Brain Harbor No Intrinsic Neuroectodermal but High Mesodermal Differentiation Potential. <i>Stem Cells Translational Medicine</i> , 2015 , 4, 1223-33	6.9	16
116	To die or not to die SGK1-sensitive ORAI/STIM in cell survival. <i>Cell Calcium</i> , 2018 , 74, 29-34	4	16
115	Alterations of red cell membrane properties in neuroacanthocytosis. <i>PLoS ONE</i> , 2013 , 8, e76715	3.7	16
114	Combined anti-platelet therapy with aspirin and clopidogrel: risk factor for thrombolysis-related intracerebral hemorrhage in acute ischemic stroke?. <i>Journal of the Neurological Sciences</i> , 2009 , 284, 155-7	3.2	16
113	SQSTM1/p62 variants in 486 patients with familial ALS from Germany and Sweden. <i>Neurobiology of Aging</i> , 2020 , 87, 139.e9-139.e15	5.6	16
112	Initiation of dopaminergic differentiation of Nurr1(-) mesencephalic precursor cells depends on activation of multiple mitogen-activated protein kinase pathways. <i>Stem Cells</i> , 2009 , 27, 2009-21	5.8	15
111	KCNC1-related disorders: new de novo variants expand the phenotypic spectrum. <i>Annals of Clinical and Translational Neurology</i> , 2019 , 6, 1319-1326	5.3	14
110	Usability of eyetracking computer systems and impact on psychological wellbeing in patients with advanced amyotrophic lateral sclerosis. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2018 , 19, 212-219	3.6	14
109	Prion-like properties of disease-relevant proteins in amyotrophic lateral sclerosis. <i>Journal of Neural Transmission</i> , 2018 , 125, 591-613	4.3	13
108	Neurons, Erythrocytes and Beyond -The Diverse Functions of Chorein. <i>NeuroSignals</i> , 2017 , 25, 117-126	1.9	13
107	High content organelle trafficking enables disease state profiling as powerful tool for disease modelling. <i>Scientific Data</i> , 2018 , 5, 180241	8.2	13
106	Effect of high-caloric nutrition on serum neurofilament light chain levels in amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020 , 91, 1007-1009	5.5	12
105	Neurotropic growth factors and glycosaminoglycan based matrices to induce dopaminergic tissue formation. <i>Biomaterials</i> , 2015 , 67, 205-13	15.6	11

104	Early Postnatal but Not Late Adult Neurogenesis Is Impaired in the Pitx3-Mutant Animal Model of Parkinson's Disease. <i>Frontiers in Neuroscience</i> , 2017 , 11, 471	5.1	11
103	Norepinephrine is a negative regulator of the adult periventricular neural stem cell niche. <i>Stem Cells</i> , 2020 , 38, 1188-1201	5.8	10
102	Defective mitochondrial and lysosomal trafficking in chorea-acanthocytosis is independent of Src-kinase signaling. <i>Molecular and Cellular Neurosciences</i> , 2018 , 92, 137-148	4.8	10
101	Proteostasis regulators modulate proteasomal activity and gene expression to attenuate multiple phenotypes in Fabry disease. <i>Biochemical Journal</i> , 2020 , 477, 359-380	3.8	9
100	Axonal Transport, Phase-Separated Compartments, and Neuron Mechanics - A New Approach to Investigate Neurodegenerative Diseases. <i>Frontiers in Cellular Neuroscience</i> , 2018 , 12, 358	6.1	9
99	Impairment in Respiratory Function Contributes to Olfactory Impairment in Amyotrophic Lateral Sclerosis. <i>Frontiers in Neurology</i> , 2018 , 9, 79	4.1	8
98	Serum creatine kinase and creatinine in adult spinal muscular atrophy under nusinersen treatment. <i>Annals of Clinical and Translational Neurology</i> , 2021 , 8, 1049-1063	5.3	8
97	Informal Caregiving in Amyotrophic Lateral Sclerosis (ALS): A High Caregiver Burden and Drastic Consequences on Caregivers' Lives. <i>Brain Sciences</i> , 2021 , 11,	3.4	8
96	Symptomatic pharmacotherapy in ALS: data analysis from a platform-based medication management programme. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020 , 91, 783-785	5.5	7
95	"Silenced" polydendrocytes: a new cell type within the oligodendrocyte progenitor cell population?. <i>Cell and Tissue Research</i> , 2010 , 340, 45-50	4.2	7
94	FUS (fused in sarcoma) is a component of the cellular response to topoisomerase I-induced DNA breakage and transcriptional stress. <i>Life Science Alliance</i> , 2019 , 2,	5.8	7
93	Deep brain stimulation in the globus pallidus compensates response inhibition deficits: evidence from pantothenate kinase-associated neurodegeneration. <i>Brain Structure and Function</i> , 2016 , 221, 2251-4	4	6
92	Hemiplegic migraine with reversible cerebral vasoconstriction caused by ATP1A2 mutations. <i>Journal of Neurology</i> , 2013 , 260, 2172-4	5.5	6
91	Communicating hydrocephalus following eosinophilic meningitis is pathogenic for chronic Viliuisk encephalomyelitis in Northeastern Siberia. <i>PLoS ONE</i> , 2014 , 9, e84670	3.7	6
90	Acanthocyte Sedimentation Rate as a Diagnostic Biomarker for Neuroacanthocytosis Syndromes: Experimental Evidence and Physical Justification. <i>Cells</i> , 2021 , 10,	7.9	6
89	Impairment of mitochondrial oxidative phosphorylation in skin fibroblasts of SALS and FALS patients is rescued by in vitro treatment with ROS scavengers. <i>Experimental Neurology</i> , 2021 , 339, 113620	5.7	6
88	Routine Cerebrospinal Fluid (CSF) Parameters in Patients With Spinal Muscular Atrophy (SMA) Treated With Nusinersen. <i>Frontiers in Neurology</i> , 2019 , 10, 1179	4.1	6
87	Drug-induced endovesiculation of erythrocytes is modulated by the dynamics in the cytoskeleton/membrane interaction. <i>Blood Cells, Molecules, and Diseases</i> , 2017 , 64, 15-22	2.1	5

86	The diagnostic value of midbrain hyperechogenicity in ALS is limited for discriminating key ALS differential diagnoses. <i>BMC Neurology</i> , 2015 , 15, 33	3.1	5
85	Assessment of Gene Variant Amenability for Pharmacological Chaperone Therapy with 1-Deoxygalactonojirimycin in Fabry Disease. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	5
84	Putaminal Dopamine Turnover in de novo Parkinson's Disease Predicts Later Neuropsychiatric Fluctuations but Not Other Major Health Outcomes. <i>Journal of Parkinson's Disease</i> , 2019 , 9, 693-704	5.3	5
83	Generation of the Niemann-Pick type C2 patient-derived iPSC line AKOSi001-A. <i>Stem Cell Research</i> , 2019 , 41, 101606	1.6	5
82	Vocal cord paralysis and rapid progressive motor neuron disease by the I113F mutation in SOD1 gene. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2011 , 12, 382-4		5
81	Safety and Effectiveness of Long-term Intravenous Administration of Edaravone for Treatment of Patients With Amyotrophic Lateral Sclerosis.. <i>JAMA Neurology</i> , 2022 ,	17.2	5
80	One nerve suffices: A clinically guided nerve ultrasound protocol for the differentiation of multifocal motor neuropathy (MMN) and amyotrophic lateral sclerosis (ALS). <i>Journal of Neurology</i> , 2021 , 268, 1495-1507	5.5	5
79	Neurofilament light chain in serum is significantly increased in chorea-acanthocytosis. <i>Parkinsonism and Related Disorders</i> , 2020 , 80, 28-31	3.6	5
78	Oxidative Stress and Alterations in the Antioxidative Defense System in Neuronal Cells Derived from NPC1 Patient-Specific Induced Pluripotent Stem Cells. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	5
77	Cognitive reserve and regional brain volume in amyotrophic lateral sclerosis. <i>Cortex</i> , 2021 , 139, 240-248	3.8	5
76	Stimulation of mGluR1/5 Improves Defective Internalization of AMPA Receptors in NPC1 Mutant Mouse. <i>Cerebral Cortex</i> , 2020 , 30, 1465-1480	5.1	5
75	TDP-43 as structure-based biomarker in amyotrophic lateral sclerosis. <i>Annals of Clinical and Translational Neurology</i> , 2021 , 8, 271-277	5.3	5
74	Treatment satisfaction in 5q-spinal muscular atrophy under nusinersen therapy. <i>Therapeutic Advances in Neurological Disorders</i> , 2021 , 14, 1756286421998902	6.6	5
73	Off-Label Treatment of 4 Amyotrophic Lateral Sclerosis Patients With 4-Aminopyridine. <i>Journal of Clinical Pharmacology</i> , 2019 , 59, 1400-1404	2.9	4
72	Patient-Reported Prevalence of Non-motor Symptoms Is Low in Adult Patients Suffering From 5q Spinal Muscular Atrophy. <i>Frontiers in Neurology</i> , 2019 , 10, 1098	4.1	4
71	New regulator for energy signaling pathway in plants highlights conservation among species. <i>Science Signaling</i> , 2010 , 3, jc5	8.8	4
70	Stage-dependent vulnerability of fetal mesencephalic neuroprogenitors towards dopaminergic neurotoxins. <i>NeuroToxicology</i> , 2008 , 29, 714-21	4.4	4
69	Human Spinal Motor Neurons Are Particularly Vulnerable to Cerebrospinal Fluid of Amyotrophic Lateral Sclerosis Patients. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	4

68	Therapeutic targeting of Lyn kinase to treat chorea-acanthocytosis. <i>Acta Neuropathologica Communications</i> , 2021 , 9, 81	7.3	4
67	Pathomechanisms of ALS8: altered autophagy and defective RNA binding protein (RBP) homeostasis due to the VAPB P56S mutation. <i>Cell Death and Disease</i> , 2021 , 12, 466	9.8	4
66	The Erythrocyte Sedimentation Rate and Its Relation to Cell Shape and Rigidity of Red Blood Cells from Chorea-Acanthocytosis Patients in an Off-Label Treatment with Dasatinib. <i>Biomolecules</i> , 2021 , 11,	5.9	4
65	Factor-Reduced Human Induced Pluripotent Stem Cells Efficiently Differentiate into Neurons Independent of the Number of Reprogramming Factors. <i>Stem Cells International</i> , 2016 , 2016, 4736159	5	4
64	Alteration of GABAergic Input Precedes Neurodegeneration of Cerebellar Purkinje Cells of NPC1-Deficient Mice. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	4
63	Practically applicable nerve ultrasound models for the diagnosis of axonal and demyelinating hereditary motor and sensory neuropathies (HMSN). <i>Journal of Neurology</i> , 2018 , 265, 165-177	5.5	4
62	Combined Dendritic and Axonal Deterioration Are Responsible for Motoneuronopathy in Patient-Derived Neuronal Cell Models of Chorea-Acanthocytosis. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	3
61	Determination of the Pathological Features of NPC1 Variants in a Cellular Complementation Test. <i>International Journal of Molecular Sciences</i> , 2019 , 20,	6.3	3
60	Lack of vascular endothelial growth factor receptor-2/Flk1 signaling does not affect substantia nigra development. <i>Neuroscience Letters</i> , 2013 , 553, 142-7	3.3	3
59	A Case of Beta-propeller Protein-associated Neurodegeneration due to a Heterozygous Deletion of. <i>Tremor and Other Hyperkinetic Movements</i> , 2017 , 7, 465	2	3
58	Combined fluorescence, optical diffraction tomography and Brillouin microscopy		3
57	Catecholaminergic Innervation of Periventricular Neurogenic Regions of the Developing Mouse Brain. <i>Frontiers in Neuroanatomy</i> , 2020 , 14, 558435	3.6	3
56	Targeting Lyn Kinase in Chorea-Acanthocytosis: A Translational Treatment Approach in a Rare Disease. <i>Journal of Personalized Medicine</i> , 2021 , 11,	3.6	3
55	DDX17 is involved in DNA damage repair and modifies FUS toxicity in an RGG-domain dependent manner. <i>Acta Neuropathologica</i> , 2021 , 142, 515-536	14.3	3
54	9-Methyl- β -carboline inhibits monoamine oxidase activity and stimulates the expression of neurotrophic factors by astrocytes. <i>Journal of Neural Transmission</i> , 2020 , 127, 999-1012	4.3	3
53	Pluripotent Stem Cells for Disease Modeling and Drug Discovery in Niemann-Pick Type C1. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	3
52	A Nation-Wide, Multi-Center Study on the Quality of Life of ALS Patients in Germany. <i>Brain Sciences</i> , 2021 , 11,	3.4	3
51	Interleukin-17 and Th17 Lymphocytes Directly Impair Motoneuron Survival of Wildtype and FUS-ALS Mutant Human iPSCs. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	3

50	Palatal Tremor with Progressive Ataxia Secondary to A Dural Arteriovenous Fistula. <i>Movement Disorders Clinical Practice</i> , 2019 , 6, 327-329	2.2	2
49	Reduced LRRK2-positive neurons in the striatum of Parkinson disease patients hypothesize a retrograde disease mechanism?. <i>Basal Ganglia</i> , 2012 , 2, 67-72		2
48	Loss of "insight" into behavioral changes in ALS: Differences across cognitive profiles. <i>Brain and Behavior</i> , 2021 , 12, e2439	3.4	2
47	MDS criteria for the diagnosis of progressive supranuclear palsy overemphasize Richardson syndrome. <i>Annals of Clinical and Translational Neurology</i> , 2020 , 7, 1702-1707	5.3	2
46	Genome Wide Analysis Points towards Subtype-Specific Diseases in Different Genetic Forms of Amyotrophic Lateral Sclerosis. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	2
45	Functional and Molecular Properties of DYT-SGCE Myoclonus-Dystonia Patient-Derived Striatal Medium Spiny Neurons. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	2
44	Cerebellar atrophy on top of motor neuron compromise as indicator of late-onset GM2 gangliosidosis. <i>Journal of Neurology</i> , 2021 , 268, 2259-2262	5.5	2
43	Reconditioning the Neurogenic Niche of Adult Non-human Primates by Antisense Oligonucleotide-Mediated Attenuation of TGFβ Signaling. <i>Neurotherapeutics</i> , 2021 , 18, 1963-1979	6.4	2
42	A serum microRNA sequence reveals fragile X protein pathology in amyotrophic lateral sclerosis. <i>Brain</i> , 2021 , 144, 1214-1229	11.2	2
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40	Co-condensation of proteins with single- and double-stranded DNA.. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022 , 119, e2107871119	11.5	2
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33	High content live profiling reveals concomitant gain and loss of function pathomechanisms in C9ORF72 amyotrophic lateral sclerosis		1

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31	Generation of an iPSC line (AKOSi006-A) from fibroblasts of an NPC1 patient, carrying the homozygous mutation p.I1061T (c.3182T>C) and a control iPSC line (AKOSi007-A) using a non-integrating Sendai virus system. <i>Stem Cell Research</i> , 2020 , 49, 102056	1.6	1
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21	Intersektoralit� 2021 , 93-107		1
20	Concomitant gain and loss of function pathomechanisms in C9ORF72 amyotrophic lateral sclerosis. <i>Life Science Alliance</i> , 2021 , 4,	5.8	1
19	Co-condensation of proteins with single- and double-stranded DNA		1
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