## Angelo Quattrini

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/6332323/publications.pdf

Version: 2024-02-01

29994 39575 9,770 149 54 94 citations h-index g-index papers 152 152 152 10631 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Injection of adult neurospheres induces recovery in a chronic model of multiple sclerosis. Nature, 2003, 422, 688-694.	13.7	1,057
2	Lentiviral haemopoietic stem-cell gene therapy in early-onset metachromatic leukodystrophy: an ad-hoc analysis of a non-randomised, open-label, phase 1/2 trial. Lancet, The, 2016, 388, 476-487.	6.3	393
3	Conditional disruption of $\hat{l}^21$ integrin in Schwann cells impedes interactions with axons. Journal of Cell Biology, 2002, 156, 199-210.	2.3	294
4	Correction of metachromatic leukodystrophy in the mouse model by transplantation of genetically modified hematopoietic stem cells. Journal of Clinical Investigation, 2004, 113, 1118-1129.	3.9	256
5	Ablation of the UPR-Mediator CHOP Restores MotorÂFunction and Reduces Demyelination inÂCharcot-Marie-Tooth 1B Mice. Neuron, 2008, 57, 393-405.	3 <b>.</b> 8	245
6	Axonal degeneration in paraplegin-deficient mice is associated with abnormal mitochondria and impairment of axonal transport. Journal of Clinical Investigation, 2004, 113, 231-242.	3.9	241
7	Polyneuropathy in POEMS syndrome: role of angiogenic factors in the pathogenesis. Brain, 2005, 128, 1911-1920.	3.7	216
8	In vivo gene therapy of metachromatic leukodystrophy by lentiviral vectors: correction of neuropathology and protection against learning impairments in affected mice. Nature Medicine, 2001, 7, 310-316.	15.2	198
9	Gene therapy of metachromatic leukodystrophy reverses neurological damage and deficits in mice. Journal of Clinical Investigation, 2006, $116$ , $3070-3082$ .	3.9	197
10	Identification of Hematopoietic Stem Cell–Specific miRNAs Enables Gene Therapy of Globoid Cell Leukodystrophy. Science Translational Medicine, 2010, 2, 58ra84.	5.8	180
11	Loss of glial fibrillary acidic protein (GFAP) impairs Schwann cell proliferation and delays nerve regeneration after damage. Journal of Cell Science, 2006, 119, 3981-3993.	1.2	174
12	Evidence of peripheral axonal neuropathy in primary restless legs syndrome. Movement Disorders, 1995, 10, 2-9.	2.2	170
13	Brain conditioning is instrumental for successful microglia reconstitution following hematopoietic stem cell transplantation. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 15018-15023.	3.3	168
14	Disruption of Mtmr2 produces CMT4B1-like neuropathy with myelin outfolding and impaired spermatogenesis. Journal of Cell Biology, 2004, 167, 711-721.	2.3	167
15	$\hat{l}^21$ integrin activates Rac1 in Schwann cells to generate radial lamellae during axonal sorting and myelination. Journal of Cell Biology, 2007, 177, 1063-1075.	2.3	163
16	Gene therapy augments the efficacy of hematopoietic cell transplantation and fully corrects mucopolysaccharidosis type I phenotype in the mouse model. Blood, 2010, 116, 5130-5139.	0.6	159
17	Alpha-lipoic acid prevents mitochondrial damage and neurotoxicity in experimental chemotherapy neuropathy. Experimental Neurology, 2008, 214, 276-284.	2.0	158
18	PO Glycoprotein Overexpression Causes Congenital Hypomyelination of Peripheral Nerves. Journal of Cell Biology, 2000, 148, 1021-1034.	2.3	145

#	Article	lF	Citations
19	Different Intracellular Pathomechanisms Produce Diverse Myelin Protein Zero Neuropathies in Transgenic Mice. Journal of Neuroscience, 2006, 26, 2358-2368.	1.7	144
20	Axonal degeneration in paraplegin-deficient mice is associated with abnormal mitochondria and impairment of axonal transport. Journal of Clinical Investigation, 2004, 113, 231-242.	3.9	144
21	TACE (ADAM17) inhibits Schwann cell myelination. Nature Neuroscience, 2011, 14, 857-865.	7.1	136
22	Mitochondrial biogenesis and fission in axons in cell culture and animal models of diabetic neuropathy. Acta Neuropathologica, 2010, 120, 477-489.	3.9	125
23	Role of integrins in the peripheral nervous system. Progress in Neurobiology, 2001, 64, 35-49.	2.8	123
24	Heterogeneity of autoantibodies in stiff-man syndrome. Annals of Neurology, 1993, 34, 57-64.	2.8	121
25	Correction of metachromatic leukodystrophy in the mouse model by transplantation of genetically modified hematopoietic stem cells. Journal of Clinical Investigation, 2004, 113, 1118-1129.	3.9	117
26	Lab-on-Chip for Exosomes and Microvesicles Detection and Characterization. Sensors, 2018, 18, 3175.	2.1	107
27	iPSC-derived neural precursors exert a neuroprotective role in immune-mediated demyelination via the secretion of LIF. Nature Communications, 2013, 4, 2597.	5.8	104
28	Relief of inflammatory pain in rats by local use of the selective P2X7 ATP receptor inhibitor, oxidized ATP. Arthritis and Rheumatism, 2002, 46, 3378-3385.	6.7	101
29	Dlg1, Sec8, and Mtmr2 Regulate Membrane Homeostasis in Schwann Cell Myelination. Journal of Neuroscience, 2009, 29, 8858-8870.	1.7	101
30	Expression of Laminin Receptors in Schwann Cell Differentiation: Evidence for Distinct Roles. Journal of Neuroscience, 2003, 23, 5520-5530.	1.7	100
31	Haploinsufficiency of <i> AFG3L2 </i> ), the Gene Responsible for Spinocerebellar Ataxia Type 28, Causes Mitochondria-Mediated Purkinje Cell Dark Degeneration. Journal of Neuroscience, 2009, 29, 9244-9254.	1.7	99
32	Loss of Mtmr2 Phosphatase in Schwann Cells But Not in Motor Neurons Causes Charcot-Marie-Tooth Type 4B1 Neuropathy with Myelin Outfoldings. Journal of Neuroscience, 2005, 25, 8567-8577.	1.7	95
33	The Mitochondrial Protease AFG3L2 Is Essential for Axonal Development. Journal of Neuroscience, 2008, 28, 2827-2836.	1.7	92
34	Hypogonadotropic hypogonadism and peripheral neuropathy inEbf2-null mice. Development (Cambridge), 2003, 130, 401-410.	1.2	89
35	Genetic Interaction between MTMR2 and FIG4 Phospholipid Phosphatases Involved in Charcot-Marie-Tooth Neuropathies. PLoS Genetics, 2011, 7, e1002319.	1.5	87
36	CRYOGLOBULINAEMIC NEUROPATHY. Brain, 1988, 111, 541-552.	3.7	85

#	Article	IF	CITATIONS
37	$\hat{l}^2$ 4 integrin and other Schwann cell markers in axonal neuropathy. , 1996, 17, 294-306.		82
38	Antinociceptive effect of a new P2Z/P2X7 antagonist, oxidized ATP, in arthritic rats. Neuroscience Letters, 2002, 327, 87-90.	1.0	81
39	Â6Â4 Integrin and Dystroglycan Cooperate to Stabilize the Myelin Sheath. Journal of Neuroscience, 2008, 28, 6714-6719.	1.7	78
40	Sox2 expression in Schwann cells inhibits myelination in vivo and induces influx of macrophages to the nerve. Development (Cambridge), 2017, 144, 3114-3125.	1.2	75
41	Patterns of reactivity of human anti-GM1 antibodies with spinal cord and motor neurons. Annals of Neurology, 1992, 32, 487-493.	2.8	71
42	Myotubularin-related 2 protein phosphatase and neurofilament light chain protein, both mutated in CMT neuropathies, interact in peripheral nerve. Human Molecular Genetics, 2003, 12, 1713-1723.	1.4	67
43	Subventricular zone neural progenitors protect striatal neurons from glutamatergic excitotoxicity. Brain, 2012, 135, 3320-3335.	3.7	67
44	Purkinje neuron Ca2+ influx reduction rescues ataxia in SCA28 model. Journal of Clinical Investigation, 2015, 125, 263-274.	3.9	67
45	Recent advances in amyotrophic lateral sclerosis. Journal of Neurology, 2016, 263, 1241-1254.	1.8	67
46	Corneal confocal microscopy reveals trigeminal small sensory fiber neuropathy in amyotrophic lateral sclerosis. Frontiers in Aging Neuroscience, 2014, 6, 278.	1.7	66
47	Prostaglandin D2 synthase/GPR44: a signaling axis in PNS myelination. Nature Neuroscience, 2014, 17, 1682-1692.	7.1	66
48	Immune response in peripheral axons delays disease progression in SOD1G93A mice. Journal of Neuroinflammation, 2016, 13, 261.	3.1	63
49	Charcot–Marie–Tooth type 4B demyelinating neuropathy: deciphering the role of MTMR phosphatases. Expert Reviews in Molecular Medicine, 2007, 9, 1-16.	1.6	62
50	Antibodies to sulfatide and to chondroitin sulfate C in patients with chronic sensory neuropathy. Journal of Neuroimmunology, 1993, 43, 79-85.	1.1	61
51	Vimentin regulates peripheral nerve myelination. Development (Cambridge), 2012, 139, 1359-1367.	1.2	58
52	The gp $120\mathrm{glycoprotein}$ of human immunodeficiency virus type $1\mathrm{binds}$ to sensory ganglion neurons. Annals of Neurology, $1993, 34, 855-863.$	2.8	57
53	A novel POglycoprotein transgene activates expression oflacZ in myelin-forming Schwann cells. European Journal of Neuroscience, 1999, 11, 1577-1586.	1.2	57
54	Effect of chronic treatment with recombinant interleukin-2 on the central nervous system of adult and old mice. Brain Research, 1992, 591, 248-252.	1.1	56

#	Article	IF	Citations
55	Genetic interaction between the m -AAA protease isoenzymes reveals novel roles in cerebellar degeneration. Human Molecular Genetics, 2009, 18, 2001-2013.	1.4	55
56	Non-redundant function of dystroglycan and $\hat{l}^21$ integrins in radial sorting of axons. Development (Cambridge), 2011, 138, 4025-4037.	1.2	55
57	Structural and functional brain signatures of C9orf72 in motor neuron disease. Neurobiology of Aging, 2017, 57, 206-219.	1.5	54
58	Epitope-Tagged POGlycoprotein Causes Charcot-Marie-Tooth–Like Neuropathy in Transgenic Mice. Journal of Cell Biology, 2000, 151, 1035-1046.	2.3	53
59	Anti-sulfatide antibodies in neurological disease: binding to rat dorsal root ganglia neurons. Journal of the Neurological Sciences, 1992, 112, 152-159.	0.3	52
60	DDIT4/REDD1/RTP801 Is a Novel Negative Regulator of Schwann Cell Myelination. Journal of Neuroscience, 2013, 33, 15295-15305.	1.7	51
61	$\hat{l}$ ± $6\hat{l}^21$ and $\hat{l}$ ± $7\hat{l}^21$ Integrins Are Required in Schwann Cells to Sort Axons. Journal of Neuroscience, 2013, 33, 17995-18007.	1.7	49
62	Increased expression of Myosin binding protein H in the skeletal muscle of amyotrophic lateral sclerosis patients. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2014, 1842, 99-106.	1.8	49
63	Analyzing Histopathological Features of Rare Charcot-Marie-Tooth Neuropathies to Unravel Their Pathogenesis. Archives of Neurology, 2010, 67, 1498-505.	4.9	48
64	Efficacy of silver coated surgical sutures on bacterial contamination, cellular response and wound healing. Materials Science and Engineering C, 2016, 69, 884-893.	3.8	48
65	Intramuscular viral delivery of paraplegin rescues peripheral axonopathy in a model of hereditary spastic paraplegia. Journal of Clinical Investigation, 2005, $116$ , 202-208.	3.9	48
66	Retromer stabilization results in neuroprotection in a model of Amyotrophic Lateral Sclerosis. Nature Communications, 2020, 11, 3848.	5.8	44
67	Motor nerve biopsy studies in motor neuropathy and motor neuron disease. Muscle and Nerve, 1997, 20, 15-21.	1.0	42
68	Neural Stem Cells of the Subventricular Zone Contribute to Neuroprotection of the Corpus Callosum after Cuprizone-Induced Demyelination. Journal of Neuroscience, 2019, 39, 5481-5492.	1.7	42
69	Counteracting roles of MHCI and CD8+ T cells in the peripheral and central nervous system of ALS SOD1G93A mice. Molecular Neurodegeneration, 2018, 13, 42.	4.4	40
70	$\hat{l}\pm6\hat{l}^24$ and $\hat{l}\pm6\hat{l}^21$ Integrins in Astrocytomas and Other CNS Tumors. Journal of Neuropathology and Experimental Neurology, 1996, 55, 456-465.	0.9	39
71	Peripheral nerve morphogenesis induced by scaffold micropatterning. Biomaterials, 2014, 35, 4035-4045.	5.7	39
72	Motor nerve biopsy: Clinical usefulness and histopathological criteria. Annals of Neurology, 2011, 69, 197-201.	2.8	38

#	Article	IF	CITATIONS
73	<i>TBK1</i> mutations in Italian patients with amyotrophic lateral sclerosis: genetic and functional characterisation. Journal of Neurology, Neurosurgery and Psychiatry, 2017, 88, 869-875.	0.9	38
74	A minimal human MBP Promoter-lacZ transgene is appropriately regulated in developing brain and after optic enucleation, but not in shiverer mutant mice. Journal of Neurobiology, 1998, 34, 10-26.	3.7	37
75	Docetaxel neuropathy: a distal axonopathy. Acta Neuropathologica, 1999, 98, 651-653.	3.9	35
76	Jab1 regulates Schwann cell proliferation and axonal sorting through p27. Journal of Experimental Medicine, 2014, 211, 29-43.	4.2	35
77	Cxcl10 enhances blood cells migration in the sub-ventricular zone of mice affected by experimental autoimmune encephalomyelitis. Molecular and Cellular Neurosciences, 2010, 43, 268-280.	1.0	34
78	MR Imaging of Brachial Plexus and Limb-Girdle Muscles in Patients with Amyotrophic Lateral Sclerosis. Radiology, 2016, 279, 553-561.	3.6	32
79	Diet, Microbiota and Brain Health: Unraveling the Network Intersecting Metabolism and Neurodegeneration. International Journal of Molecular Sciences, 2020, 21, 7471.	1.8	32
80	Human IgM anti-GM1 autoantibodies modulate intracellular calcium homeostasis in neuroblastoma cells. Journal of Neuroimmunology, 2001, 114, 213-219.	1.1	31
81	Impaired turnover of hyperfused mitochondria in severe axonal neuropathy due to a novel DRP1 mutation. Human Molecular Genetics, 2020, 29, 177-188.	1.4	30
82	Two factor-based reprogramming of rodent and human fibroblasts into Schwann cells. Nature Communications, 2017, 8, 14088.	5.8	28
83	The Peripheral Nervous System in Amyotrophic Lateral Sclerosis: Opportunities for Translational Research. Frontiers in Neuroscience, 2019, 13, 601.	1.4	28
84	Impaired flickering of the permeability transition pore causes SPG7 spastic paraplegia. EBioMedicine, 2020, 61, 103050.	2.7	28
85	Neutrophils predominate the immune signature of cerebral thrombi in COVID-19 stroke patients. Acta Neuropathologica Communications, 2022, 10, 14.	2.4	27
86	Vocal cord paralysis in Charcot–Marie–Tooth type 4b1 disease associated with a novel mutation in the myotubularin-related protein 2 gene: A case report and review of the literature. Neuromuscular Disorders, 2017, 27, 487-491.	0.3	26
87	Serum phosphorylated neurofilament heavy-chain levels reflect phenotypic heterogeneity and are an independent predictor of survival in motor neuron disease. Journal of Neurology, 2020, 267, 2272-2280.	1.8	26
88	Integrated evaluation of a panel of neurochemical biomarkers to optimize diagnosis and prognosis in amyotrophic lateral sclerosis. European Journal of Neurology, 2022, 29, 1930-1939.	1.7	25
89	Axonal neuropathy with monoclonal IgG kappa that binds to a neurofilament protein. Annals of Neurology, 1990, 28, 361-364.	2.8	24
90	Unraveling gene expression profiles in peripheral motor nerve from amyotrophic lateral sclerosis patients: insights into pathogenesis. Scientific Reports, 2016, 6, 39297.	1.6	24

#	Article	IF	CITATIONS
91	Schwann cell overexpression of the GPR7 receptor in inflammatory and painful neuropathies. Molecular and Cellular Neurosciences, 2005, 28, 55-63.	1.0	23
92	A novel composite type I collagen scaffold with micropatterned porosity regulates the entrance of phagocytes in a severe model of spinal cord injury., 2017, 105, 1040-1053.		23
93	Phosphorylated TDP-43 aggregates in peripheral motor nerves of patients with amyotrophic lateral sclerosis. Brain, 2022, 145, 276-284.	3.7	22
94	An update on the diagnosis and management of the polyneuropathy of POEMS syndrome. Journal of Neurology, 2019, 266, 258-267.	1.8	21
95	A longitudinal DTI and histological study of the spinal cord reveals early pathological alterations in G93A-SOD1 mouse model of amyotrophic lateral sclerosis. Experimental Neurology, 2017, 293, 43-52.	2.0	19
96	Selective loss of microvesicles is a major issue of the differential centrifugation isolation protocols. Scientific Reports, 2021, 11, 3589.	1.6	19
97	Defining Peripheral Nervous System Dysfunction in the SOD-1 <sup>G93A</sup> Transgenic Rat Model of Amyotrophic Lateral Sclerosis. Journal of Neuropathology and Experimental Neurology, 2014, 73, 658-670.	0.9	18
98	Early detection of skin and muscular involvement in lafora disease. Journal of Neurology, 1991, 238, 217-220.	1.8	17
99	Churg Strauss syndrome presenting as acute neuropathy resembling Guillain Barré syndrome. Journal of Neurology, 2008, 255, 1843-1844.	1.8	17
100	Chronic motor axonal neuropathy. Journal of the Peripheral Nervous System, 2011, 16, 341-346.	1.4	17
101	Synthesis and Preliminary Evaluation in Tumor Bearing Mice of New <sup>18</sup> F-Labeled Arylsulfone Matrix Metalloproteinase Inhibitors as Tracers for Positron Emission Tomography. Journal of Medicinal Chemistry, 2013, 56, 2676-2689.	2.9	17
102	Current application of neurofilaments in amyotrophic lateral sclerosis and future perspectives. Neural Regeneration Research, 2021, 16, 1985.	1.6	17
103	lgG monoclonal proteins from patients with axonal peripheral neuropathies bind to different epitopes of the 68 kDa neurofilament protein. Journal of Neuroimmunology, 1992, 36, 97-104.	1.1	16
104	Acute presentation of Tangier polyneuropathy: a clinical and morphological study. Acta Neuropathologica, 1993, 86, 90-94.	3.9	16
105	Urokinase Plasminogen Receptor and the Fibrinolytic Complex Play a Role in Nerve Repair after Nerve Crush in Mice, and in Human Neuropathies. PLoS ONE, 2012, 7, e32059.	1.1	16
106	Axonal neuropathy in a patient with monoclonal IgM kappa reactive with Schmidt-Lantermann incisures. Journal of Neuroimmunology, 1991, 33, 73-79.	1.1	15
107	Distinct Protein Expression Networks are Activated in Microglia Cells after Stimulation with IFN- $\hat{l}^3$ and IL-4. Cells, 2019, 8, 580.	1.8	15
108	Loss of function <scp>MPZ</scp> mutation causes milder <scp>CMT1B</scp> neuropathy. Journal of the Peripheral Nervous System, 2021, 26, 177-183.	1.4	15

#	Article	IF	CITATIONS
109	Autoimmunity in the Peripheral Nervous System. Critical Reviews in Neurobiology, 2003, 15, 1-39.	3.3	15
110	Clinical features and outcomes of the flail arm and flail leg and pure lower motor neuron MND variants: a multicentre Italian study. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 1001-1003.	0.9	14
111	Thermosensitive chitosan-based hydrogels supporting motor neuron-like NSC-34 cell differentiation. Biomaterials Science, 2021, 9, 7492-7503.	2.6	14
112	Effect of hypothyroidism on rat peripheral nervous system. NeuroReport, 1993, 4, 499-502.	0.6	13
113	Foot pad skin biopsy in mouse models of hereditary neuropathy. Glia, 2010, 58, 2005-2016.	2.5	13
114	Monoclonal Antibodies Conjugated with Superparamagnetic Iron Oxide Particles Allow Magnetic Resonance Imaging Detection of Lymphocytes in the Mouse Brain. Molecular Imaging, 2012, 11, 7290.2011.00032.	0.7	13
115	Prostaglandin D2 synthase modulates macrophage activity and accumulation in injured peripheral nerves. Glia, 2020, 68, 95-110.	2.5	13
116	Functioning and quality of life in patients with neuropathy associated with anti-MAG antibodies. Journal of Neurology, 2018, 265, 2927-2933.	1.8	12
117	A nonsense mutation in myelin protein zero causes congenital hypomyelination neuropathy through altered PO membrane targeting and gain of abnormal function. Human Molecular Genetics, 2019, 28, 124-132.	1.4	12
118	JAB1 deletion in oligodendrocytes causes senescence-induced inflammation and neurodegeneration in mice. Journal of Clinical Investigation, 2022, 132, .	3.9	12
119	Balance exercise in patients with chronic sensory ataxic neuropathy: a pilot study. Journal of the Peripheral Nervous System, 2014, 19, 145-151.	1.4	11
120	Development of Injectable Thermosensitive Chitosan-Based Hydrogels for Cell Encapsulation. Applied Sciences (Switzerland), 2020, 10, 6550.	1.3	11
121	Burden of Rare Variants in ALS and Axonal Hereditary Neuropathy Genes Influence Survival in ALS: Insights from a Next Generation Sequencing Study of an Italian ALS Cohort. International Journal of Molecular Sciences, 2020, 21, 3346.	1.8	11
122	X-ray phase contrast tomography for the investigation of amyotrophic lateral sclerosis. Journal of Synchrotron Radiation, 2020, 27, 1042-1048.	1.0	11
123	In vivo modulation of myelin gene expression by human recombinant IL-2. Molecular Brain Research, 1992, 12, 331-334.	2.5	10
124	A fatal case of Churg–Strauss syndrome presenting with acute polyneuropathy mimicking Guillain–Barré syndrome. Neurological Sciences, 2011, 32, 937-940.	0.9	10
125	Neonatal combination therapy improves some of the clinical manifestations in the Mucopolysaccharidosis type I murine model. Molecular Genetics and Metabolism, 2020, 130, 197-208.	0.5	10
126	Intramuscular viral delivery of paraplegin rescues peripheral axonopathy in a model of hereditary spastic paraplegia. Journal of Clinical Investigation, 2014, 124, 871-871.	3.9	10

#	Article	IF	CITATIONS
127	Proteomic expression profile of injured rat peripheral nerves revealed biological networks and processes associated with nerve regeneration. Journal of Cellular Physiology, 2018, 233, 6207-6223.	2.0	9
128	NEK1 Variants in a Cohort of Italian Patients With Amyotrophic Lateral Sclerosis. Frontiers in Neuroscience, 2022, 16, 833051.	1.4	9
129	The brachial plexus branches to the pectoral muscles in adult rats: morphological aspects and morphometric normative data. Frontiers in Neuroanatomy, 2012, 6, 41.	0.9	8
130	Neurofilament light chain as a biological marker for amyotrophic lateral sclerosis: a meta-analysis study. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2022, 23, 446-457.	1.1	8
131	Laminin receptor ?6?4 integrin is highly expressed in ENU-induced glioma in rat. Glia, 1999, 26, 55-63.	2.5	7
132	From pathogenesis to personalized treatments of neuropathies in hematological malignancies. Journal of the Peripheral Nervous System, 2020, 25, 212-221.	1.4	7
133	Corneal and Epidermal Nerve Quantification in Chemotherapy Induced Peripheral Neuropathy. Frontiers in Medicine, 2022, 9, 832344.	1.2	7
134	Morphological and Functional Evaluation of Peripheral Nerve Regeneration in the Rat Using an Expanded Polytetrafluoroethylene (PTFE) Microprosthesis. Journal of Investigative Surgery, 1991, 4, 437-443.	0.6	6
135	Neurovascular signals in amyotrophic lateral sclerosis. Current Opinion in Biotechnology, 2022, 74, 75-83.	3.3	6
136	Peripheral Nerve Dysmyelination Due to POGlycoprotein Overexpression Is Dose-Dependent. Annals of the New York Academy of Sciences, 1999, 883, 294-301.	1.8	5
137	Limitations in daily activities and general perception of quality of life: Long term followâ€up in patients with antiâ€myelinâ€glycoprotein antibody polyneuropathy. Journal of the Peripheral Nervous System, 2019, 24, 276-282.	1.4	5
138	Diffuse intraneural leiomyoma in a case of sensorimotor neuropathy. Acta Neuropathologica, 2009, 117, 595-597.	3.9	3
139	Ablation of neuronal ADAM17 impairs oligodendrocyte differentiation and myelination. Glia, 2020, 68, 1148-1164.	2.5	2
140	Sox2 expression in Schwann cells inhibits myelination in vivo and induces influx of macrophages to the nerve. Journal of Cell Science, 2017, 130, e1.2-e1.2.	1.2	2
141	Stiff-Man Syndrome. , 2014, , 1465-1477.		2
142	ADAM17 Regulates p75 <sup>NTR</sup> -Mediated Fibrinolysis and Nerve Remyelination. Journal of Neuroscience, 2022, 42, 2433-2447.	1.7	2
143	Concurrence of NMOSD and ALS in a patient with hexanucleotide repeat expansions of C9orf72. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2019, 20, 449-452.	1.1	1
144	Normal structure and pathological features in peripheral neuropathies. Journal of the Peripheral Nervous System, 2021, 26, S11-S20.	1.4	1

## Angelo Quattrini

#	Article	IF	CITATIONS
145	Primary Lateral Sclerosis Presenting With Focal Onset Spreading Through Contiguous Neuroanatomic Regions. Neurology, 2022, , 10.1212/WNL.00000000000011.	1.5	1
146	Neuromuscular weakness., 0,, 317-331.		0
147	Nerve pathology in animal models of neuropathies. Journal of the Peripheral Nervous System, 2021, 26 Suppl 2, S61-S68.	1.4	O
148	Vimentin regulates peripheral nerve myelination. Journal of Cell Science, 2012, 125, e1-e1.	1.2	0
149	Jab1 regulates Schwann cell proliferation and axonal sorting through p27. Journal of Cell Biology, 2013, 203, 2036OIA155.	2.3	0