Massimiliano Mirabella

List of Publications by Year in descending order

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210 papers 7,357 citations

45 h-index 74163 75 g-index

215 all docs

215 docs citations

215 times ranked

9680 citing authors

#	Article	IF	CITATIONS
1	Risk of multiple sclerosis relapses when switching from fingolimod to cell-depleting agents: the role of washout duration. Journal of Neurology, 2022, 269, 1463-1469.	3.6	4
2	Immunogenicity and safety of mRNA COVID-19 vaccines in people with multiple sclerosis treated with different disease-modifying therapies. Neurotherapeutics, 2022, 19, 325-333.	4.4	32
3	Cognitive Reserve in Early Manifest Huntington Disease Patients: Leisure Time Is Associated with Lower Cognitive and Functional Impairment. Journal of Personalized Medicine, 2022, 12, 36.	2.5	7
4	Hsa-miR223-3p circulating level is upregulated in Friedreich's ataxia and inversely associated with HCLS1 associated protein X-1, HAX-1. Human Molecular Genetics, 2022, , .	2.9	1
5	A TLR/CD44 axis regulates T cell trafficking in experimental and human multiple sclerosis. IScience, 2022, 25, 103763.	4.1	12
6	Influence of Previous Disease-Modifying Drug Exposure on T-Lymphocyte Dynamic in Patients With Multiple Sclerosis Treated With Ocrelizumab. Neurology: Neuroimmunology and NeuroInflammation, 2022, 9, .	6.0	9
7	Update on Multiple Sclerosis Molecular Biomarkers to Monitor Treatment Effects. Journal of Personalized Medicine, 2022, 12, 549.	2.5	4
8	The Expanding Role of the Infectious Disease Expert in the Context of the MS Centre. Journal of Personalized Medicine, 2022, 12, 591.	2.5	0
9	Treatment Challenges in Multiple Sclerosis – A Continued Role for Glatiramer Acetate?. Frontiers in Neurology, 2022, 13, 844873.	2.4	4
10	Shift of multiple sclerosis onset towards older age. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, 1137-1139.	1.9	12
11	Heterozygous <i>KIF1A</i> variants underlie a wide spectrum of neurodevelopmental and neurodegenerative disorders. Journal of Medical Genetics, 2021, 58, 475-483.	3.2	21
12	A voxel-based lesion symptom mapping analysis of chronic pain in multiple sclerosis. Neurological Sciences, 2021, 42, 1941-1947.	1.9	3
13	Predictors of lymphocyte count recovery after dimethyl fumarate-induced lymphopenia in people with multiple sclerosis. Journal of Neurology, 2021, 268, 2238-2245.	3.6	15
14	Real world experience with teriflunomide in multiple sclerosis: the TER-Italy study. Journal of Neurology, 2021, 268, 2922-2932.	3.6	18
15	Efficacy and Safety of Bimagrumab in Sporadic Inclusion Body Myositis. Neurology, 2021, 96, e1595-e1607.	1.1	25
16	Exit Strategies in Natalizumab-Treated RRMS at High Risk of Progressive Multifocal Leukoencephalopathy: a Multicentre Comparison Study. Neurotherapeutics, 2021, 18, 1166-1174.	4.4	24
17	Defining the disease course of TNFα blockers-associated Multiple Sclerosis. Journal of Neuroimmunology, 2021, 353, 577525.	2.3	4
18	Long-term Follow-up and Muscle Imaging Findings in Brachio-Cervical Inflammatory Myopathy. Neurology: Neuroimmunology and NeuroInflammation, 2021, 8, .	6.0	4

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19	Anti-cN1A Antibodies Are Associated with More Severe Dysphagia in Sporadic Inclusion Body Myositis. Cells, 2021, 10, 1146.	4.1	23
20	MRI activity and extended interval of Natalizumab dosing regimen: a multicentre Italian study. Journal of the Neurological Sciences, 2021, 424, 117385.	0.6	9
21	The neurobiological underpinning of the social cognition impairments in patients with spinocerebellar ataxia type 2. Cortex, 2021, 138, 101-112.	2.4	22
22	Cognitive and behavioral associated changes in manifest Huntington disease: A retrospective crossâ€sectional study. Brain and Behavior, 2021, 11, e02151.	2.2	12
23	Lower urinary tract disorders in multiple sclerosis patients: prevalence, clinical features, and response to treatments. Neurourology and Urodynamics, 2021, 40, 1500-1508.	1.5	8
24	The Disease-Modifying Therapies of Relapsing-Remitting Multiple Sclerosis and Liver Injury: A Narrative Review. CNS Drugs, 2021, 35, 861-880.	5.9	29
25	DMTs and Covidâ€19 severity in MS: a pooled analysis from Italy and France. Annals of Clinical and Translational Neurology, 2021, 8, 1738-1744.	3.7	86
26	Intestinal Permeability and Circulating CD161+CCR6+CD8+T Cells in Patients With Relapsing–Remitting Multiple Sclerosis Treated With Dimethylfumarate. Frontiers in Neurology, 2021, 12, 683398.	2.4	5
27	Disease Reactivation after Fingolimod Discontinuation in Pregnant Multiple Sclerosis Patients. Neurotherapeutics, 2021, 18, 2598-2607.	4.4	12
28	F19â \in Cognitive reserve: the leisure time concurs to the cognition performance and to the independence of early huntington disease patients., 2021,,.		0
29	Comment on: Clinico-radiologic features and therapeutic strategies in tumefactive demyelination: a retrospective analysis of 50 consecutive cases. Therapeutic Advances in Neurological Disorders, 2021, 14, 175628642110500.	3.5	2
30	CSF CXCL13 and chitinase 3-like-1 concentrations predict disease course in relapsing multiple sclerosis. Journal of the Neurological Sciences, 2021, 429, 118114.	0.6	0
31	Extending the Interval of Natalizumab Dosing: Is Efficacy Preserved?. Neurotherapeutics, 2020, 17, 200-207.	4.4	39
32	Exit strategies for "needle fatigue―in multiple sclerosis: a propensity score-matched comparison study. Journal of Neurology, 2020, 267, 694-702.	3.6	6
33	Alemtuzumab-induced lung injury in multiple sclerosis: Learning from adversity in three patients. Multiple Sclerosis and Related Disorders, 2020, 37, 101450.	2.0	8
34	Natalizumab is associated with early improvement of working ability in relapsing-remitting multiple sclerosis patients: WANT observational study results. Neurological Sciences, 2020, 42, 2837-2845.	1.9	11
35	Dimethyl fumarate vs Teriflunomide: an Italian time-to-event data analysis. Journal of Neurology, 2020, 267, 3008-3020.	3.6	19
36	Induction Versus Escalation in Multiple Sclerosis: A 10-Year Real World Study. Neurotherapeutics, 2020, 17, 994-1004.	4.4	34

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37	Cost-Effectiveness Analysis of Cannabinoid Oromucosal Spray Use for the Management of Spasticity in Subjects with Multiple Sclerosis. Clinical Drug Investigation, 2020, 40, 319-326.	2.2	5
38	Dominus effect: challenging complications of alemtuzumab-related thyroid autoimmunity. Journal of Endocrinological Investigation, 2020, 43, 1159-1161.	3.3	2
39	Is serological response to SARS-CoV-2 preserved in MS patients on ocrelizumab treatment? A case report. Multiple Sclerosis and Related Disorders, 2020, 44, 102323.	2.0	34
40	An Italian Neurology Outpatient Clinic Facing SARS-CoV-2 Pandemic: Data From 2,167 Patients. Frontiers in Neurology, 2020, 11, 564.	2.4	30
41	Muscle involvement in myasthenia gravis: Expanding the clinical spectrum of Myasthenia-Myositis association from a large cohort of patients. Autoimmunity Reviews, 2020, 19, 102498.	5.8	32
42	Frataxin deficiency in Friedreich's ataxia is associated with reduced levels of HAX-1, a regulator of cardiomyocyte death and survival. Human Molecular Genetics, 2020, 29, 471-482.	2.9	8
43	A pilot study of lncRNAs expression profile in serum of progressive multiple sclerosis patients. European Review for Medical and Pharmacological Sciences, 2020, 24, 3267-3273.	0.7	11
44	Estimating the impact of COVID-19 pandemic on services provided by Italian Neuromuscular Centers: an Italian Association of Myology survey of the acute phase. Acta Myologica, 2020, 39, 57-66.	1.5	24
45	A method to compare prospective and historical cohorts to evaluate drug effects. Application to the analysis of early treatment effectiveness of intramuscular interferon- \hat{l}^21a in multiple sclerosis patients. Multiple Sclerosis and Related Disorders, 2020, 40, 101952.	2.0	O
46	Hereditary inclusion-body myopathies. , 2020, , 479-489.		0
47	DNA damage signatures in peripheral blood cells as biomarkers in prodromal huntington disease. Annals of Neurology, 2019, 85, 296-301.	5. 3	28
48	Safety and efficacy of intravenous bimagrumab in inclusion body myositis (RESILIENT): a randomised, double-blind, placebo-controlled phase 2b trial. Lancet Neurology, The, 2019, 18, 834-844.	10.2	91
49	The predictive value of CSF multiple assay in multiple sclerosis: A single center experience. Multiple Sclerosis and Related Disorders, 2019, 35, 176-181.	2.0	13
50	Drug Holiday of Interferon Beta 1b in Multiple Sclerosis: A Pilot, Randomized, Single Blind Study of Non-inferiority. Frontiers in Neurology, 2019, 10, 695.	2.4	5
51	A unique case of multiphasic ADEM or what else?. Multiple Sclerosis and Related Disorders, 2019, 35, 73-75.	2.0	1
52	The influence of physiotherapy intervention on patients with multiple sclerosis–related spasticity treated with nabiximols (THC:CBD oromucosal spray). PLoS ONE, 2019, 14, e0219670.	2.5	7
53	P.20Expanding the myasthenia-myositis association spectrum: clinical, morphological and immunological data form a large case series. Neuromuscular Disorders, 2019, 29, S47.	0.6	O
54	The Contribution of Gut Barrier Changes to Multiple Sclerosis Pathophysiology. Frontiers in Immunology, 2019, 10, 1916.	4.8	39

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55	Cardiovascular autonomic individual profile of relapsing-remitting multiple sclerosis patients and risk of extending cardiac monitoring after first dose fingolimod. Journal of the Neurological Sciences, 2019, 405, 116423.	0.6	6
56	A phase 3 randomized study evaluating sialic acid extended-release for GNE myopathy. Neurology, 2019, 92, e2109-e2117.	1.1	40
57	Genome-Wide Multiple Sclerosis Association Data and Coagulation. Frontiers in Neurology, 2019, 10, 95.	2.4	7
58	Different regimen of natalizumab treatment in multiple sclerosis patients: A real world study in Italy. Journal of the Neurological Sciences, 2019, 405, 338-339.	0.6	1
59	The Italian multiple sclerosis register. Neurological Sciences, 2019, 40, 155-165.	1.9	59
60	Novel homozygous GBA2 mutation in a patient with complicated spastic paraplegia. Clinical Neurology and Neurosurgery, 2018, 168, 60-63.	1.4	9
61	The Prevalence of Multiple Sclerosis in the Metropolitan Area of Rome: A Capture-Recapture Analysis. Neuroepidemiology, 2018, 50, 105-110.	2.3	4
62	Sporadic late-onset nemaline myopathy: clinical, pathology and imaging findings in a single center cohort. Journal of Neurology, 2018, 265, 542-551.	3 . 6	36
63	Siponimod versus placebo in secondary progressive multiple sclerosis (EXPAND): a double-blind, randomised, phase 3 study. Lancet, The, 2018, 391, 1263-1273.	13.7	684
64	Low reliability of anti-KIR4.183–120 peptide auto-antibodies in multiple sclerosis patients. Multiple Sclerosis Journal, 2018, 24, 910-918.	3.0	5
65	Systematic assessment and characterization of chronic pain in multiple sclerosis patients. Neurological Sciences, 2018, 39, 445-453.	1.9	39
66	Personalized, bilateral whole-body somatosensory cortex stimulation to relieve fatigue in multiple sclerosis. Multiple Sclerosis Journal, 2018, 24, 1366-1374.	3.0	51
67	Bridging the gap between vaccination with Bacille Calmette-Gu \tilde{A} \otimes rin (BCG) and immunological tolerance: the cases of type 1 diabetes and multiple sclerosis. Current Opinion in Immunology, 2018, 55, 89-96.	5.5	45
68	Analysis of coding and non-coding transcriptome of peripheral B cells reveals an altered interferon response factor (IRF)-1 pathway in multiple sclerosis patients. Journal of Neuroimmunology, 2018, 324, 165-171.	2.3	10
69	BDNF rs6265 polymorphism methylation in Multiple Sclerosis: A possible marker of disease progression. PLoS ONE, 2018, 13, e0206140.	2.5	24
70	Abortion induces reactivation of inflammation in relapsing-remitting multipleÂsclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 1272-1278.	1.9	10
71	The cerebellar topography of attention sub-components in spinocerebellar ataxia type 2. Cortex, 2018, 108, 35-49.	2.4	14
72	Potential Effect of Cyclophosphamide on Bleb Survival in Five Patients with Multiple Sclerosis Who Underwent Glaucoma Surgery. Ophthalmology and Therapy, 2018, 7, 431-436.	2.3	0

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73	Safety and Efficacy of Dimethyl Fumarate in Multiple Sclerosis: An Italian, Multicenter, Real-World Study. CNS Drugs, 2018, 32, 963-970.	5.9	35
74	Liver injury after pulsed methylprednisolone therapy in multiple sclerosis patients. Brain and Behavior, 2018, 8, e00968.	2.2	24
75	Fingolimod vs dimethyl fumarate in multiple sclerosis. Neurology, 2018, 91, e153-e161.	1.1	35
76	Altered intestinal permeability in patients with relapsing–remitting multiple sclerosis: A pilot study. Multiple Sclerosis Journal, 2017, 23, 442-446.	3.0	107
77	Response to the letter to the Editor for the manuscript "Sleep and Fatigue in Multiple Sclerosis: A questionnaire-based, cross-sectional, cohort study―by Tomoyuki Kawada. Journal of the Neurological Sciences, 2017, 373, 142.	0.6	О
78	Association study reveals novel risk loci for sporadic inclusion body myositis. European Journal of Neurology, 2017, 24, 572-577.	3.3	11
79	Focal muscle vibration, an effective rehabilitative approach in severe gait impairment due to multiple sclerosis. Journal of the Neurological Sciences, 2017, 372, 33-39.	0.6	30
80	A phase 3 randomized, double blind, placebo-controlled study to evaluate the efficacy and safety of sialic acid extended-release tablets in patients with GNE myopathy (GNEM). Neuromuscular Disorders, 2017, 27, S150.	0.6	0
81	Identifying Relapses in Multiple Sclerosis Patients through Administrative Data: A Validation Study in the Lazio Region, Italy. Neuroepidemiology, 2017, 48, 171-178.	2.3	6
82	Sativex \hat{A}^{\otimes} effects on promoter methylation and on CNR1 / CNR2 expression in peripheral blood mononuclear cells of progressive multiple sclerosis patients. Journal of the Neurological Sciences, 2017, 379, 298-303.	0.6	11
83	Prevalence and severity of liver injury after pulsed methylprednisolone therapy in multiple sclerosis patients. Journal of Hepatology, 2017, 66, S399-S400.	3.7	O
84	Novel <i>SEC61G</i> â€" <i>EGFR</i> Fusion Gene in Pediatric Ependymomas Discovered by Clonal Expansion of Stem Cells in Absence of Exogenous Mitogens. Cancer Research, 2017, 77, 5860-5872.	0.9	21
85	PLEC gene mutations cause familial disto-proximal myopathy and long QT syndrome mimicking mitochondrial disease. Neuromuscular Disorders, 2017, 27, S150-S151.	0.6	1
86	Sporadic inclusion body myositis: A polygenic disorder?. Neuromuscular Disorders, 2017, 27, S155.	0.6	0
87	Sleep and fatigue in multiple sclerosis: A questionnaire-based, cross-sectional, cohort study. Journal of the Neurological Sciences, 2017, 372, 387-392.	0.6	37
88	Real-world effectiveness of natalizumab and fingolimod compared with self-injectable drugs in non-responders and in treatment-na \tilde{A} ve patients with multiple sclerosis. Journal of Neurology, 2017, 264, 284-294.	3.6	44
89	Sativex in resistant multiple sclerosis spasticity: Discontinuation study in a large population of Italian patients (SA.FE. study). PLoS ONE, 2017, 12, e0180651.	2.5	24
90	Depression in multiple sclerosis: effect of brain derived neurotrophic factor Val66Met polymorphism and disease perception. European Journal of Neurology, 2016, 23, 630-640.	3.3	14

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91	Efficacy and safety of cannabinoid oromucosal spray for multiple sclerosis spasticity. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, 944-951.	1.9	88
92	Expression Profile of Long Non-Coding RNAs in Serum of Patients with Multiple Sclerosis. Journal of Molecular Neuroscience, 2016, 59, 18-23.	2.3	104
93	Huntington's disease and suicidal behavior: The importance of lithium treatment. Clinical Neurology and Neurosurgery, 2016, 145, 108-109.	1.4	5
94	Severe dyspnoea with alteration of the diffusion capacity of the lung associated with fingolimod treatment. Multiple Sclerosis and Related Disorders, 2016, 9, 11-13.	2.0	4
95	D6â€Dna damage in lymphocytes as a predictor of illness evolution in pre-manifest and overt huntington's disease. Journal of Neurology, Neurosurgery and Psychiatry, 2016, 87, A35.3-A36.	1.9	0
96	Riluzole in patients with hereditary cerebellar ataxia – Authors' reply. Lancet Neurology, The, 2016, 15, 789.	10.2	5
97	Transient hair loss during treatment with dimethyl-fumarate for multiple sclerosis. Multiple Sclerosis and Related Disorders, 2016, 7, 68-69.	2.0	6
98	Prevalence of multiple sclerosis in the Lazio region, Italy: use of an algorithm based on health information systems. Journal of Neurology, 2016, 263, 751-759.	3.6	35
99	An atypical case of acute disseminated encephalomyelitis associated with cytomegalovirus infection. Multiple Sclerosis and Related Disorders, 2016, 5, 70-72.	2.0	4
100	Effects of rehabilitation treatment of the upper limb in multiple sclerosis patients and predictive value of neurophysiological measures. European Journal of Physical and Rehabilitation Medicine, 2016, 52, 819-826.	2.2	2
101	Blepharoptosis onset after topical prostaglandin therapy for glaucoma. Clinical and Experimental Ophthalmology, 2015, 43, 689-690.	2.6	1
102	Idiopathic inflammatory myopathies evaluated by near-infrared spectroscopy. Muscle and Nerve, 2015, 51, 830-837.	2.2	3
103	Hereditary Inclusion-Body Myopathies. , 2015, , 1145-1152.		0
104	Analyzing the Effects of a G137V Mutation in the FXN Gene. Frontiers in Molecular Neuroscience, 2015, 8, 66.	2.9	14
105	IFN-Î ² Therapy Regulates TLR7-Mediated Response in Plasmacytoid Dendritic Cells of Multiple Sclerosis Patients Influencing an Anti-Inflammatory Status. Journal of Interferon and Cytokine Research, 2015, 35, 668-681.	1.2	10
106	Second-Line Therapy with Fingolimod for Relapsing-Remitting Multiple Sclerosis in Clinical Practice: The Effect of Previous Exposure to Natalizumab. European Neurology, 2015, 73, 57-65.	1.4	20
107	Primary fibroblasts cultures reveal TDP-43 abnormalities in amyotrophic lateral sclerosis patients with and without SOD1 mutations. Neurobiology of Aging, 2015, 36, 2005.e5-2005.e13.	3.1	42
108	Oculopharyngeal muscular dystrophy: Clinical and neurophysiological features. Clinical Neurophysiology, 2015, 126, 2406-2408.	1.5	12

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109	Acyclovir-related kidney injury during alemtuzumab infusion. Journal of Neurology, 2015, 262, 1772-1774.	3.6	2
110	Epstein-Barr virus genetic variants are associated with multiple sclerosis. Neurology, 2015, 84, 1362-1368.	1.1	44
111	Magnetic resonance imaging pattern recognition in sporadic inclusionâ€body myositis. Muscle and Nerve, 2015, 52, 956-962.	2.2	93
112	A Case of Hemiabdominal Myoclonus. Clinical EEG and Neuroscience, 2015, 46, 331-334.	1.7	2
113	Riluzole in patients with hereditary cerebellar ataxia: a randomised, double-blind, placebo-controlled trial. Lancet Neurology, The, 2015, 14, 985-991.	10.2	163
114	Rituximab as a first-line treatment in pediatric neuromyelitis optica spectrum disorder. Neurological Sciences, 2015, 36, 2301-2302.	1.9	7
115	Hereditary inclusion-body myopathies. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2015, 1852, 644-650.	3.8	33
116	A shared haplotype for dentatorubropallidoluysian atrophy (DRPLA) in Italian families testifies of the recent introduction of the mutation. Journal of Human Genetics, 2014, 59, 153-157.	2.3	6
117	Case of postpartum Parsonage‶urner syndrome. Muscle and Nerve, 2014, 49, 294-295.	2.2	2
118	Moving to Fingolimod From Natalizumab in Multiple Sclerosis: The ENIGM Is Not Solved. JAMA Neurology, 2014, 71, 924.	9.0	2
119	Muscle biopsy features of idiopathic inflammatory myopathies and differential diagnosis. Autoimmunity Highlights, 2014, 5, 77-85.	3.9	63
120	Rasmussen encephalitis: an unusual cause for intractable seizures in elderly. Neurological Sciences, 2014, 35, 143-145.	1.9	4
121	Lower motor neuron involvement in longitudinally extensive transverse myelitis with and without aquaporin-4 antibodies. Clinical Neurophysiology, 2014, 125, 1925-1926.	1.5	1
122	Sleep disorder associated with antibodies to IgLON5: parasomnia or agrypnia?. Lancet Neurology, The, 2014, 13, 864.	10.2	3
123	Distinctive clinical and neuroimaging characteristics of longitudinally extensive transverse myelitis associated with aquaporin-4 autoantibodies. Journal of Neurology, 2013, 260, 2396-2402.	3.6	44
124	Severe Disability in Patients with Relapsing-Remitting Multiple Sclerosis Is Associated with Profound Changes in the Regulation of Leptin Secretion. NeuroImmunoModulation, 2013, 20, 341-347.	1.8	26
125	Cerebellar degeneration associated with mGluR1 autoantibodies as a paraneoplastic manifestation of prostate adenocarcinoma. Journal of Neuroimmunology, 2013, 263, 155-158.	2.3	49
126	Circulating CD56dim NK cells expressing perforin are increased in progressive multiple sclerosis. Journal of Neuroimmunology, 2013, 265, 124-127.	2.3	27

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127	Sleep disordered breathing in a cohort of patients with sporadic inclusion body myositis. Clinical Neurophysiology, 2013, 124, 1615-1621.	1.5	13
128	Mutations in the $3\hat{a}\in^2$ untranslated region of FUS causing FUS overexpression are associated with amyotrophic lateral sclerosis. Human Molecular Genetics, 2013, 22, 4748-4755.	2.9	94
129	Bilateral thoracic long nerve involvement in motor multifocal neuropathy. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 584-584.	1.9	1
130	Atrophy, Fibrosis, and Increased PAX7-Positive Cells in Pharyngeal Muscles of Oculopharyngeal Muscular Dystrophy Patients. Journal of Neuropathology and Experimental Neurology, 2013, 72, 234-243.	1.7	47
131	TWEAK in Inclusion-Body Myositis Muscle. American Journal of Pathology, 2012, 180, 1603-1613.	3.8	30
132	Multi-organ investigation in 16 CADASIL families from central Italy sharing the same R1006C mutation. Neuroscience Letters, 2012, 506, 116-120.	2.1	8
133	Intravascular large B-cell lymphoma presenting as slowly progressive paraparesis with normal MRI features. Journal of the Neurological Sciences, 2012, 314, 171-174.	0.6	4
134	Different Molecular Signatures in Magnetic Resonance Imaging-Staged Facioscapulohumeral Muscular Dystrophy Muscles. PLoS ONE, 2012, 7, e38779.	2.5	106
135	Muscle imaging findings in GNE myopathy. Journal of Neurology, 2012, 259, 1358-1365.	3.6	57
136	The recovery of platelet cyclooxygenase activity explains interindividual variability in responsiveness to lowâ€dose aspirin in patients with and without diabetes. Journal of Thrombosis and Haemostasis, 2012, 10, 1220-1230.	3.8	211
137	Muscle <scp>MRI</scp> in female carriers of dystrophinopathy. European Journal of Neurology, 2012, 19, 1256-1260.	3.3	31
138	Mosaic caveolin-3 expression in acquired rippling muscle disease without evidence of myasthenia gravis or acetylcholine receptor autoantibodies. Neuromuscular Disorders, 2011, 21, 194-203.	0.6	16
139	Mesoangioblasts from Facioscapulohumeral Muscular Dystrophy Display in Vivo a Variable Myogenic Ability Predictable by their in Vitro Behavior. Cell Transplantation, 2011, 20, 1299-1313.	2.5	26
140	CD8+ T Cells in Facioscapulohumeral Muscular Dystrophy Patients with Inflammatory Features at Muscle MRI. Journal of Clinical Immunology, 2011, 31, 155-166.	3.8	113
141	Mixed connective tissue disease presenting as a peculiar myositis with poor muscle regeneration. Neurological Sciences, 2011, 32, 171-174.	1.9	4
142	Pilot trial of simvastatin in the treatment of sporadic inclusion-body myositis. Neurological Sciences, 2011, 32, 841-847.	1.9	33
143	Cerebellar degeneration and ocular myasthenia gravis in a patient with recurring ovarian carcinoma. Neurological Sciences, 2010, 31, 79-81.	1.9	6
144	Analysis of NCAM helps identify unusual phenotypes of hereditary inclusion-body myopathy. Neurology, 2010, 75, 265-272.	1.1	28

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145	Increased aging in primary muscle cultures of sporadic inclusion-body myositis. Neurobiology of Aging, 2010, 31, 1205-1214.	3.1	35
146	An Italian case of hereditary myopathy with early respiratory failure (HMERF) not associated with the titin kinase domain R279W mutation. Neuromuscular Disorders, 2010, 20, 730-734.	0.6	15
147	Vessel-associated stem cells from skeletal muscle: From biology to future uses in cell therapy. World Journal of Stem Cells, 2010, 2, 39.	2.8	15
148	T-bet, pSTAT1 and pSTAT3 expression in peripheral blood mononuclear cells during pregnancy correlates with post-partum activation of multiple sclerosis. Clinical Immunology, 2009, 131, 70-83.	3.2	21
149	Hereditary inclusionâ€body myopathy: Clues on pathogenesis and possible therapy. Muscle and Nerve, 2009, 40, 340-349.	2.2	26
150	Regulatory T cells fail to suppress CD4 ⁺ Tâ€bet ⁺ T cells in relapsing multiple sclerosis patients. Immunology, 2009, 127, 418-428.	4.4	78
151	Progressive multifocal leukoencephalopathy in a patient with Franklin disease and hypogammaglobulinemia. Journal of the Neurological Sciences, 2009, 284, 203-204.	0.6	9
152	An Italian family with inclusionâ€body myopathy and frontotemporal dementia due to mutation in the <i>VCP</i> gene. Muscle and Nerve, 2008, 37, 111-114.	2.2	44
153	Hyposialylation of neprilysin possibly affects its expression and enzymatic activity in hereditary inclusionâ€body myopathy muscle. Journal of Neurochemistry, 2008, 105, 971-981.	3.9	53
154	The persistency of high levels of pSTAT3 expression in circulating CD4+ T cells from CIS patients favors the early conversion to clinically defined multiple sclerosis. Journal of Neuroimmunology, 2008, 205, 126-134.	2.3	22
155	IL17 and IFNÎ ³ production by peripheral blood mononuclear cells from clinically isolated syndrome to secondary progressive multiple sclerosis. Cytokine, 2008, 44, 22-25.	3.2	53
156	Increased expression of T-bet in circulating B cells from a patient with multiple sclerosis and celiac disease. Human Immunology, 2008, 69, 837-839.	2.4	27
157	The ER-Bound RING Finger Protein 5 (RNF5/RMA1) Causes Degenerative Myopathy in Transgenic Mice and Is Deregulated in Inclusion Body Myositis. PLoS ONE, 2008, 3, e1609.	2.5	57
158	Chronic autoimmune autonomic neuropathy responsive to immunosuppressive therapy. Neurology, 2007, 68, 161-162.	1.1	27
159	Gene expression profiling in the early phases of DMD: a constant molecular signature characterizes DMD muscle from early postnatal life throughout disease progression. FASEB Journal, 2007, 21, 1210-1226.	0.5	209
160	A human anti-neuronal autoantibody against GABAB receptor induces experimental autoimmune agrypnia. Experimental Neurology, 2007, 204, 808-818.	4.1	20
161	Isolation and Characterization of Mesoangioblasts from Facioscapulohumeral Muscular Dystrophy Muscle Biopsies. Stem Cells, 2007, 25, 3173-3182.	3.2	37
162	Glucocorticoid treatment reduces T-bet and pSTAT1 expression in mononuclear cells from relapsing remitting multiple sclerosis patients. Clinical Immunology, 2007, 124, 284-293.	3.2	25

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163	The effect of disease activity on leptin, leptin receptor and suppressor of cytokine signalling-3 expression in relapsing–remitting multiple sclerosis. Journal of Neuroimmunology, 2007, 192, 174-183.	2.3	74
164	Brainstem signs with progressing atrophy of medulla oblongata and upper cervical spinal cord. Lancet Neurology, The, 2007, 6, 562-570.	10.2	21
165	Integrated Backscatter in Becker Muscular Dystrophy Patients With Functionally Normal Heart: Myocardial Ultrasound Tissue Characterization Study. Journal of the American College of Cardiology, 2006, 47, 686-688.	2.8	9
166	Neurotrophic factors in relapsing remitting and secondary progressive multiple sclerosis patients during interferon beta therapy. Clinical Immunology, 2006, 118 , 77-82.	3. 2	58
167	Skeletal muscle apoptosis is not increased in gastric cancer patients with mild–moderate weight loss. International Journal of Biochemistry and Cell Biology, 2006, 38, 1561-1570.	2.8	18
168	Reliability of the Hammersmith functional motor scale for spinal muscular atrophy in a multicentric study. Neuromuscular Disorders, 2006, 16, 93-98.	0.6	64
169	Neprilysin participates in skeletal muscle regeneration and is accumulated in abnormal muscle fibres of inclusion body myositis. Journal of Neurochemistry, 2006, 96, 777-789.	3.9	35
170	Coeliac disease presenting with acute disseminated encephalomyelitis. European Journal of Neurology, 2006, 13, 202-203.	3.3	6
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