

Renato Mantegazza

List of Publications by Year in Descending Order

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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.

273
papers

9,053
citations

52
h-index

82
g-index

304
ext. papers

10,625
ext. citations

5.4
avg, IF

5.63
L-index

#	Paper	IF	Citations
273	Antibody Therapies in Autoimmune Neuromuscular Junction Disorders: Approach to Myasthenic Crisis and Chronic Management.. <i>Neurotherapeutics</i> , 2022 , 1	6.4	2
272	Novel Cell-Based Assay for Alpha-3 Nicotinic Receptor Antibodies Detects Antibodies Exclusively in Autoimmune Autonomic Ganglionopathy.. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2022 , 9,	9.1	1
271	Quantitative Muscle MRI Protocol as Possible Biomarker in Becker Muscular Dystrophy. <i>Clinical Neuroradiology</i> , 2021 , 31, 257-266	2.7	7
270	Therapeutic efficacy of 3,4-Diaminopyridine phosphate on neuromuscular junction in Pompe disease. <i>Biomedicine and Pharmacotherapy</i> , 2021 , 137, 111357	7.5	1
269	Dysregulation of Muscle-Specific MicroRNAs as Common Pathogenic Feature Associated with Muscle Atrophy in ALS, SMA and SBMA: Evidence from Animal Models and Human Patients. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	4
268	MyomiRs and their multifaceted regulatory roles in muscle homeostasis and amyotrophic lateral sclerosis. <i>Journal of Cell Science</i> , 2021 , 134,	5.3	3
267	Safety, efficacy, and tolerability of efgartigimod in patients with generalised myasthenia gravis (ADAPT): a multicentre, randomised, placebo-controlled, phase 3 trial. <i>Lancet Neurology</i> , 2021 , 20, 526-536	24.1	29
266	Post-intervention Status in Patients With Refractory Myasthenia Gravis Treated With Eculizumab During REGAIN and Its Open-Label Extension. <i>Neurology</i> , 2021 , 96, e610-e618	6.5	16
265	Next-generation sequencing application to investigate skeletal muscle channelopathies in a large cohort of Italian patients. <i>Neuromuscular Disorders</i> , 2021 , 31, 336-347	2.9	4
264	Immune Soluble Factors in the Cerebrospinal Fluid of Progressive Multiple Sclerosis Patients Segregate Into Two Groups. <i>Frontiers in Immunology</i> , 2021 , 12, 633167	8.4	3
263	Patient-reported impact of myasthenia gravis in the real world: protocol for a digital observational study (MyRealWorld MG). <i>BMJ Open</i> , 2021 , 11, e048198	3	0
262	Eculizumab in refractory generalized myasthenia gravis previously treated with rituximab: subgroup analysis of REGAIN and its extension study. <i>Muscle and Nerve</i> , 2021 , 64, 662-669	3.4	1
261	Iatrogenic Kaposi's sarcoma in myasthenia gravis: learnings from two case reports. <i>Neurological Sciences</i> , 2021 , 42, 2081-2083	3.5	1
260	Multiomic elucidation of a coding 99-mer repeat-expansion skeletal muscle disease. <i>Acta Neuropathologica</i> , 2020 , 140, 231-235	14.3	6
259	miR-146a in Myasthenia Gravis Thymus Bridges Innate Immunity With Autoimmunity and Is Linked to Therapeutic Effects of Corticosteroids. <i>Frontiers in Immunology</i> , 2020 , 11, 142	8.4	17
258	Cytokine Profile in Striated Muscle Laminopathies: New Promising Biomarkers for Disease Prediction. <i>Cells</i> , 2020 , 9,	7.9	2
257	Eculizumab for the treatment of myasthenia gravis. <i>Expert Opinion on Biological Therapy</i> , 2020 , 20, 991-998	9.4	4

256	Employment in Myasthenia Gravis: A Systematic Literature Review and Meta-Analysis. <i>Neuroepidemiology</i> , 2020 , 54, 304-312	5.4	2
255	Circulating MyomiRs as Potential Biomarkers to Monitor Response to Nusinersen in Pediatric SMA Patients. <i>Biomedicines</i> , 2020 , 8,	4.8	15
254	Guidance for the management of myasthenia gravis (MG) and Lambert-Eaton myasthenic syndrome (LEMS) during the COVID-19 pandemic. <i>Journal of the Neurological Sciences</i> , 2020 , 412, 116803	3.2	81
253	Complement Inhibition for the Treatment of Myasthenia Gravis. <i>ImmunoTargets and Therapy</i> , 2020 , 9, 317-331	9	11
252	Glycogen storage in a zebrafish Pompe disease model is reduced by 3-BrPA treatment. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2020 , 1866, 165662	6.9	4
251	COVID-19-associated risks and effects in myasthenia gravis (CARE-MG). <i>Lancet Neurology</i> , 2020 , 19, 970-971	24.1	45
250	Clinical and Molecular Spectrum of Myotonia and Periodic Paralysis Associated With Mutations in a Large Cohort of Italian Patients. <i>Frontiers in Neurology</i> , 2020 , 11, 646	4.1	3
249	Consistent improvement with eculizumab across muscle groups in myasthenia gravis. <i>Annals of Clinical and Translational Neurology</i> , 2020 , 7, 1327-1339	5.3	6
248	From Traditional to Targeted Immunotherapy in Myasthenia Gravis: Prospects for Research. <i>Frontiers in Neurology</i> , 2020 , 11, 981	4.1	7
247	Reliability and Repeatability Analysis of Indices to Measure Gait Deterioration in MS Patients during Prolonged Walking. <i>Sensors</i> , 2020 , 20,	3.8	3
246	Pharmacogenetic and pharmaco-miR biomarkers for tailoring and monitoring myasthenia gravis treatments. <i>Expert Review of Precision Medicine and Drug Development</i> , 2020 , 5, 317-329	1.6	2
245	Nusinersen safety and effects on motor function in adult spinal muscular atrophy type 2 and 3. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020 , 91, 1166-1174	5.5	32
244	2019 ,		3
243	Long-term effect of thymectomy plus prednisone versus prednisone alone in patients with non-thymomatous myasthenia gravis: 2-year extension of the MGTX randomised trial. <i>Lancet Neurology</i> , 2019 , 18, 259-268	24.1	74
242	Randomized phase 2 study of FcRn antagonist efgartigimod in generalized myasthenia gravis. <i>Neurology</i> , 2019 , 92, e2661-e2673	6.5	90
241	Eculizumab improves fatigue in refractory generalized myasthenia gravis. <i>Quality of Life Research</i> , 2019 , 28, 2247-2254	3.7	19
240	FM19G11-Loaded Gold Nanoparticles Enhance the Proliferation and Self-Renewal of Ependymal Stem Progenitor Cells Derived from ALS Mice. <i>Cells</i> , 2019 , 8,	7.9	13
239	Aging-associated genes and microRNAs: a contribution to myogenic program dysregulation in oculopharyngeal muscular dystrophy. <i>FASEB Journal</i> , 2019 , 33, 7155-7167	0.9	6

238	Hyperexcitability in Cultured Cortical Neuron Networks from the G93A-SOD1 Amyotrophic Lateral Sclerosis Model Mouse and its Molecular Correlates. <i>Neuroscience</i> , 2019 , 416, 88-99	3.9	10
237	MicroRNA signature associated with treatment response in myasthenia gravis: A further step towards precision medicine. <i>Pharmacological Research</i> , 2019 , 148, 104388	10.2	10
236	Long-term efficacy and safety of eculizumab in Japanese patients with generalized myasthenia gravis: A subgroup analysis of the REGAIN open-label extension study. <i>Journal of the Neurological Sciences</i> , 2019 , 407, 116419	3.2	8
235	Amifampridine tablets for the treatment of Lambert-Eaton myasthenic syndrome. <i>Expert Review of Clinical Pharmacology</i> , 2019 , 12, 1013-1018	3.8	7
234	Italian recommendations for the diagnosis and treatment of myasthenia gravis. <i>Neurological Sciences</i> , 2019 , 40, 1111-1124	3.5	23
233	Long-term safety and efficacy of eculizumab in generalized myasthenia gravis. <i>Muscle and Nerve</i> , 2019 , 60, 14-24	3.4	76
232	Therapeutic Effect of Bifidobacterium Administration on Experimental Autoimmune Myasthenia Gravis in Lewis Rats. <i>Frontiers in Immunology</i> , 2019 , 10, 2949	8.4	7
231	Diagnosis and treatment of myasthenia gravis. <i>Current Opinion in Rheumatology</i> , 2019 , 31, 623-633	5.3	15
230	Italian recommendations for diagnosis and management of congenital myasthenic syndromes. <i>Neurological Sciences</i> , 2019 , 40, 457-468	3.5	16
229	Longitudinal evaluation of SMN levels as biomarker for spinal muscular atrophy: results of a phase IIb double-blind study of salbutamol. <i>Journal of Medical Genetics</i> , 2019 , 56, 293-300	5.8	16
228	Pharmacogenetics of myotonic hNav1.4 sodium channel variants situated near the fast inactivation gate. <i>Pharmacological Research</i> , 2019 , 141, 224-235	10.2	12
227	Older age, higher perceived disability and depressive symptoms predict the amount and severity of work-related difficulties in persons with multiple sclerosis. <i>Disability and Rehabilitation</i> , 2019 , 41, 2255-2263	2.4	3
226	Botulinum toxin type A affects the transcriptome of cell cultures derived from muscle biopsies of controls and spastic patients. <i>Toxicology in Vitro</i> , 2018 , 50, 124-136	3.6	1
225	Toll-like receptors 7 and 9 in myasthenia gravis thymus: amplifiers of autoimmunity?. <i>Annals of the New York Academy of Sciences</i> , 2018 , 1413, 11-24	6.5	18
224	Gut microbiota and probiotics: novel immune system modulators in myasthenia gravis?. <i>Annals of the New York Academy of Sciences</i> , 2018 , 1413, 49-58	6.5	22
223	When myasthenia gravis is deemed refractory: clinical signposts and treatment strategies. <i>Therapeutic Advances in Neurological Disorders</i> , 2018 , 11, 1756285617749134	6.6	74
222	Exosomes and exosomal miRNAs from muscle-derived fibroblasts promote skeletal muscle fibrosis. <i>Matrix Biology</i> , 2018 , 74, 77-100	11.4	37
221	Up-regulation of Toll-like receptors 7 and 9 and its potential implications in the pathogenic mechanisms of LMNA-related myopathies. <i>Nucleus</i> , 2018 , 9, 398-409	3.9	9

220	Severe multiple sclerosis reactivation during prolonged lymphopenia after dimethyl fumarate discontinuation. <i>Acta Neurologica Scandinavica</i> , 2018 , 137, 623-625	3.8	13
219	Paraneoplastic autoimmune diseases in patients with thymic malignancies: a favorable, but not independent, prognostic factor. <i>Mediastinum</i> , 2018 , 2, 41-41	0.5	
218	Administration of bifidobacterium and lactobacillus strains modulates experimental myasthenia gravis and experimental encephalomyelitis in Lewis rats. <i>Oncotarget</i> , 2018 , 9, 22269-22287	3.3	22
217	Amifampridine phosphate in the treatment of muscle-specific kinase myasthenia gravis: a phase IIb, randomized, double-blind, placebo-controlled, double crossover study. <i>SAGE Open Medicine</i> , 2018 , 6, 2050312118819013	2.4	18
216	Myasthenia gravis: from autoantibodies to therapy. <i>Current Opinion in Neurology</i> , 2018 , 31, 517-525	7.1	25
215	Atypical Post-Injection Reactions with Delayed Onset Following Glatiramer Acetate 40mg: Need for Titration?. <i>CNS Drugs</i> , 2018 , 32, 653-660	6.7	1
214	Difficulties in adjustment to multiple sclerosis: vulnerability and unpredictability of illness in the foreground. <i>Disability and Rehabilitation</i> , 2017 , 39, 897-903	2.4	19
213	Long-term cardiovascular autonomic and clinical changes after immunoglobulin G immunoadsorption therapy in autoimmune autonomic ganglionopathy. <i>Journal of Hypertension</i> , 2017 , 35, 1513-1520	1.9	5
212	A propensity score analysis for comparison of T-3b and VATET in myasthenia gravis. <i>Neurology</i> , 2017 , 89, 189-195	6.5	8
211	Lambert-Eaton myasthenic syndrome (LEMS): a rare autoimmune presynaptic disorder often associated with cancer. <i>Journal of Neurology</i> , 2017 , 264, 1854-1863	5.5	38
210	A longitudinal DTI and histological study of the spinal cord reveals early pathological alterations in G93A-SOD1 mouse model of amyotrophic lateral sclerosis. <i>Experimental Neurology</i> , 2017 , 293, 43-52	5.7	16
209	Validation of the italian version of the 15-item Myasthenia Gravis Quality-of-Life questionnaire. <i>Muscle and Nerve</i> , 2017 , 56, 716-720	3.4	8
208	Coexistence of CLCN1 and SCN4A mutations in one family suffering from myotonia. <i>Neurogenetics</i> , 2017 , 18, 219-225	3	13
207	Safety and efficacy of eculizumab in anti-acetylcholine receptor antibody-positive refractory generalised myasthenia gravis (REGAIN): a phase 3, randomised, double-blind, placebo-controlled, multicentre study. <i>Lancet Neurology</i> , 2017 , 16, 976-986	24.1	278
206	A novel ABCC6 haplotype is associated with azathioprine drug response in myasthenia gravis. <i>Pharmacogenetics and Genomics</i> , 2017 , 27, 51-56	1.9	5
205	Validity, reliability, and sensitivity to change of the myasthenia gravis activities of daily living profile in a sample of Italian myasthenic patients. <i>Neurological Sciences</i> , 2017 , 38, 1927-1931	3.5	4
204	Epstein-Barr virus in tumor-infiltrating B cells of myasthenia gravis thymoma: an innocent bystander or an autoimmunity mediator?. <i>Oncotarget</i> , 2017 , 8, 95432-95449	3.3	13
203	Biobank of Cells, Tissues and DNA from Patients with Neuromuscular Diseases: An Indispensable link between Clinical Centers and the Scientific Community. <i>Open Journal of Bioresources</i> , 2017 , 4,	0.9	3

202	Severe articular and musculoskeletal pain: An unexpected side effect of dimethyl-fumarate therapy for multiple sclerosis. <i>Journal of the Neurological Sciences</i> , 2016 , 369, 139-140	3.2	9
201	A novel infection- and inflammation-associated molecular signature in peripheral blood of myasthenia gravis patients. <i>Immunobiology</i> , 2016 , 221, 1227-36	3.4	22
200	Psychosocial difficulties of individuals with multiple sclerosis: the PARADISE-24 questionnaire. <i>International Journal of Rehabilitation Research</i> , 2016 , 39, 339-345	1.8	7
199	Multidisciplinary study of a new CIC-1 mutation causing myotonia congenita: a paradigm to understand and treat ion channelopathies. <i>FASEB Journal</i> , 2016 , 30, 3285-3295	0.9	15
198	Increased expression of Toll-like receptors 7 and 9 in myasthenia gravis thymus characterized by active Epstein-Barr virus infection. <i>Immunobiology</i> , 2016 , 221, 516-27	3.4	36
197	Titin antibodies in "seronegative" myasthenia gravis--A new role for an old antigen. <i>Journal of Neuroimmunology</i> , 2016 , 292, 108-15	3.5	37
196	Multiple Sclerosis Questionnaire for Job Difficulties (MSQ-Job): definition of the cut-off score. <i>Neurological Sciences</i> , 2016 , 37, 777-80	3.5	2
195	Anti-fibrotic effect of pirfenidone in muscle derived-fibroblasts from Duchenne muscular dystrophy patients. <i>Life Sciences</i> , 2016 , 145, 127-36	6.8	19
194	Comparison of Diffusion MRI Acquisition Protocols for the In Vivo Characterization of the Mouse Spinal Cord: Variability Analysis and Application to an Amyotrophic Lateral Sclerosis Model. <i>PLoS ONE</i> , 2016 , 11, e0161646	3.7	5
193	Animal models of myasthenia gravis: utility and limitations. <i>International Journal of General Medicine</i> , 2016 , 9, 53-64	2.3	22
192	Cognitive function alone is a poor predictor of health-related quality of life in employed patients with MS: results from a cross-sectional study. <i>Clinical Neuropsychologist</i> , 2016 , 30, 201-15	4.4	7
191	Validation of the MG-DIS: a disability assessment for myasthenia gravis. <i>Journal of Neurology</i> , 2016 , 263, 871-882	5.5	13
190	Early effect of dalfampridine in patients with MS: A multi-instrumental approach to better investigate responsiveness. <i>Journal of the Neurological Sciences</i> , 2016 , 368, 402-7	3.2	12
189	Identification of a gene expression signature in peripheral blood of multiple sclerosis patients treated with disease-modifying therapies. <i>Clinical Immunology</i> , 2016 , 173, 133-146	9	4
188	Rare variants in SQSTM1 and VCP genes and risk of sporadic inclusion body myositis. <i>Neurobiology of Aging</i> , 2016 , 47, 218.e1-218.e9	5.6	29
187	Validation of the Besta Neurological Institute rating scale for myasthenia gravis. <i>Muscle and Nerve</i> , 2016 , 53, 32-7	3.4	10
186	Up-regulation of neural and cell cycle-related microRNAs in brain of amyotrophic lateral sclerosis mice at late disease stage. <i>Molecular Brain</i> , 2015 , 8, 5	4.5	36
185	MuSK autoantibodies in myasthenia gravis detected by cell based assay--A multinational study. <i>Journal of Neuroimmunology</i> , 2015 , 284, 10-7	3.5	48

184	Opposing roles of miR-21 and miR-29 in the progression of fibrosis in Duchenne muscular dystrophy. <i>Biochimica Et Biophysica Acta - Molecular Basis of Disease</i> , 2015 , 1852, 1451-64	6.9	60
183	CLC-1 mutations in myotonia congenita patients: insights into molecular gating mechanisms and genotype-phenotype correlation. <i>Journal of Physiology</i> , 2015 , 593, 4181-99	3.9	21
182	CHARACTERIZATION BY MOLECULAR MARKERS OF POMPIANA NATURAL CITRUS HYBRID CULTIVATED IN SARDINIA. <i>Acta Horticulturae</i> , 2015 , 165-172	0.3	9
181	Development and validation of the multiple sclerosis questionnaire for the evaluation of job difficulties (MSQ-Job). <i>Acta Neurologica Scandinavica</i> , 2015 , 132, 226-34	3.8	8
180	CLC-1 chloride channels: state-of-the-art research and future challenges. <i>Frontiers in Cellular Neuroscience</i> , 2015 , 9, 156	6.1	38
179	The effects of an intronic polymorphism in TOMM40 and APOE genotypes in sporadic inclusion body myositis. <i>Neurobiology of Aging</i> , 2015 , 36, 1766.e1-1766.e3	5.6	13
178	Home-based palliative approach for people with severe multiple sclerosis and their carers: study protocol for a randomized controlled trial. <i>Trials</i> , 2015 , 16, 184	2.8	24
177	The European LEMS Registry: Baseline Demographics and Treatment Approaches. <i>Neurology and Therapy</i> , 2015 , 4, 105-24	4.6	23
176	In vivo quantitative magnetization transfer imaging correlates with histology during de- and remyelination in cuprizone-treated mice. <i>NMR in Biomedicine</i> , 2015 , 28, 327-37	4.4	61
175	Human CD14+ cells loaded with Paclitaxel inhibit in vitro cell proliferation of glioblastoma. <i>Cytotherapy</i> , 2015 , 17, 310-9	4.8	11
174	Autoimmune Ion Channel Disorders of the Peripheral Nervous System 2015 , 457-468		
173	Italian recommendations for Lambert-Eaton myasthenic syndrome (LEMS) management. <i>Neurological Sciences</i> , 2014 , 35, 515-20	3.5	16
172	Altered miRNA expression is associated with neuronal fate in G93A-SOD1 ependymal stem progenitor cells. <i>Experimental Neurology</i> , 2014 , 253, 91-101	5.7	27
171	A comprehensive analysis of the epidemiology and clinical characteristics of anti-LRP4 in myasthenia gravis. <i>Journal of Autoimmunity</i> , 2014 , 52, 139-45	15.5	187
170	Fibrosis and inflammation are greater in muscles of beta-sarcoglycan-null mouse than mdx mouse. <i>Cell and Tissue Research</i> , 2014 , 356, 427-43	4.2	18
169	Safety of the first dose of fingolimod for multiple sclerosis: results of an open-label clinical trial. <i>BMC Neurology</i> , 2014 , 14, 65	3.1	43
168	Innate immunity in myasthenia gravis thymus: pathogenic effects of Toll-like receptor 4 signaling on autoimmunity. <i>Journal of Autoimmunity</i> , 2014 , 52, 74-89	15.5	52
167	Autophagy, inflammation and innate immunity in inflammatory myopathies. <i>PLoS ONE</i> , 2014 , 9, e111490	3.7	30

166	European database for myasthenia gravis: a model for an international disease registry. <i>Neurology</i> , 2014 , 83, 189-91	6.5	7
165	Development of the MG-DIS: an ICF-based disability assessment instrument for myasthenia gravis. <i>Disability and Rehabilitation</i> , 2014 , 36, 546-55	2.4	12
164	LMNA-associated myopathies: the Italian experience in a large cohort of patients. <i>Neurology</i> , 2014 , 83, 1634-44	6.5	45
163	VAV1 and BAFF, via NFB pathway, are genetic risk factors for myasthenia gravis. <i>Annals of Clinical and Translational Neurology</i> , 2014 , 1, 329-39	5.3	22
162	LRP4 antibodies in serum and CSF from amyotrophic lateral sclerosis patients. <i>Annals of Clinical and Translational Neurology</i> , 2014 , 1, 80-7	5.3	65
161	Experience of an information aid for newly diagnosed multiple sclerosis patients: a qualitative study on the SIMS-Trial. <i>Health Expectations</i> , 2014 , 17, 36-48	3.7	11
160	Association of increased progression-free survival in primary glioblastomas with lymphopenia at baseline and activation of NK and NKT cells after dendritic cell immunotherapy.. <i>Journal of Clinical Oncology</i> , 2014 , 32, 2087-2087	2.2	1
159	Orphan drugs to treat myasthenia gravis. <i>Expert Opinion on Orphan Drugs</i> , 2013 , 1, 373-384	1.1	1
158	Duchenne muscular dystrophy fibroblast nodules: a cell-based assay for screening anti-fibrotic agents. <i>Cell and Tissue Research</i> , 2013 , 352, 659-70	4.2	13
157	Exacerbation of experimental autoimmune encephalomyelitis by passive transfer of IgG antibodies from a multiple sclerosis patient responsive to immunoadsorption. <i>Journal of Neuroimmunology</i> , 2013 , 262, 19-26	3.5	9
156	Etiology of myasthenia gravis: innate immunity signature in pathological thymus. <i>Autoimmunity Reviews</i> , 2013 , 12, 863-74	13.6	64
155	Marked phenotypic variability in two siblings with congenital myasthenic syndrome due to mutations in MUSK. <i>Journal of Neurology</i> , 2013 , 260, 2894	5.5	22
154	Percutaneous vertebroplasty in a series of myasthenic patients with steroid-induced symptomatic vertebral fractures. <i>Neurological Sciences</i> , 2013 , 34, 773-6	3.5	2
153	Differential targeting of immune-cells by Pixantrone in experimental myasthenia gravis. <i>Journal of Neuroimmunology</i> , 2013 , 258, 41-50	3.5	3
152	P20.4 Antithetic role of miR-21 and miR-29 in the progression of fibrosis in Duchenne muscular dystrophy. <i>Neuromuscular Disorders</i> , 2013 , 23, 839-840	2.9	2
151	A large cohort of myotonia congenita probands: novel mutations and a high-frequency mutation region in exons 4 and 5 of the CLCN1 gene. <i>Journal of Human Genetics</i> , 2013 , 58, 581-7	4.3	32
150	Clinical and molecular cross-sectional study of a cohort of adult type III spinal muscular atrophy patients: clues from a biomarker study. <i>European Journal of Human Genetics</i> , 2013 , 21, 630-6	5.3	32
149	Complete stable remission and autoantibody specificity in myasthenia gravis. <i>Neurology</i> , 2013 , 80, 188-96.5		43

148	Factors related to difficulties with employment in patients with multiple sclerosis: a review of 2002-2011 literature. <i>International Journal of Rehabilitation Research</i> , 2013 , 36, 105-11	1.8	86
147	A new thiopurine s-methyltransferase haplotype associated with intolerance to azathioprine. <i>Journal of Clinical Pharmacology</i> , 2013 , 53, 67-74	2.9	20
146	The natural killer cell response and tumor debulking are associated with prolonged survival in recurrent glioblastoma patients receiving dendritic cells loaded with autologous tumor lysates. <i>Oncolmmunology</i> , 2013 , 2, e23401	7.2	42
145	Comparative neuronal differentiation of self-renewing neural progenitor cell lines obtained from human induced pluripotent stem cells. <i>Frontiers in Cellular Neuroscience</i> , 2013 , 7, 175	6.1	26
144	Sleep breathing disorders in 40 Italian patients with Myotonic dystrophy type 1. <i>Neuromuscular Disorders</i> , 2012 , 22, 219-24	2.9	37
143	Patient registries: useful tools for clinical research in myasthenia gravis. <i>Annals of the New York Academy of Sciences</i> , 2012 , 1274, 107-13	6.5	5
142	Recommendations for myasthenia gravis clinical trials. <i>Muscle and Nerve</i> , 2012 , 45, 909-17	3.4	86
141	Autoimmune mechanisms in myasthenia gravis. <i>Current Opinion in Neurology</i> , 2012 , 25, 621-9	7.1	52
140	Treatment of myasthenia gravis: focus on pyridostigmine. <i>Clinical Drug Investigation</i> , 2011 , 31, 691-701	3.2	56
139	Current and emerging therapies for the treatment of myasthenia gravis. <i>Neuropsychiatric Disease and Treatment</i> , 2011 , 7, 151-60	3.1	52
138	Human neurotrophin receptor p75NTR defines differentiation-oriented skeletal muscle precursor cells: implications for muscle regeneration. <i>Journal of Neuropathology and Experimental Neurology</i> , 2011 , 70, 133-42	3.1	22
137	Osteopontin is highly expressed in severely dystrophic muscle and seems to play a role in muscle regeneration and fibrosis. <i>Histopathology</i> , 2011 , 59, 1215-28	7.3	41
136	Fibroblasts from the muscles of Duchenne muscular dystrophy patients are resistant to cell detachment apoptosis. <i>Experimental Cell Research</i> , 2011 , 317, 2536-47	4.2	25
135	Variable disease severity in Saudi Arabian and Sudanese families with c.3924 + 2 T > C mutation of LAMA2. <i>BMC Research Notes</i> , 2011 , 4, 534	2.3	7
134	Hind limb muscle atrophy precedes cerebral neuronal degeneration in G93A-SOD1 mouse model of amyotrophic lateral sclerosis: a longitudinal MRI study. <i>Experimental Neurology</i> , 2011 , 231, 30-7	5.7	67
133	Pilot trial of simvastatin in the treatment of sporadic inclusion-body myositis. <i>Neurological Sciences</i> , 2011 , 32, 841-7	3.5	30
132	The thymus in myasthenia gravis: Site of "innate autoimmunity"?. <i>Muscle and Nerve</i> , 2011 , 44, 467-84	3.4	46
131	Epstein-Barr virus in myasthenia gravis thymus: a matter of debate. <i>Annals of Neurology</i> , 2011 , 70, 519	9.4	7

130	Importance of Shank3 protein in regulating metabotropic glutamate receptor 5 (mGluR5) expression and signaling at synapses. <i>Journal of Biological Chemistry</i> , 2011 , 286, 34839-50	5.4	145
129	Type I interferon and Toll-like receptor expression characterizes inflammatory myopathies. <i>Neurology</i> , 2011 , 76, 2079-88	6.5	64
128	Myasthenia gravis. <i>Autoimmune Diseases</i> , 2011 , 2011, 697575	2.9	1
127	Inflammation and epstein-barr virus infection are common features of myasthenia gravis thymus: possible roles in pathogenesis. <i>Autoimmune Diseases</i> , 2011 , 2011, 213092	2.9	18
126	The Multiple Sclerosis Knowledge Questionnaire: a self-administered instrument for recently diagnosed patients. <i>Multiple Sclerosis Journal</i> , 2010 , 16, 100-11	5	38
125	Naturally occurring CD4+CD25+ regulatory T cells prevent but do not improve experimental myasthenia gravis. <i>Journal of Immunology</i> , 2010 , 185, 5656-67	5.3	21
124	Development and validation of a patient self-assessed questionnaire on satisfaction with communication of the multiple sclerosis diagnosis. <i>Multiple Sclerosis Journal</i> , 2010 , 16, 1237-47	5	22
123	Detection of poliovirus-infected macrophages in thymus of patients with myasthenia gravis. <i>Neurology</i> , 2010 , 74, 1118-26	6.5	48
122	Calsequestrin and junctin immunoreactivity in hexagonally cross-linked tubular arrays myopathy. <i>Neuromuscular Disorders</i> , 2010 , 20, 326-9	2.9	3
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