

Michael Swash

List of Publications by Year in descending order

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Version: 2024-02-01

548
papers

31,151
citations

8181

76
h-index

6836

155
g-index

566
all docs

566
docs citations

566
times ranked

15604
citing authors

#	ARTICLE	IF	CITATIONS
1	Respiratory onset in amyotrophic lateral sclerosis: clinical features and spreading pattern. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2023, 24, 40-44.	1.7	7
2	Henry Head's lifelong studies of cutaneous sensation. <i>Journal of Medical Biography</i> , 2022, 30, 57-63.	0.1	0
3	Exercise following immobility increases lower motor neuron excitability: F-wave and H-reflex studies. <i>Neurophysiologie Clinique</i> , 2022, , .	2.2	2
4	Respiratory function tests in amyotrophic lateral sclerosis: The role of maximal voluntary ventilation. <i>Journal of the Neurological Sciences</i> , 2022, 434, 120143.	0.6	4
5	The senile hand: Age effects on intrinsic hand muscle <scp>CMAP</scp> amplitudes influence splitâ€hand index calculations. <i>Muscle and Nerve</i> , 2022, 65, 463-467.	2.2	8
6	Vulvodynia: a neuroinflammatory pain syndrome originating in pelvic visceral nerve plexuses due to mechanical factors. <i>Archives of Gynecology and Obstetrics</i> , 2022, 306, 1411-1415.	1.7	4
7	Modulation of spinal inhibition in amyotrophic lateral sclerosis. <i>Acta Physiologica</i> , 2022, 234, e13801.	3.8	1
8	Thyroid dysfunction in Portuguese amyotrophic lateral sclerosis patients. <i>Neurological Sciences</i> , 2022, 43, 5625-5627.	1.9	1
9	Mouth occlusion pressure at 100ms (PO.1) as a respiratory biomarker in amyotrophic lateral sclerosis. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2021, 22, 53-60.	1.7	7
10	The cutaneous silent period in motor neuron disease. <i>Clinical Neurophysiology</i> , 2021, 132, 660-665.	1.5	8
11	Cardiovascular comorbidities in amyotrophic lateral sclerosis. <i>Journal of the Neurological Sciences</i> , 2021, 421, 117292.	0.6	10
12	Compensatory metabolic and central respiratory drive mechanisms in ALS. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2021, 22, 1-3.	1.7	2
13	Motor neuron disease beginning with frontotemporal dementia: clinical features and progression. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2021, 22, 508-516.	1.7	7
14	FOWLER'S SYNDROME: WHAT IT IS AND WHAT IT'S NOT. <i>Pelviperrineology</i> , 2021, 39, 107-114.	0.1	0
15	Benign fasciculations: A follow-up study with electrophysiological studies. <i>Muscle and Nerve</i> , 2021, 64, 670-675.	2.2	3
16	Levosimendan for amyotrophic lateral sclerosis. <i>Lancet Neurology</i> , The, 2021, 20, 775-777.	10.2	0
17	Electromyographic findings in primary lateral sclerosis during disease progression. <i>Clinical Neurophysiology</i> , 2021, 132, 2996-3001.	1.5	6
18	Delayed Diagnosis and Diagnostic Pathway of ALS Patients in Portugal: Where Can We Improve?. <i>Frontiers in Neurology</i> , 2021, 12, 761355.	2.4	12

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19	Professor Henry Ulrich 1916-2015: A London neuropathologist and his Polish heritage. <i>Journal of Medical Biography</i> , 2020, 28, 8-15.	0.1	0
20	Cervical muscle weakness is a marker of respiratory dysfunction in amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020, 91, 323-324.	1.9	3
21	Fasciculations: Opening Pandora's box. <i>Clinical Neurophysiology</i> , 2020, 131, 239-240.	1.5	0
22	Measuring spinal presynaptic inhibition in human subjects. <i>Clinical Neurophysiology</i> , 2020, 131, 1966-1967.	1.5	0
23	Immobility and F waves: Impact on lower motor neuron excitability. <i>Muscle and Nerve</i> , 2020, 61, 480-484.	2.2	3
24	Respiratory Neurophysiology in Intensive Care Unit. <i>Journal of Clinical Neurophysiology</i> , 2020, 37, 208-210.	1.7	0
25	Intensive Care Unit-Acquired Weakness: Neuropathology. <i>Journal of Clinical Neurophysiology</i> , 2020, 37, 197-199.	1.7	8
26	Sir William Osler and the Schorstein Memorial lectures at the London Hospital. <i>Journal of Medical Biography</i> , 2020, , 096777202092451.	0.1	0
27	Spreading in ALS: The relative impact of upper and lower motor neuron involvement. <i>Annals of Clinical and Translational Neurology</i> , 2020, 7, 1181-1192.	3.7	34
28	Motor unit estimation by MRI: Integrating old and new ideas. <i>Clinical Neurophysiology</i> , 2020, 131, 1379-1380.	1.5	1
29	Hypothesis: amyotrophic lateral sclerosis and environmental pollutants. <i>Muscle and Nerve</i> , 2020, 62, 187-191.	2.2	5
30	Occasional essay: Upper motor neuron syndrome in amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020, 91, 227-234.	1.9	26
31	Chitinases, neuroinflammation and biomarkers in ALS. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020, 91, 338-338.	1.9	6
32	Diaphragmatic CMAP amplitude from phrenic nerve stimulation predicts functional decline in ALS. <i>Journal of Neurology</i> , 2020, 267, 2123-2129.	3.6	5
33	A proposal for new diagnostic criteria for ALS. <i>Clinical Neurophysiology</i> , 2020, 131, 1975-1978.	1.5	268
34	The "split-leg" syndrome in ALS: specific or variable?. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2019, 20, 615-616.	1.7	1
35	Face-making: task-specific facial tensions and grimacing in musicians. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 1180-1182.	1.9	1
36	Testing electrolyte supplementation for muscle cramp. <i>Muscle and Nerve</i> , 2019, 60, 499-500.	2.2	1

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37	Interleukin-6 and amyotrophic lateral sclerosis. <i>Journal of the Neurological Sciences</i> , 2019, 398, 50-53.	0.6	29
38	The α -neurophysiological index™ predicts survival in amyotrophic lateral sclerosis. <i>Clinical Neurophysiology</i> , 2019, 130, 1684-1685.	1.5	1
39	The split hand in amyotrophic lateral sclerosis: a possible role for the neuromuscular junction. <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2019, 20, 368-375.	1.7	16
40	Motor unit recruitment in myopathy: The myopathic EMG reconsidered. <i>Journal of Electromyography and Kinesiology</i> , 2019, 45, 41-45.	1.7	2
41	Diaphragmatic Neurophysiology and Respiratory Markers in ALS. <i>Frontiers in Neurology</i> , 2019, 10, 143.	2.4	38
42	Sensory modulation of fasciculation discharge frequency. <i>Muscle and Nerve</i> , 2019, 59, 688-693.	2.2	4
43	Neurology and the homeless. <i>Neurology</i> , 2019, 92, 1131-1132.	1.1	0
44	Clinical trials in the ALS syndrome: it is time for change. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, jnnp-2019-321411.	1.9	4
45	Muscular cramp: causes and management. <i>European Journal of Neurology</i> , 2019, 26, 214-221.	3.3	27
46	Reading the palm with MUNIX: A α -reversed split hand™ in spinal muscular atrophy. <i>Clinical Neurophysiology</i> , 2019, 130, 305-306.	1.5	5
47	Diagnostic criteria for amyotrophic lateral sclerosis: A multicentre study of inter-rater variation and sensitivity. <i>Clinical Neurophysiology</i> , 2019, 130, 307-314.	1.5	46
48	Medical conferences: value for money?. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2019, 90, 483-484.	1.9	6
49	Sensorimotor integration is problematic in amyotrophic lateral sclerosis. <i>Clinical Neurophysiology</i> , 2018, 129, 849-850.	1.5	0
50	Diaphragm motor responses to phrenic nerve stimulation in ALS: Surface and needle recordings. <i>Clinical Neurophysiology</i> , 2018, 129, 349-353.	1.5	11
51	Relationships between neurologists and industry. <i>Neurology</i> , 2018, 90, 1047-1048.	1.1	16
52	Physical activity as a risk factor in ALS. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018, 89, 793-793.	1.9	1
53	Plasma level of clubcell (CC α 16) predicts outcome in amyotrophic lateral sclerosis. <i>Acta Neurologica Scandinavica</i> , 2018, 137, 233-237.	2.1	4
54	Kinnier Wilson™s puzzling features of amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018, 89, 657-666.	1.9	4

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55	Retention of urine in women is alleviated by uterosacral ligament repair: implications for Fowler's syndrome. <i>Central European Journal of Urology</i> , 2018, 71, 436-443.	0.3	10
56	Concentric or monopolar electrode for jitter determination in orbicularis oculi. <i>Clinical Neurophysiology</i> , 2018, 129, 2552-2556.	1.5	4
57	Motor unit number estimation (MUNE): Where are we now?. <i>Clinical Neurophysiology</i> , 2018, 129, 1507-1516.	1.5	79
58	The covert recording of medico-legal consultations. <i>Medico-Legal Journal</i> , 2018, 86, 202-207.	0.5	3
59	MUNIX in the clinic in ALS: MUNE comes of age. <i>Clinical Neurophysiology</i> , 2017, 128, 482-483.	1.5	9
60	20th Anniversary Meeting of the Meryon Society Worcester College, Oxford. <i>Neuromuscular Disorders</i> , 2017, 27, 298-303.	0.6	0
61	New ideas on the ALS Functional Rating Scale. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017, 88, 371-372.	1.9	4
62	The generator site in acquired autoimmune neuromyotonia. <i>Clinical Neurophysiology</i> , 2017, 128, 643-646.	1.5	5
63	Fasciculation in amyotrophic lateral sclerosis: origin and pathophysiological relevance. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017, 88, 773-779.	1.9	76
64	Six issues in muscle disease. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017, 88, 603-607.	1.9	1
65	Pseudobulbar affective disorder, emotion and the brain. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2017, 88, 809-810.	1.9	2
66	Changing cortical inhibition in the course of amyotrophic lateral sclerosis. <i>Clinical Neurophysiology</i> , 2017, 128, 1032-1033.	1.5	0
67	Interplay of upper and lower motor neuron degeneration in amyotrophic lateral sclerosis. <i>Clinical Neurophysiology</i> , 2017, 128, 2200-2204.	1.5	4
68	Homozygous mutation in <i>HSPB1</i> causing distal vacuolar myopathy and motor neuropathy. <i>Neurology: Genetics</i> , 2017, 3, e168.	1.9	18
69	Risk factors for onset of amyotrophic lateral sclerosis. <i>European Journal of Neurology</i> , 2017, 24, 9-10.	3.3	1
70	Phrenic nerve stimulation is more sensitive than ultrasound measurement of diaphragm thickness in assessing early ALS progression. <i>Neurophysiologie Clinique</i> , 2017, 47, 69-73.	2.2	25
71	Physiology of the fasciculation potentials in amyotrophic lateral sclerosis: which motor units fasciculate?. <i>Journal of Physiological Sciences</i> , 2017, 67, 569-576.	2.1	12
72	Modulation of fasciculation frequency in amyotrophic lateral sclerosis: Table 1. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, jnnp-2014-309686.	1.9	11

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73	Lower motor neuron dysfunction in ALS. <i>Clinical Neurophysiology</i> , 2016, 127, 2670-2681.	1.5	62
74	Awaji criteria improves the diagnostic sensitivity in amyotrophic lateral sclerosis: A systematic review using individual patient data. <i>Clinical Neurophysiology</i> , 2016, 127, 2684-2691.	1.5	74
75	Dietary Factors and Amyotrophic Lateral Sclerosis. <i>JAMA Neurology</i> , 2016, 73, 1398.	9.0	1
76	Origin of fasciculations in root lesions. <i>Clinical Neurophysiology</i> , 2016, 127, 870-873.	1.5	5
77	Post-traumatic amnesia and confusional state: hazards of retrospective assessment. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, 1068-1074.	1.9	18
78	Fasciculation discharge frequency in amyotrophic lateral sclerosis and related disorders. <i>Clinical Neurophysiology</i> , 2016, 127, 2257-2262.	1.5	32
79	Ultrasound for assessment of diaphragm in ALS. <i>Clinical Neurophysiology</i> , 2016, 127, 892-897.	1.5	76
80	Brown-Vialetto-Van Laere syndrome: a 28-year follow-up. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2016, 87, 681-682.	1.9	10
81	Alcohol and amyotrophic lateral sclerosis: a possible neuroprotective effect. <i>European Journal of Neurology</i> , 2016, 23, 221-222.	3.3	2
82	Comment on: The Awaji criteria are not always superior to the previous criteria: A meta-analysis. <i>Muscle and Nerve</i> , 2015, 52, 467-468.	2.2	1
83	Measuring change in amyotrophic lateral sclerosis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 1169-1170.	1.9	3
84	John Hughlings Jackson (1835-1911): An adornment to the London Hospital. <i>Journal of Medical Biography</i> , 2015, 23, 2-8.	0.1	2
85	Clinical neurology: a changing role?. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 123-123.	1.9	0
86	The expanding syndrome of amyotrophic lateral sclerosis: a clinical and molecular odyssey. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2015, 86, 667-673.	1.9	104
87	Dissociated lower limb muscle involvement in amyotrophic lateral sclerosis. <i>Journal of Neurology</i> , 2015, 262, 1424-1432.	3.6	47
88	Motoneuron firing in amyotrophic lateral sclerosis (ALS). <i>Frontiers in Human Neuroscience</i> , 2014, 8, 719.	2.0	30
89	Theodore Leon "Ted" Munsat MD (1930-2013). <i>Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration</i> , 2014, 15, 473-474.	1.7	0
90	Diet and Risk of Amyotrophic Lateral Sclerosis. <i>JAMA Neurology</i> , 2014, 71, 1085.	9.0	4

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91	Does surgery accelerate progression of amyotrophic lateral sclerosis?. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 643-646.	1.9	48
92	Amyotrophic lateral sclerosis: a long preclinical period?. Journal of Neurology, Neurosurgery and Psychiatry, 2014, 85, 1232-1238.	1.9	120
93	Vascular endothelial growth factor and amyotrophic lateral sclerosis: The interplay with exercise and noninvasive ventilation. Muscle and Nerve, 2014, 49, 545-550.	2.2	12
94	Early diagnosis of amyotrophic lateral sclerosis – a way forward?. European Journal of Neurology, 2014, 21, 1435-1435.	3.3	1
95	Neurologic complications of craniocervical dislocation. Handbook of Clinical Neurology / Edited By PJ Vinken and G W Bruyn, 2014, 119, 435-448.	1.8	16
96	Delayed diagnosis in ALS: The problem continues. Journal of the Neurological Sciences, 2014, 343, 173-175.	0.6	63
97	Sensitivity of MUP parameters in detecting change in early ALS. Clinical Neurophysiology, 2014, 125, 166-169.	1.5	23
98	Congenital myopathy with focal loss of cross-striations revisited. Neuromuscular Disorders, 2013, 23, 160-164.	0.6	1
99	Fasciculation potentials and earliest changes in motor unit physiology in ALS. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 963-968.	1.9	93
100	Controversies and priorities in amyotrophic lateral sclerosis. Lancet Neurology, The, 2013, 12, 310-322.	10.2	454
101	Ventilation in ALS. European Journal of Neurology, 2013, 20, 1508-1509.	3.3	4
102	How does ALS spread between neurones in the CNS?. Journal of Neurology, Neurosurgery and Psychiatry, 2013, 84, 116-117.	1.9	10
103	Primary lateral sclerosis: Predicting functional outcome. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2013, 14, 141-145.	1.7	14
104	Origin of Fasciculations in Amyotrophic Lateral Sclerosis and Benign Fasciculation Syndrome. JAMA Neurology, 2013, 70, 1562-5.	9.0	32
105	Case: Failure to diagnose myocarditis leading to stroke and hemiplegia. Clinical Risk, 2012, 18, 33-35.	0.1	0
106	Why are upper motor neuron signs difficult to elicit in amyotrophic lateral sclerosis?: Figure 1. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 659-662.	1.9	108
107	Survey of non-invasive ventilation use in ALS in Britain. Journal of Neurology, Neurosurgery and Psychiatry, 2012, 83, 370-370.	1.9	3
108	Defecation 1: Testing a hypothesis for pelvic striated muscle action to open the anorectum. Techniques in Coloproctology, 2012, 16, 437-443.	1.8	22

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109	Defecation 2: Internal anorectal resistance is a critical factor in defecatory disorders. Techniques in Coloproctology, 2012, 16, 445-450.	1.8	21
110	Patrikios syndrome in two patients with treatable flail-leg weakness. Journal of Clinical Neuroscience, 2012, 19, 318-321.	1.5	0
111	Fasciculation potentials: Still mysterious. Clinical Neurophysiology, 2012, 123, 227-228.	1.5	6
112	Motor unit firing in amyotrophic lateral sclerosis and other upper and lower motor neurone disorders. Clinical Neurophysiology, 2012, 123, 2312-2318.	1.5	27
113	Awaji Criteria for the Diagnosis of Amyotrophic Lateral Sclerosis. Archives of Neurology, 2012, 69, 1410.	4.5	211
114	Respiratory exercise in amyotrophic lateral sclerosis. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2012, 13, 33-43.	2.1	56
115	Does the motor cortex influence denervation in ALS? EMG studies of muscles with both contralateral and bilateral corticospinal innervation. Clinical Neurophysiology, 2011, 122, 629-635.	1.5	15
116	Fasciculation-cramp syndrome preceding anterior horn cell disease: an intermediate syndrome?. Journal of Neurology, Neurosurgery and Psychiatry, 2011, 82, 459-461.	1.9	19
117	Amyotrophic lateral sclerosis. Current Opinion in Neurology, 2011, 24, 497-503.	3.6	44
118	Sacrocolpopexy may cause difficult defecation by inhibiting the external opening out mechanism. International Urogynecology Journal, 2011, 22, 255-255.	1.4	2
119	The Awaji criteria for diagnosis of ALS. Muscle and Nerve, 2011, 44, 456-456.	2.2	22
120	Nitric oxide and muscle weakness. Neurology, 2011, 76, 940-941.	1.1	3
121	Muscle ultrasound detects fasciculations and facilitates diagnosis in ALS. Neurology, 2011, 77, 1508-1509.	1.1	15
122	Love lies bleedingâ€”those who are left behind salute you. Neurology, 2011, 77, 1770-1770.	1.1	0
123	Sensitivity of electrophysiological tests for upper and lower motor neuron dysfunction in ALS: A sixâ€”month longitudinal study. Muscle and Nerve, 2010, 41, 208-211.	2.2	37
124	Lithium time-to-event trial in amyotrophic lateral sclerosis stops early for futility. Lancet Neurology, The, 2010, 9, 449-451.	10.2	7
125	Apraxia contributes to the motor deficit in Parkinsonâ€™s Disease and Multiple System Atrophy. European Journal of Neurology, 2010, 17, 346-347.	3.3	1
126	Internet facilitated management improves home ventilation in amyotrophic lateral sclerosis. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, 1180-1180.	1.9	1

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127	Meetings at the RSM. Journal of the Royal Society of Medicine, 2010, 103, 432-432.	2.0	0
128	Coping with motor neuron disease: how do people adapt to the devastating reality?. Journal of Neurology, Neurosurgery and Psychiatry, 2010, 81, 826-826.	1.9	2
129	Lockhart Clarke's contribution to the description of amyotrophic lateral sclerosis. Brain, 2010, 133, 3470-3479.	7.6	22
130	The onset of ALS?. Clinical Neurophysiology, 2010, 121, 1709-1710.	1.5	4
131	CSF markers in amyotrophic lateral sclerosis. Neurology, 2010, 74, 949-950.	1.1	6
132	Association of paraspinal and diaphragm denervation in ALS. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2010, 11, 63-66.	2.1	43
133	The Pelvic Floor: Functional Concepts and Neurocontrol. , 2010, , 33-42.		0
134	The Integral Theory: A Musculo-elastic Theory of Pelvic Floor Function and Dysfunction. , 2010, , 17-23.		0
135	What Is Really New in Progressive Muscle Atrophy?. Archives of Neurology, 2009, 66, 1427.	4.5	16
136	Clinical questions need perceptive answers. Practical Neurology, 2009, 9, 117-117.	1.1	0
137	Money and medicine. Neurology, 2009, 72, 766-768.	1.1	2
138	Peer review and "openness"™. Journal of the Royal Society of Medicine, 2009, 102, 507-508.	2.0	0
139	Motor unit changes in thoracic paraspinal muscles in amyotrophic lateral sclerosis. Muscle and Nerve, 2009, 39, 83-86.	2.2	15
140	Stratifying disease stages with different progression rates determined by electrophysiological tests in patients with amyotrophic lateral sclerosis. Muscle and Nerve, 2009, 40, 318-318.	2.2	1
141	Late-onset axial myopathy with cores due to a novel heterozygous dominant mutation in the skeletal muscle ryanodine receptor (RYR1) gene. Neuromuscular Disorders, 2009, 19, 344-347.	0.6	103
142	Predicting respiratory insufficiency in amyotrophic lateral sclerosis: The role of phrenic nerve studies. Clinical Neurophysiology, 2009, 120, 941-946.	1.5	72
143	An error of self-diagnosis--but what was the real diagnosis?. Practical Neurology, 2009, 9, 284-288.	1.1	1
144	Awaji diagnostic algorithm increases sensitivity of El Escorial criteria for ALS diagnosis. Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders, 2009, 10, 53-57.	2.1	196

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145	Paraspinal and limb motor neuron involvement within homologous spinal segments in ALS. <i>Clinical Neurophysiology</i> , 2008, 119, 1607-1613.	1.5	31
146	The innervation of muscle and the neuron theory. <i>Neuromuscular Disorders</i> , 2008, 18, 426-430.	0.6	2
147	Cervical Cord Compression in Mucopolysaccharidosis. <i>Developmental Medicine and Child Neurology</i> , 2008, 15, 194-199.	2.1	23
148	Electrodiagnostic criteria for diagnosis of ALS. <i>Clinical Neurophysiology</i> , 2008, 119, 497-503.	1.5	927
149	WHAT IS NEXT IN ALS CLINICAL TRIALS?. <i>Neurology</i> , 2008, 70, 1365-1366.	1.1	1
150	WHAT IS NEXT IN ALS CLINICAL TRIALS?. <i>Neurology</i> , 2008, 70, 1366-1367.	1.1	1
151	Henry Head and the development of clinical neuroscience. <i>Brain</i> , 2008, 131, 3453-3456.	7.6	4
152	Sphincter Disorders and the Nervous System. , 2008, , 633-650.		3
153	Comparison of the 40-item Amyotrophic Lateral Sclerosis Assessment Questionnaire (ALSAQ-40) with a short-form five-item version (ALSAQ-5) in a longitudinal survey. <i>Clinical Rehabilitation</i> , 2007, 21, 266-272.	2.2	14
154	Homozygous mutation in MYH7 in myosin storage myopathy and cardiomyopathy. <i>Neurology</i> , 2007, 68, 962-962.	1.1	56
155	We have a problem: Why have ALS trials been negative?. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2007, 8, 259-259.	2.1	7
156	Molecular mechanisms and phenotypic variation in RYR1-related congenital myopathies. <i>Brain</i> , 2007, 130, 2024-2036.	7.6	161
157	LIMB-KINETIC APRAXIA IN PARKINSON DISEASE. <i>Neurology</i> , 2007, 69, 810-811.	1.1	2
158	Phase II/III randomized trial of TCH346 in patients with ALS. <i>Neurology</i> , 2007, 69, 776-784.	1.1	112
159	And Lord Brain said <i>Practical Neurology</i> , 2007, 7, 250-251.	1.1	0
160	Myosin storage myopathy with cardiomyopathy. <i>Neuromuscular Disorders</i> , 2007, 17, 725.	0.6	12
161	Monomelic neurogenic syndromes: A prospective study. <i>Journal of the Neurological Sciences</i> , 2007, 263, 26-34.	0.6	21
162	Clinical patterns in progressive muscular atrophy (PMA): A prospective study. <i>Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders</i> , 2007, 8, 296-299.	2.1	28

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163	The phagocytic capacity of neurones. <i>European Journal of Neuroscience</i> , 2007, 25, 2947-2955.	2.6	41
164	Learning from failed trials in ALS. <i>Lancet Neurology</i> , The, 2007, 6, 1034-1035.	10.2	13
165	Hereditary Spastic Paraplegia and Primary Lateral Sclerosis. , 2007, , 537-544.		2
166	Hemicrania Continua. <i>Cephalalgia</i> , 2006, 26, 341-344.	3.9	41
167	Can Selection of Rapidly Progressing Patients Shorten Clinical Trials in Amyotrophic Lateral Sclerosis?. <i>Archives of Neurology</i> , 2006, 63, 557.	4.5	32
168	Hughlings Jackson's clinical research: Evidence from contemporary documents. <i>Neurology</i> , 2006, 67, 666-672.	1.1	2
169	Multifocal motor neuropathy. <i>Neurology</i> , 2006, 67, 558-559.	1.1	27
170	Rectal Hyposensitivity. <i>American Journal of Gastroenterology</i> , 2006, 101, 1140-1151.	0.4	137
171	The anal reflex in idiopathic faecal incontinence; an electrophysiological study. <i>British Journal of Surgery</i> , 2005, 67, 781-783.	0.3	56
172	Physiological studies of the anal sphincter musculature in faecal incontinence and rectal prolapse. <i>British Journal of Surgery</i> , 2005, 68, 531-536.	0.3	310
173	The pelvic floor musculature in the descending perineum syndrome. <i>British Journal of Surgery</i> , 2005, 69, 470-472.	0.3	291
174	Slowed conduction in the pudendal nerves in idiopathic (neurogenic) faecal incontinence. <i>British Journal of Surgery</i> , 2005, 71, 614-616.	0.3	444
175	Faecal incontinence after anal dilatation. <i>British Journal of Surgery</i> , 2005, 71, 617-618.	0.3	81
176	Electrophysiological and manometric assessment of the pelvic floor in the solitary rectal ulcer syndrome. <i>British Journal of Surgery</i> , 2005, 72, 131-133.	0.3	78
177	Anorectal incontinence: Electrophysiological tests. <i>British Journal of Surgery</i> , 2005, 72, s14-s15.	0.3	20
178	Risk factors in childbirth causing damage to the pelvic floor innervation. <i>British Journal of Surgery</i> , 2005, 72, s15-s17.	0.3	77
179	Neural control of internal anal sphincter function. <i>British Journal of Surgery</i> , 2005, 74, 668-670.	0.3	121
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