Baziel van Engelen

List of Publications by Year
in descending order

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508 papers	21,037 citations	9254 74 h-index	19169 118 g-index
527 all docs	527 docs citations	527 times ranked	17009 citing authors

#	Article	IF	Citations
1	Development and validation of the patient-reported â€œFacial Function Scaleâ€•for facioscapulohumeral muscular dystrophy. Disability and Rehabilitation, 2023, 45, 1530-1535.	0.9	2
2	Chromosome 10q-linked FSHD identifies DUX4 as principal disease gene. Journal of Medical Genetics, 2022, 59, 180-188.	1.5	18
3	Facioscapulohumeral muscular dystrophyâ€”Reproductive counseling, pregnancy, and delivery in a complex multigenetic disease. Clinical Genetics, 2022, 101, 149-160.	1.0	5
4	High-resolution breakpoint junction mapping of proximally extended D4Z4 deletions in FSHD1 reveals evidence for a founder effect. Human Molecular Genetics, 2022, 31, 748-760.	1.4	8
5	Experiences of patients with facioscapulohumeral dystrophy with facial weakness: a qualitative study. Disability and Rehabilitation, 2022, 44, 6775-6782.	0.9	3
6	N-of-1 Trials in Neurology. Neurology, 2022, 98, .	1.5	7
7	Facioscapulohumeral dystrophy transcriptome signatures correlate with different stages of disease and are marked by different MRI biomarkers. Scientific Reports, 2022, 12, 1426.	1.6	14
8	Long-term follow-up of respiratory function in facioscapulohumeral muscular dystrophy. Journal of Neurology, 2022, 269, 3682-3689.	1.8	2
9	Visuomotor processing is altered after peripheral nerve damage in neuralgic amyotrophy. Brain Communications, 2022, 4, fcac034.	1.5	2
10	Neurological features of Noonan syndrome and related <scp>RASopathies</scp> : Pain and nerve enlargement characterized by nerve ultrasound. American Journal of Medical Genetics, Part A, 2022, , .	0.7	3
11	Blood Transcriptome Profiling Links Immunity to Disease Severity in Myotonic Dystrophy Type 1 (DM1). International Journal of Molecular Sciences, 2022, 23, 3081.	1.8	3
12	Quantitative Muscle Analysis in Facioscapulohumeral Muscular Dystrophy Using <scp>Wholeâ€Body Fatâ€Referenced MRI</scp> : Protocol Development, Multicenter Feasibility, and Repeatability. Muscle and Nerve, 2022, , .	1.0	1
13	Reachable workspace analysis is a potential measurement for impairment of the upper extremity in neuralgic amyotrophy. Muscle and Nerve, 2022, 66, 282-288.	1.0	2
14	Respiratory muscle function in patients with nemaline myopathy. Neuromuscular Disorders, 2022, 32, 654-663.	0.3	7
15	Electrocardiographic predictors of infrahissian conduction disturbances in myotonic dystrophy type 1. Europace, 2021, 23, 298-304.	0.7	18
16	Systemic cell therapy for muscular dystrophies. Stem Cell Reviews and Reports, 2021, 17, 878-899.	1.7	11
17	Characterizing the face in facioscapulohumeral muscular dystrophy. Journal of Neurology, 2021, 268, 1342-1350.	1.8	13
18	Reduced specific force in patients with mild and severe facioscapulohumeral muscular dystrophy. Muscle and Nerve, 2021, 63, 60-67.	1.0	9

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19	Semiâ€automated Rasch analysis using inâ€plusâ€outâ€ofâ€questionnaire log likelihood. British Journal of Mathematical and Statistical Psychology, 2021, 74, 313-339.	1.0	3
20	New Insights in Adherence and Survival in Myotonic Dystrophy Patients Using Home Mechanical Ventilation. Respiration, 2021, 100, 154-163.	1.2	6
21	Noninvasive Home Mechanical Ventilation in Adult Myotonic Dystrophy Type 1: A Systematic Review. Respiration, 2021, 100, 816-825.	1.2	5
22	Human brain pathology in myotonic dystrophy type 1: A systematic review. Neuropathology, 2021, 41, 3-20.	0.7	21
23	NEM6, KBTBD13-Related Congenital Myopathy: Myopathological Analysis in 18 Dutch Patients Reveals Ring Rods Fibers, Cores, Nuclear Clumps, and Granulo-Filamentous Protein Material. Journal of Neuropathology and Experimental Neurology, 2021, 80, 366-376.	0.9	4
24	Associations Between Variant Repeat Interruptions and Clinical Outcomes in Myotonic Dystrophy Type 1. Neurology: Genetics, 2021, 7, e572.	0.9	10
25	Respiratory muscle imaging by ultrasound and MRI in neuromuscular disorders. European Respiratory Journal, 2021, 58, 2100137.	3.1	5
26	Second intravenous immunoglobulin dose in patients with Guillain-BarrÃ© syndrome with poor prognosis (SID-GBS): a double-blind, randomised, placebo-controlled trial. Lancet Neurology, The, 2021, 20, 275-283.	4.9	34
27	The socioeconomic burden of facioscapulohumeral muscular dystrophy. Journal of Neurology, 2021, 268, 4778-4788.	1.8	5
28	Antiâ€“Cytosolic 5â€²â€Nucleotidase 1A Autoantibodies Are Absent in Juvenile Dermatomyositis. Arthritis and Rheumatology, 2021, 73, 1329-1333.	2.9	2
29	Phase 1 clinical trial of losmapimod in facioscapulohumeral dystrophy: Safety, tolerability, pharmacokinetics, and target engagement. British Journal of Clinical Pharmacology, 2021, 87, 4658-4669.	1.1	20
30	The facioscapulohumeral muscular dystrophy Raschâ€built overall disability scale (FSHDâ€RODS). European Journal of Neurology, 2021, 28, 2339-2348.	1.7	8
31	N-of-1 trial of salbutamol in hyperkalaemic periodic paralysis. Journal of Neurology, Neurosurgery and Psychiatry, 2021, 92, jnnp-2021-326347.	0.9	1
32	Behavioural Impairment and Frontotemporal Dementia in Oculopharyngeal Muscular Dystrophy. Journal of Neuromuscular Diseases, 2021, , 1-7.	1.1	1
33	Clinical Outcome Evaluations and CBT Response Prediction in Myotonic Dystrophy. Journal of Neuromuscular Diseases, 2021, 8, 1031-1046.	1.1	4
34	Exploring the influence of smoking and alcohol consumption on clinical severity in patients with facioscapulohumeral muscular dystrophy. Neuromuscular Disorders, 2021, 31, 824-828.	0.3	1
35	Longitudinal Assessment of Strength, Functional Capacity, Oropharyngeal Function, and Quality of Life in Oculopharyngeal Muscular Dystrophy. Neurology, 2021, 97, e1475-e1483.	1.5	11
36	Mixed methods evaluation of a self-management group programme for patients with neuromuscular disease and chronic fatigue. BMJ Open, 2021, 11, e048890.	0.8	3

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37	The neuromuscular and multisystem features of RYR1-related malignant hyperthermia and rhabdomyolysis. Medicine (United States), 2021, 100, e26999.	0.4	8
38	Natural history, outcome measures and trial readiness in LAMA2-related muscular dystrophy and SELENON-related myopathy in children and adults: protocol of the LAST STRONG study. BMC Neurology, 2021, 21, 313.	0.8	12
39	Clinical, genetic, and histological features of centronuclear myopathy in the Netherlands. Clinical Genetics, 2021, 100, 692-702.	1.0	7
40	Profiling Serum Antibodies Against Muscle Antigens in Facioscapulohumeral Muscular Dystrophy Finds No Disease-Specific Autoantibodies. Journal of Neuromuscular Diseases, 2021, 8, 801-814.	1.1	6
41	MYOTONIC DYSTROPHY. Neuromuscular Disorders, 2021, 31, S121.	0.3	0
42	FSHD. Neuromuscular Disorders, 2021, 31, S99-S100.	0.3	0
43	FSHD. Neuromuscular Disorders, 2021, 31, S100.	0.3	0
44	CONGENITAL MUSCULAR DYSTROPHIES. Neuromuscular Disorders, 2021, 31, S70.	0.3	0
45	CONGENITAL MYOPATHIES. Neuromuscular Disorders, 2021, 31, S65.	0.3	0
46	Rasch analysis to evaluate the motor function measure for patients with facioscapulohumeral muscular dystrophy. International Journal of Rehabilitation Research, 2021, 44, 38-44.	0.7	13
47	Natural History of Facioscapulohumeral Dystrophy in Children. Neurology, 2021, 97, e2103-e2113.	1.5	7
48	Care for capabilities: Implementing the capability approach in rehabilitation of patients with neuromuscular diseases. Study protocol of the controlled before-after ReCap-NMD study. PLoS ONE, 2021, 16, e0261475.	1.1	1
49	Short fatigue questionnaire: Screening for severe fatigue.. Journal of Psychosomatic Research, 2020, 137, 110229.	1.2	17
50	Swallowing, Chewing and Speaking: Frequently Impaired in Oculopharyngeal Muscular Dystrophy. Journal of Neuromuscular Diseases, 2020, 7, 483-494.	1.1	8
51	CONGENITAL MYOPATHIES 1 â€“ NEMALINE. Neuromuscular Disorders, 2020, 30, S53.	0.3	0
52	CONGENITAL MYOPATHIES 1 â€“ NEMALINE. Neuromuscular Disorders, 2020, 30, S55.	0.3	0
53	FSHD / OPMD / MYOTONIC DYSTROPHY. Neuromuscular Disorders, 2020, 30, S112-S113.	0.3	0
54	FSHD / OPMD / MYOTONIC DYSTROPHY. Neuromuscular Disorders, 2020, 30, S113.	0.3	0

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55	FSHD / OPMD / MYOTONIC DYSTROPHY. Neuromuscular Disorders, 2020, 30, S113.	0.3	0
56	Deep phenotyping of facioscapulohumeral muscular dystrophy type 2 by magnetic resonance imaging. European Journal of Neurology, 2020, 27, 2604-2615.	1.7	16
57	Clinical, morphological and genetic characterization of Brody disease: an international study of 40 patients. Brain, 2020, 143, 452-466.	3.7	22
58	Age-Associated Salivary MicroRNA Biomarkers for Oculopharyngeal Muscular Dystrophy. International Journal of Molecular Sciences, 2020, 21, 6059.	1.8	9
59	Correlation Between Quantitative MRI and Muscle Histopathology in Muscle Biopsies from Healthy Controls and Patients with IBM, FSHD and OPMD. Journal of Neuromuscular Diseases, 2020, 7, 495-504.	1.1	25
60	Swallowing, Chewing and Speaking: Frequently Impaired in Oculopharyngeal Muscular Dystrophy. Journal of Neuromuscular Diseases, 2020, 7, 1-12.	1.1	12
61	Quantitative Muscle MRI Depicts Increased Muscle Mass after a Behavioral Change in Myotonic Dystrophy Type 1. Radiology, 2020, 297, 132-142.	3.6	11
62	Inclusion body myositis in patients with spinocerebellar ataxia types 3 and 6. Journal of Neurology, Neurosurgery and Psychiatry, 2020, 91, 876-878.	0.9	2
63	Altered sensorimotor representations after recovery from peripheral nerve damage in neuralgic amyotrophy. Cortex, 2020, 127, 180-190.	1.1	10
64	Muscle ultrasound is a responsive biomarker in facioscapulohumeral dystrophy. Neurology, 2020, 94, e1488-e1494.	1.5	23
65	N-of-1 Trials: Evidence-Based Clinical Care or Medical Research that Requires IRB Approval? A Practical Flowchart Based on an Ethical Framework. Healthcare (Switzerland), 2020, 8, 49.	1.0	18
66	Characterization of EEG-based functional brain networks in myotonic dystrophy type 1. Clinical Neurophysiology, 2020, 131, 1886-1895.	0.7	1
67	Consequences of epigenetic derepression in facioscapulohumeral muscular dystrophy. Clinical Genetics, 2020, 97, 799-814.	1.0	40
68	Preserved single muscle fiber specific force in facioscapulohumeral muscular dystrophy. Neurology, 2020, 94, e1157-e1170.	1.5	8
69	Continued misuse of orphan drug legislation: a life-threatening risk for mexiletine. European Heart Journal, 2020, 41, 614-617.	1.0	15
70	Guidelines on clinical presentation and management of nondystrophic myotonias. Muscle and Nerve, 2020, 62, 430-444.	1.0	53
71	248th ENMC International Workshop: Myotonic dystrophies: Molecular approaches for clinical purposes, framing a European molecular research network, Hoofddorp, the Netherlands, 11â€“13 October 2019. Neuromuscular Disorders, 2020, 30, 521-531.	0.3	1
72	KBTBD13 is an actin-binding protein that modulates muscle kinetics. Journal of Clinical Investigation, 2020, 130, 754-767.	3.9	25

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73	NA-CONTROL: a study protocol for a randomised controlled trial to compare specific outpatient rehabilitation that targets cerebral mechanisms through relearning motor control and uses self-management strategies to improve functional capability of the upper extremity, to usual care in patients with neuralgic amyotrophy. Trials, 2019, 20, 482.	0.7	9
74	SMCHD1 mutation spectrum for facioscapulohumeral muscular dystrophy type 2 (FSHD2) and Bosma arhinia microphthalmia syndrome (BAMS) reveals disease-specific localisation of variants in the ATPase domain. Journal of Medical Genetics, 2019, 56, 693-700.	1.5	27
75	Ophthalmological findings in facioscapulohumeral dystrophy. Brain Communications, 2019, 1, fcz023.	1.5	14
76	O.25Phase 1 clinical trial of losmapimod in FSHD: safety, tolerability and target engagement. Neuromuscular Disorders, 2019, 29, S123.	0.3	1
77	P.40Ophthalmological findings in facioscapulohumeral dystrophy. Neuromuscular Disorders, 2019, 29, S52-S53.	0.3	0
78	P.46Clinical trial readiness to solve barriers to drug development in FSHD (ReSolve): protocol of a large, multi-center prospective study. Neuromuscular Disorders, 2019, 29, S54.	0.3	0
79	P.63Chronic progressive external ophthalmoplegia (CPEO) and CPEO-plus cohort of 54 patients from the Netherlands. Neuromuscular Disorders, 2019, 29, S59.	0.3	0
80	P.162Novel Kbtbd13R408C-knockin mouse model phenocopies NEM6 myopathy. Neuromuscular Disorders, 2019, 29, S95.	0.3	0
81	O.13Nemaline myopathy patients with mutations in KBTBD13 display a cardiac phenotype. Neuromuscular Disorders, 2019, 29, S118.	0.3	0
82	P.247Capturing disease progression in oculopharyngeal muscular dystrophy (OPMD). Neuromuscular Disorders, 2019, 29, S139.	0.3	0
83	P.306Multicentric MRI study in a cohort of FSHD2 patients: pattern definition and differences between FSHD1 and FSHD2. Neuromuscular Disorders, 2019, 29, S156.	0.3	0
84	High incidence of falls in patients with myotonic dystrophy type 1 and 2: A prospective study. Neuromuscular Disorders, 2019, 29, 758-765.	0.3	10
85	Scapular dyskinesis in myotonic dystrophy type 1: clinical characteristics and genetic investigations. Journal of Neurology, 2019, 266, 2987-2996.	1.8	1
86	Clinical trial readiness to solve barriers to drug development in FSHD (ReSolve): protocol of a large, international, multi-center prospective study. BMC Neurology, 2019, 19, 224.	0.8	28
87	Effects of weakness of orofacial muscles on swallowing and communication in FSHD. Neurology, 2019, 92, e957-e963.	1.5	25
88	Lower extremity muscle pathology in myotonic dystrophy type 1 assessed by quantitative MRI. Neurology, 2019, 92, e2803-e2814.	1.5	34
89	MSH3 modifies somatic instability and disease severity in Huntingtonâ€™s and myotonic dystrophy type 1. Brain, 2019, 142, 1876-1886.	3.7	114
90	Panel-Based Exome Sequencing for Neuromuscular Disorders as a Diagnostic Service. Journal of Neuromuscular Diseases, 2019, 6, 241-258.	1.1	32

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91	Muscle fiber dysfunction contributes to weakness in inclusion body myositis. Neuromuscular Disorders, 2019, 29, 468-476.	0.3	3
92	Autoantibody testing in idiopathic inflammatory myopathies. Practical Neurology, 2019, 19, 284-294.	0.5	16
93	Affective symptoms and apathy in myotonic dystrophy type 1 a systematic review and meta-analysis. Journal of Affective Disorders, 2019, 250, 260-269.	2.0	23
94	Health-Related Quality of Life in Patients with Adult-Onset Myotonic Dystrophy Type 1: A Systematic Review. Patient, 2019, 12, 365-373.	1.1	11
95	The Position of Neuromuscular Patients in Shared Decision Making. Report from the 235th ENMC Workshop: Milan, Italy, January 19-20, 2018. Journal of Neuromuscular Diseases, 2019, 6, 161-172.	1.1	7
96	Genetic determinants of disease severity in the myotonic dystrophy type 1 OPTIMISTIC cohort. Neurology, 2019, 93, e995-e1009.	1.5	71
97	Self-management program improves participation in patients with neuromuscular disease. Neurology, 2019, 93, e1720-e1731.	1.5	23
98	Insulin Signaling as a Key Moderator in Myotonic Dystrophy Type 1. Frontiers in Neurology, 2019, 10, 1229.	1.1	17
99	Strength training and aerobic exercise training for muscle disease. The Cochrane Library, 2019, 2019, CD003907.	1.5	44
100	Early onset as a marker for disease severity in facioscapulohumeral muscular dystrophy. Neurology, 2019, 92, e378-e385.	1.5	30
101	Structural white matter networks in myotonic dystrophy type 1. NeuroImage: Clinical, 2019, 21, 101615.	1.4	23
102	Single-cell RNA sequencing in facioscapulohumeral muscular dystrophy disease etiology and development. Human Molecular Genetics, 2019, 28, 1064-1075.	1.4	46
103	Functional impairments, fatigue and quality of life in RYR1-related myopathies: A questionnaire study. Neuromuscular Disorders, 2019, 29, 30-38.	0.3	20
104	Diagnostics of short tandem repeat expansion variants using massively parallel sequencing and componential tools. European Journal of Human Genetics, 2019, 27, 400-407.	1.4	12
105	Muscle MRI in a large cohort of patients with oculopharyngeal muscular dystrophy. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 576-585.	0.9	38
106	Reference values of maximum performance tests of speech production. International Journal of Speech-Language Pathology, 2019, 21, 56-64.	0.6	29
107	Ultrasound: A Potential Tool for Detecting of Fasciitis in Dermatomyositis and Polymyositis. Journal of Rheumatology, 2018, 45, 441.1-442.	1.0	5
108	Lifetime endogenous estrogen exposure and disease severity in female patients with facioscapulohumeral muscular dystrophy. Neuromuscular Disorders, 2018, 28, 508-511.	0.3	21

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109	Hearing impairment in patients with myotonic dystrophy type 2. Neurology, 2018, 90, e615-e622.	1.5	11
110	Falls and resulting fractures in Myotonic Dystrophy: Results from a multinational retrospective survey. Neuromuscular Disorders, 2018, 28, 229-235.	0.3	19
111	Electrical impedance myography in facioscapulohumeral muscular dystrophy: A 1â€year followâ€up study. Muscle and Nerve, 2018, 58, 213-218.	1.0	15
112	Qualitative and Quantitative Aspects of Pain in Patients With Myotonic Dystrophy Type 2. Journal of Pain, 2018, 19, 920-930.	0.7	19
113	Prevalence and mutation spectrum of skeletal muscle channelopathies in the Netherlands. Neuromuscular Disorders, 2018, 28, 402-407.	0.3	40
114	MRI in sarcoglycanopathies: a large international cohort study. Journal of Neurology, Neurosurgery and Psychiatry, 2018, 89, 72-77.	0.9	55
115	Deep characterization of a common D4Z4 variant identifies biallelic DUX4 expression as a modifier for disease penetrance in FSHD2. European Journal of Human Genetics, 2018, 26, 94-106.	1.4	22
116	Specific muscle strength is reduced in facioscapulohumeral dystrophy: An MRI based musculoskeletal analysis. Neuromuscular Disorders, 2018, 28, 238-245.	0.3	11
117	Evidence of ER stress and UPR activation in patients with Brody disease and Brody syndrome. Neuropathology and Applied Neurobiology, 2018, 44, 533-536.	1.8	1
118	CONGENITAL MYOPATHIES: NEMALINE AND TITINOPATHIES. Neuromuscular Disorders, 2018, 28, S100-S101.	0.3	0
119	Effect of Mexiletine on Muscle Stiffness in Patients With Nondystrophic Myotonia Evaluated Using Aggregated N-of-1 Trials. JAMA - Journal of the American Medical Association, 2018, 320, 2344.	3.8	81
120	Quantitative muscle MRI and ultrasound for facioscapulohumeral muscular dystrophy: complementary imaging biomarkers. Journal of Neurology, 2018, 265, 2646-2655.	1.8	54
121	Consensus-based care recommendations for adults with myotonic dystrophy type 1. Neurology: Clinical Practice, 2018, 8, 507-520.	0.8	115
122	Facioscapulohumeral Dystrophy in Childhood: A Nationwide Natural History Study. Annals of Neurology, 2018, 84, 627-637.	2.8	21
123	NEW GENES, FUNCTIONS AND BIOMARKERS. Neuromuscular Disorders, 2018, 28, S31.	0.3	0
124	Phenotypeâ€genotype relations in facioscapulohumeral muscular dystrophy type 1. Clinical Genetics, 2018, 94, 521-527.	1.0	25
125	Eight years after an international workshop on myotonic dystrophy patient registries: case study of a global collaboration for a rare disease. Orphanet Journal of Rare Diseases, 2018, 13, 155.	1.2	19
126	Second IVIg course in Guillainâ€BarrÃ© syndrome patients with poor prognosis (SIDâ€GBS trial): Protocol for a doubleâ€blind randomized, placeboâ€controlled clinical trial. Journal of the Peripheral Nervous System, 2018, 23, 210-215.	1.4	36

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127	A 22-year follow-up reveals a variable disease severity in early-onset facioscapulohumeral dystrophy. European Journal of Paediatric Neurology, 2018, 22, 782-785.	0.7	8
128	Cis D4Z4 repeat duplications associated with facioscapulohumeral muscular dystrophy type 2. Human Molecular Genetics, 2018, 27, 3488-3497.	1.4	27
129	Autoantibodies to Cytosolic 5â€²-Nucleotidase 1A in Primary SjÃ¶grenâ€™s Syndrome and Systemic Lupus Erythematosus. Frontiers in Immunology, 2018, 9, 1200.	2.2	32
130	A family-based study into penetrance in facioscapulohumeral muscular dystrophy type 1. Neurology, 2018, 91, e444-e454.	1.5	33
131	FSHD type 2 and Bosma arhinia microphthalmia syndrome. Neurology, 2018, 91, e562-e570.	1.5	24
132	MRI-Guided Biopsy as a Tool for Diagnosis and Research of Muscle Disorders. Journal of Neuromuscular Diseases, 2018, 5, 315-319.	1.1	24
133	Repeatability and reliability of muscle relaxation properties induced by motor cortical stimulation. Journal of Applied Physiology, 2018, 124, 1597-1604.	1.2	5
134	Fatigue, not self-rated motor symptom severity, affects quality of life in functional motor disorders. Journal of Neurology, 2018, 265, 1803-1809.	1.8	48
135	Cognitive behavioural therapy with optional graded exercise therapy in patients with severe fatigue with myotonic dystrophy type 1: a multicentre, single-blind, randomised trial. Lancet Neurology, The, 2018, 17, 671-680.	4.9	95
136	Is fatigue a disease-specific or generic symptom in chronic medical conditions?. Health Psychology, 2018, 37, 530-543.	1.3	79
137	AB0776â€…Muscle ultrasonography: a potential new diagnostic tool for inflammatory myopathies. , 2018, , .		0
138	PGM1 deficiency: Substrate use during exercise and effect of treatment with galactose. Neuromuscular Disorders, 2017, 27, 370-376.	0.3	31
139	Cytosolic 5â€²-nucleotidase 1A autoantibody profile and clinical characteristics in inclusion body myositis. Annals of the Rheumatic Diseases, 2017, 76, 862-868.	0.5	71
140	225th ENMC international workshop:. Neuromuscular Disorders, 2017, 27, 782-790.	0.3	20
141	The assessment of fatigue: Psychometric qualities and norms for the Checklist individual strength. Journal of Psychosomatic Research, 2017, 98, 40-46.	1.2	222
142	Respiratory function in facioscapulohumeral muscular dystrophy 1. Neuromuscular Disorders, 2017, 27, 526-530.	0.3	14
143	Respiratory pattern in a FSHD pediatric population. Respiratory Medicine, 2017, 126, 130-131.	1.3	0
144	Oculopharyngeal muscular dystrophy with frontotemporal dementia. European Geriatric Medicine, 2017, 8, 81-83.	1.2	3

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145	The yield of diagnostic work-up of patients presenting with myalgia, exercise intolerance, or fatigue. Neuromuscular Disorders, 2017, 27, 243-250.	0.3	10
146	Involvement of pelvic girdle and proximal leg muscles in early oculopharyngeal muscular dystrophy. Neuromuscular Disorders, 2017, 27, 1099-1105.	0.3	17
147	Muscle fiber dysfunction contributes to clinical muscle weakness in inclusion body myositis. Neuromuscular Disorders, 2017, 27, S154.	0.3	0
148	The Brody disease cohort study: clarification of the phenotype. Neuromuscular Disorders, 2017, 27, S164.	0.3	0
149	Adding quantitative muscle MRI to the FSHD clinical trial toolbox. Neuromuscular Disorders, 2017, 27, S122.	0.3	1
150	Specific strength is reduced in facioscapulohumeral dystrophy muscles. An MRI-based musculoskeletal analysis. Neuromuscular Disorders, 2017, 27, S200.	0.3	0
151	Adding quantitative muscle MRI to the FSHD clinical trial toolbox. Neurology, 2017, 89, 2057-2065.	1.5	72
152	Muscle ultrasonography is a potential tool for detecting fasciitis in dermatomyositis and polymyositis: comment on the article by Yoshida etÂal. Arthritis and Rheumatology, 2017, 69, 2248-2249.	2.9	4
153	Clinical phenotype and outcome of hepatitis E virus associated neuralgic amyotrophy; an international retrospective comparative cohort study. Journal of Hepatology, 2017, 66, S59.	1.8	2
154	Hepatitis E virus infection and acute non-traumatic neurological injury: A prospective multicentre study. Journal of Hepatology, 2017, 67, 925-932.	1.8	80
155	Clinical phenotype and outcome of hepatitis E virusâ€“associated neuralgic amyotrophy. Neurology, 2017, 89, 909-917.	1.5	75
156	Brain imaging in myotonic dystrophy type 1. Neurology, 2017, 89, 960-969.	1.5	76
157	The cognitive profile of myotonic dystrophy type 1:ÂA systematic review and meta-analysis. Cortex, 2017, 95, 143-155.	1.1	82
158	Immunemediated necrotizing autoimmune myopathy: Dutch and Belgian experience. Neuromuscular Disorders, 2017, 27, S151.	0.3	0
159	Muscular fat fraction correlates with functionality in myotonic dystrophy type 1. Neuromuscular Disorders, 2017, 27, S180.	0.3	0
160	Retinal abnormalities in FSHD. Neuromuscular Disorders, 2017, 27, S201.	0.3	0
161	Slow relaxation kinetics of sarcomeres contribute to muscle slowness in NEM6 patients. Neuromuscular Disorders, 2017, 27, S230.	0.3	0
162	Early onset facioscapulohumeral dystrophy â€“ a systematic review using individual patient data. Neuromuscular Disorders, 2017, 27, 1077-1083.	0.3	39

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163	Effects of Mindfulness-Based Stress Reduction on the Mental Health of Clinical Clerkship Students: A Cluster-Randomized Controlled Trial. Academic Medicine, 2017, 92, 1012-1021.	0.8	56
164	Recognizing the tenascinâ€X deficient type of Ehlersâ€“Danlos syndrome: a crossâ€sectional study in 17 patients. Clinical Genetics, 2017, 91, 411-425.	1.0	46
165	The Radboud Dysarthria Assessment: Development and Clinimetric Evaluation. Folia Phoniatrica Et Logopaedica, 2017, 69, 143-153.	0.5	35
166	Characteristics and natural history of oculopharyngeal muscular dystrophy (OPMD): The study protocol of â€˜OPMD Forteâ€™. Neuromuscular Disorders, 2016, 26, S139-S140.	0.3	3
167	The wrong end of the telescope: neuromuscular mimics of movement disorders (and vice versa). Practical Neurology, 2016, 16, 264-269.	0.5	11
168	The energetic study: Effectiveness of a self-management group programme to improve social participation in patients with neuromuscular disease and chronic fatigue. Neuromuscular Disorders, 2016, 26, S147.	0.3	0
169	Contractile dysfunction in permeabilized muscle fibers of NEM6 patients with the Dutch founder mutation in KBTBD13. Neuromuscular Disorders, 2016, 26, S134.	0.3	0
170	Monitoring creatine and phosphocreatine by 13C MR spectroscopic imaging during and after 13C4 creatine loading: a feasibility study. Amino Acids, 2016, 48, 1857-1866.	1.2	4
171	Integrating clinical and genetic observations in facioscapulohumeral muscular dystrophy. Current Opinion in Neurology, 2016, 29, 606-613.	1.8	10
172	The epidemiology of neuromuscular disorders: Age at onset and gender in the Netherlands. Neuromuscular Disorders, 2016, 26, 447-452.	0.3	17
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174	Quantitative MRI reveals decelerated fatty infiltration in muscles of active FSHD patients. Neurology, 2016, 86, 1700-1707.	1.5	61
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