

Baziel van Engelen

List of Publications by Year in descending order

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Version: 2024-02-01

508
papers

21,037
citations

10650

74
h-index

21843

118
g-index

527
all docs

527
docs citations

527
times ranked

18162
citing authors

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

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19	Semi-automated Rasch analysis using inâ€plusâ€outâ€ofâ€questionnaire log likelihood. <i>British Journal of Mathematical and Statistical Psychology</i> , 2021, 74, 313-339.	1.0	3
20	New Insights in Adherence and Survival in Myotonic Dystrophy Patients Using Home Mechanical Ventilation. <i>Respiration</i> , 2021, 100, 154-163.	1.2	6
21	Noninvasive Home Mechanical Ventilation in Adult Myotonic Dystrophy Type 1: A Systematic Review. <i>Respiration</i> , 2021, 100, 816-825.	1.2	5
22	Human brain pathology in myotonic dystrophy type 1: A systematic review. <i>Neuropathology</i> , 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. <i>Journal of Neuropathology and Experimental Neurology</i> , 2021, 80, 366-376.	0.9	4
24	Associations Between Variant Repeat Interruptions and Clinical Outcomes in Myotonic Dystrophy Type 1. <i>Neurology: Genetics</i> , 2021, 7, e572.	0.9	10
25	Respiratory muscle imaging by ultrasound and MRI in neuromuscular disorders. <i>European Respiratory Journal</i> , 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. <i>Lancet Neurology</i> , The, 2021, 20, 275-283.	4.9	34
27	The socioeconomic burden of facioscapulohumeral muscular dystrophy. <i>Journal of Neurology</i> , 2021, 268, 4778-4788.	1.8	5
28	Anti-â€Cytosolic 5â€Nucleotidase 1A Autoantibodies Are Absent in Juvenile Dermatomyositis. <i>Arthritis and Rheumatology</i> , 2021, 73, 1329-1333.	2.9	2
29	Phase 1 clinical trial of losmapimod in facioscapulohumeral dystrophy: Safety, tolerability, pharmacokinetics, and target engagement. <i>British Journal of Clinical Pharmacology</i> , 2021, 87, 4658-4669.	1.1	20
30	The facioscapulohumeral muscular dystrophy Rasch-â€built overall disability scale (FSHD-â€RODS). <i>European Journal of Neurology</i> , 2021, 28, 2339-2348.	1.7	8
31	N-of-1 trial of salbutamol in hyperkalaemic periodic paralysis. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2021, 92, jnnp-2021-326347.	0.9	1
32	Behavioural Impairment and Frontotemporal Dementia in Oculopharyngeal Muscular Dystrophy. <i>Journal of Neuromuscular Diseases</i> , 2021, , 1-7.	1.1	1
33	Clinical Outcome Evaluations and CBT Response Prediction in Myotonic Dystrophy. <i>Journal of Neuromuscular Diseases</i> , 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. <i>Neuromuscular Disorders</i> , 2021, 31, 824-828.	0.3	1
35	Longitudinal Assessment of Strength, Functional Capacity, Oropharyngeal Function, and Quality of Life in Oculopharyngeal Muscular Dystrophy. <i>Neurology</i> , 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. <i>BMJ Open</i> , 2021, 11, e048890.	0.8	3

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37	The neuromuscular and multisystem features of RYR1-related malignant hyperthermia and rhabdomyolysis. <i>Medicine (United States)</i> , 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. <i>BMC Neurology</i> , 2021, 21, 313.	0.8	12
39	Clinical, genetic, and histological features of centronuclear myopathy in the Netherlands. <i>Clinical Genetics</i> , 2021, 100, 692-702.	1.0	7
40	Profiling Serum Antibodies Against Muscle Antigens in Facioscapulohumeral Muscular Dystrophy Finds No Disease-Specific Autoantibodies. <i>Journal of Neuromuscular Diseases</i> , 2021, 8, 801-814.	1.1	6
41	MYOTONIC DYSTROPHY. <i>Neuromuscular Disorders</i> , 2021, 31, S121.	0.3	0
42	FSHD. <i>Neuromuscular Disorders</i> , 2021, 31, S99-S100.	0.3	0
43	FSHD. <i>Neuromuscular Disorders</i> , 2021, 31, S100.	0.3	0
44	CONGENITAL MUSCULAR DYSTROPHIES. <i>Neuromuscular Disorders</i> , 2021, 31, S70.	0.3	0
45	CONGENITAL MYOPATHIES. <i>Neuromuscular Disorders</i> , 2021, 31, S65.	0.3	0
46	Rasch analysis to evaluate the motor function measure for patients with facioscapulohumeral muscular dystrophy. <i>International Journal of Rehabilitation Research</i> , 2021, 44, 38-44.	0.7	13
47	Natural History of Facioscapulohumeral Dystrophy in Children. <i>Neurology</i> , 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. <i>PLoS ONE</i> , 2021, 16, e0261475.	1.1	1
49	Short fatigue questionnaire: Screening for severe fatigue.. <i>Journal of Psychosomatic Research</i> , 2020, 137, 110229.	1.2	17
50	Swallowing, Chewing and Speaking: Frequently Impaired in Oculopharyngeal Muscular Dystrophy. <i>Journal of Neuromuscular Diseases</i> , 2020, 7, 483-494.	1.1	8
51	CONGENITAL MYOPATHIES 1 " NEMALINE. <i>Neuromuscular Disorders</i> , 2020, 30, S53.	0.3	0
52	CONGENITAL MYOPATHIES 1 " NEMALINE. <i>Neuromuscular Disorders</i> , 2020, 30, S55.	0.3	0
53	FSHD / OPMD / MYOTONIC DYSTROPHY. <i>Neuromuscular Disorders</i> , 2020, 30, S112-S113.	0.3	0
54	FSHD / OPMD / MYOTONIC DYSTROPHY. <i>Neuromuscular Disorders</i> , 2020, 30, S113.	0.3	0

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55	FSHD / OPMD / MYOTONIC DYSTROPHY. <i>Neuromuscular Disorders</i> , 2020, 30, S113.	0.3	0
56	Deep phenotyping of facioscapulohumeral muscular dystrophy type 2 by magnetic resonance imaging. <i>European Journal of Neurology</i> , 2020, 27, 2604-2615.	1.7	16
57	Clinical, morphological and genetic characterization of Brody disease: an international study of 40 patients. <i>Brain</i> , 2020, 143, 452-466.	3.7	22
58	Age-Associated Salivary MicroRNA Biomarkers for Oculopharyngeal Muscular Dystrophy. <i>International Journal of Molecular Sciences</i> , 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. <i>Journal of Neuromuscular Diseases</i> , 2020, 7, 495-504.	1.1	25
60	Swallowing, Chewing and Speaking: Frequently Impaired in Oculopharyngeal Muscular Dystrophy. <i>Journal of Neuromuscular Diseases</i> , 2020, 7, 1-12.	1.1	12
61	Quantitative Muscle MRI Depicts Increased Muscle Mass after a Behavioral Change in Myotonic Dystrophy Type 1. <i>Radiology</i> , 2020, 297, 132-142.	3.6	11
62	Inclusion body myositis in patients with spinocerebellar ataxia types 3 and 6. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2020, 91, 876-878.	0.9	2
63	Altered sensorimotor representations after recovery from peripheral nerve damage in neuralgic amyotrophy. <i>Cortex</i> , 2020, 127, 180-190.	1.1	10
64	Muscle ultrasound is a responsive biomarker in facioscapulohumeral dystrophy. <i>Neurology</i> , 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. <i>Healthcare (Switzerland)</i> , 2020, 8, 49.	1.0	18
66	Characterization of EEG-based functional brain networks in myotonic dystrophy type 1. <i>Clinical Neurophysiology</i> , 2020, 131, 1886-1895.	0.7	1
67	Consequences of epigenetic derepression in facioscapulohumeral muscular dystrophy. <i>Clinical Genetics</i> , 2020, 97, 799-814.	1.0	40
68	Preserved single muscle fiber specific force in facioscapulohumeral muscular dystrophy. <i>Neurology</i> , 2020, 94, e1157-e1170.	1.5	8
69	Continued misuse of orphan drug legislation: a life-threatening risk for mexiletine. <i>European Heart Journal</i> , 2020, 41, 614-617.	1.0	15
70	Guidelines on clinical presentation and management of nondystrophic myotonias. <i>Muscle and Nerve</i> , 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. <i>Neuromuscular Disorders</i> , 2020, 30, 521-531.	0.3	1
72	KBTBD13 is an actin-binding protein that modulates muscle kinetics. <i>Journal of Clinical Investigation</i> , 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. <i>Trials</i> , 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. <i>Journal of Medical Genetics</i> , 2019, 56, 693-700.	1.5	27
75	Ophthalmological findings in facioscapulohumeral dystrophy. <i>Brain Communications</i> , 2019, 1, fcz023.	1.5	14
76	O.25Phase 1 clinical trial of losmapimod in FSHD: safety, tolerability and target engagement. <i>Neuromuscular Disorders</i> , 2019, 29, S123.	0.3	1
77	P.40Ophthalmological findings in facioscapulohumeral dystrophy. <i>Neuromuscular Disorders</i> , 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. <i>Neuromuscular Disorders</i> , 2019, 29, S54.	0.3	0
79	P.63Chronic progressive external ophthalmoplegia (CPEO) and CPEO-plus cohort of 54 patients from the Netherlands. <i>Neuromuscular Disorders</i> , 2019, 29, S59.	0.3	0
80	P.162Novel Kbtbd13R408C-knockin mouse model phenocopies NEM6 myopathy. <i>Neuromuscular Disorders</i> , 2019, 29, S95.	0.3	0
81	O.13Nemaline myopathy patients with mutations in KBTBD13 display a cardiac phenotype. <i>Neuromuscular Disorders</i> , 2019, 29, S118.	0.3	0
82	P.247Capturing disease progression in oculopharyngeal muscular dystrophy (OPMD). <i>Neuromuscular Disorders</i> , 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. <i>Neuromuscular Disorders</i> , 2019, 29, S156.	0.3	0
84	High incidence of falls in patients with myotonic dystrophy type 1 and 2: A prospective study. <i>Neuromuscular Disorders</i> , 2019, 29, 758-765.	0.3	10
85	Scapular dyskinesia in myotonic dystrophy type 1: clinical characteristics and genetic investigations. <i>Journal of Neurology</i> , 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. <i>BMC Neurology</i> , 2019, 19, 224.	0.8	28
87	Effects of weakness of orofacial muscles on swallowing and communication in FSHD. <i>Neurology</i> , 2019, 92, e957-e963.	1.5	25
88	Lower extremity muscle pathology in myotonic dystrophy type 1 assessed by quantitative MRI. <i>Neurology</i> , 2019, 92, e2803-e2814.	1.5	34
89	MSH3 modifies somatic instability and disease severity in Huntington's and myotonic dystrophy type 1. <i>Brain</i> , 2019, 142, 1876-1886.	3.7	114
90	Panel-Based Exome Sequencing for Neuromuscular Disorders as a Diagnostic Service. <i>Journal of Neuromuscular Diseases</i> , 2019, 6, 241-258.	1.1	32

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

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109	Hearing impairment in patients with myotonic dystrophy type 2. <i>Neurology</i> , 2018, 90, e615-e622.	1.5	11
110	Falls and resulting fractures in Myotonic Dystrophy: Results from a multinational retrospective survey. <i>Neuromuscular Disorders</i> , 2018, 28, 229-235.	0.3	19
111	Electrical impedance myography in facioscapulohumeral muscular dystrophy: A 1-year follow-up study. <i>Muscle and Nerve</i> , 2018, 58, 213-218.	1.0	15
112	Qualitative and Quantitative Aspects of Pain in Patients With Myotonic Dystrophy Type 2. <i>Journal of Pain</i> , 2018, 19, 920-930.	0.7	19
113	Prevalence and mutation spectrum of skeletal muscle channelopathies in the Netherlands. <i>Neuromuscular Disorders</i> , 2018, 28, 402-407.	0.3	40
114	MRI in sarcoglycanopathies: a large international cohort study. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 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. <i>European Journal of Human Genetics</i> , 2018, 26, 94-106.	1.4	22
116	Specific muscle strength is reduced in facioscapulohumeral dystrophy: An MRI based musculoskeletal analysis. <i>Neuromuscular Disorders</i> , 2018, 28, 238-245.	0.3	11
117	Evidence of ER stress and UPR activation in patients with Brody disease and Brody syndrome. <i>Neuropathology and Applied Neurobiology</i> , 2018, 44, 533-536.	1.8	1
118	CONGENITAL MYOPATHIES: NEMALINE AND TITINOPATHIES. <i>Neuromuscular Disorders</i> , 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. <i>JAMA - Journal of the American Medical Association</i> , 2018, 320, 2344.	3.8	81
120	Quantitative muscle MRI and ultrasound for facioscapulohumeral muscular dystrophy: complementary imaging biomarkers. <i>Journal of Neurology</i> , 2018, 265, 2646-2655.	1.8	54
121	Consensus-based care recommendations for adults with myotonic dystrophy type 1. <i>Neurology: Clinical Practice</i> , 2018, 8, 507-520.	0.8	115
122	Facioscapulohumeral Dystrophy in Childhood: A Nationwide Natural History Study. <i>Annals of Neurology</i> , 2018, 84, 627-637.	2.8	21
123	NEW GENES, FUNCTIONS AND BIOMARKERS. <i>Neuromuscular Disorders</i> , 2018, 28, S31.	0.3	0
124	Phenotype-genotype relations in facioscapulohumeral muscular dystrophy type 1. <i>Clinical Genetics</i> , 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. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 155.	1.2	19
126	Second IVIg course in Guillain-Barré syndrome patients with poor prognosis (SIDeCBS trial): Protocol for a double-blind randomized, placebo-controlled clinical trial. <i>Journal of the Peripheral Nervous System</i> , 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. <i>European Journal of Paediatric Neurology</i> , 2018, 22, 782-785.	0.7	8
128	Cis D4Z4 repeat duplications associated with facioscapulohumeral muscular dystrophy type 2. <i>Human Molecular Genetics</i> , 2018, 27, 3488-3497.	1.4	27
129	Autoantibodies to Cytosolic 5â€²-Nucleotidase 1A in Primary Sjögrenâ€™s Syndrome and Systemic Lupus Erythematosus. <i>Frontiers in Immunology</i> , 2018, 9, 1200.	2.2	32
130	A family-based study into penetrance in facioscapulohumeral muscular dystrophy type 1. <i>Neurology</i> , 2018, 91, e444-e454.	1.5	33
131	FSHD type 2 and Bosma arhinia microphthalmia syndrome. <i>Neurology</i> , 2018, 91, e562-e570.	1.5	24
132	MRI-Guided Biopsy as a Tool for Diagnosis and Research of Muscle Disorders. <i>Journal of Neuromuscular Diseases</i> , 2018, 5, 315-319.	1.1	24
133	Repeatability and reliability of muscle relaxation properties induced by motor cortical stimulation. <i>Journal of Applied Physiology</i> , 2018, 124, 1597-1604.	1.2	5
134	Fatigue, not self-rated motor symptom severity, affects quality of life in functional motor disorders. <i>Journal of Neurology</i> , 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. <i>Lancet Neurology</i> , The, 2018, 17, 671-680.	4.9	95
136	Is fatigue a disease-specific or generic symptom in chronic medical conditions?. <i>Health Psychology</i> , 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. <i>Neuromuscular Disorders</i> , 2017, 27, 370-376.	0.3	31
139	Cytosolic 5â€²-nucleotidase 1A autoantibody profile and clinical characteristics in inclusion body myositis. <i>Annals of the Rheumatic Diseases</i> , 2017, 76, 862-868.	0.5	71
140	225th ENMC international workshop:. <i>Neuromuscular Disorders</i> , 2017, 27, 782-790.	0.3	20
141	The assessment of fatigue: Psychometric qualities and norms for the Checklist individual strength. <i>Journal of Psychosomatic Research</i> , 2017, 98, 40-46.	1.2	222
142	Respiratory function in facioscapulohumeral muscular dystrophy 1. <i>Neuromuscular Disorders</i> , 2017, 27, 526-530.	0.3	14
143	Respiratory pattern in a FSHD pediatric population. <i>Respiratory Medicine</i> , 2017, 126, 130-131.	1.3	0
144	Oculopharyngeal muscular dystrophy with frontotemporal dementia. <i>European Geriatric Medicine</i> , 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. <i>Neuromuscular Disorders</i> , 2017, 27, 243-250.	0.3	10
146	Involvement of pelvic girdle and proximal leg muscles in early oculopharyngeal muscular dystrophy. <i>Neuromuscular Disorders</i> , 2017, 27, 1099-1105.	0.3	17
147	Muscle fiber dysfunction contributes to clinical muscle weakness in inclusion body myositis. <i>Neuromuscular Disorders</i> , 2017, 27, S154.	0.3	0
148	The Brody disease cohort study: clarification of the phenotype. <i>Neuromuscular Disorders</i> , 2017, 27, S164.	0.3	0
149	Adding quantitative muscle MRI to the FSHD clinical trial toolbox. <i>Neuromuscular Disorders</i> , 2017, 27, S122.	0.3	1
150	Specific strength is reduced in facioscapulohumeral dystrophy muscles. An MRI-based musculoskeletal analysis. <i>Neuromuscular Disorders</i> , 2017, 27, S200.	0.3	0
151	Adding quantitative muscle MRI to the FSHD clinical trial toolbox. <i>Neurology</i> , 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 et al. <i>Arthritis and Rheumatology</i> , 2017, 69, 2248-2249.	2.9	4
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