Nanna Witting

List of Publications by Year
in descending order

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58 papers	2,056 citations	279487 23 h-index	243296 44 g-index
58 all docs	58 docs citations	58 times ranked	2654 citing authors

#	Article	IF	Citations
1	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. Lancet Neurology, The, 2017, 16, 976-986.	4.9	472
2	Longâ€term safety and efficacy of eculizumab in generalized myasthenia gravis. Muscle and Nerve, 2019, 60, 14-24.	1.0	162
3	Cardiac manifestations of myotonic dystrophy type 1. International Journal of Cardiology, 2012, 160, 82-88.	0.8	146
4	A PET activation study of brush-evoked allodynia in patientswith nerve injury pain. Pain, 2006, 120, 145-154.	2.0	122
5	Congenital Titinopathy: Comprehensive characterization and pathogenic insights. Annals of Neurology, 2018, 83, 1105-1124.	2.8	93
6	A heterozygous 21-bp deletion inCAPN3causes dominantly inherited limb girdle muscular dystrophy. Brain, 2016, 139, 2154-2163.	3.7	87
7	Anoctamin 5 muscular dystrophy in Denmark: prevalence, genotypes, phenotypes, cardiac findings, and muscle protein expression. Journal of Neurology, 2013, 260, 2084-2093.	1.8	63
8	Two- and 6-minute walk tests assess walking capability equally in neuromuscular diseases. Neurology, 2016, 86, 442-445.	1.5	51
9	Clinical and genetic spectrum in limb-girdle muscular dystrophy type 2E. Neurology, 2015, 84, 1772-1781.	1.5	50
10	The antimyotonic effect of lamotrigine in non-dystrophic myotonias: a double-blind randomized study. Brain, 2017, 140, 2295-2305.	3.7	49
11	High prevalence of cardiac involvement in patients with myotonic dystrophy type 1: A cross-sectional study. International Journal of Cardiology, 2014, 174, 31-36.	0.8	44
12	Axial myopathy: an overlooked feature of muscle diseases. Brain, 2016, 139, 13-22.	3.7	44
13	Myocardial fibrosis in patients with myotonic dystrophy type 1: a cardiovascular magnetic resonance study. Journal of Cardiovascular Magnetic Resonance, 2014, 16, 59.	1.6	43
14	Effect of sildenafil on skeletal and cardiac muscle in <scp>B</scp>ecker muscular dystrophy. Annals of Neurology, 2014, 76, 550-557.	2.8	39
15	Progression of cardiac involvement in patients with limb-girdle type 2 and Becker muscular dystrophies: A 9-year follow-up study. International Journal of Cardiology, 2015, 182, 403-411.	0.8	36
16	Phenotypes, genotypes, and prevalence of congenital myopathies older than 5 years in Denmark. Neurology: Genetics, 2017, 3, e140.	0.9	34
17	Panel-Based Exome Sequencing for Neuromuscular Disorders as a Diagnostic Service. Journal of Neuromuscular Diseases, 2019, 6, 241-258.	1.1	32
18	Eculizumab improves fatigue in refractory generalized myasthenia gravis. Quality of Life Research, 2019, 28, 2247-2254.	1.5	32

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19	Phenotype and genotype of muscle ryanodine receptor rhabdomyolysis-myalgia syndrome. Acta Neurologica Scandinavica, 2018, 137, 452-461.	1.0	31
20	MSTO1 mutations cause mtDNA depletion, manifesting as muscular dystrophy with cerebellar involvement. Acta Neuropathologica, 2019, 138, 1013-1031.	3.9	31
21	Recurrent TTN metatranscriptâ€only c.39974â€“11T>G splice variant associated with autosomal recessive arthrogryposis multiplex congenita and myopathy. Human Mutation, 2020, 41, 403-411.	1.1	28
22	Pharmacologic Treatment of Downstream of Tyrosine Kinase 7 Congenital Myasthenic Syndrome. JAMA Neurology, 2014, 71, 350.	4.5	27
23	Role of neuronal nitric oxide synthase (nNOS) in Duchenne and Becker muscular dystrophies â€“ Still a possible treatment modality?. Neuromuscular Disorders, 2018, 28, 914-926.	0.3	24
24	Decreased Variability of the 6-Minute Walk Test by Heart Rate Correction in Patients with Neuromuscular Disease. PLoS ONE, 2014, 9, e114273.	1.1	24
25	Ocular, bulbar, limb, and cardiopulmonary involvement in oculopharyngeal muscular dystrophy. Acta Neurologica Scandinavica, 2014, 130, 125-130.	1.0	23
26	Severe Axial Myopathy in McArdle Disease. JAMA Neurology, 2014, 71, 88.	4.5	18
27	Prevalence and phenotypes of congenital myopathy due to Î±â€actin 1 gene mutations. Muscle and Nerve, 2016, 53, 388-393.	1.0	18
28	Long-term efficacy and safety of eculizumab in Japanese patients with generalized myasthenia gravis: A subgroup analysis of the REGAIN open-label extension study. Journal of the Neurological Sciences, 2019, 407, 116419.	0.3	18
29	Frequency and phenotype of patients carrying TPM2 and TPM3 gene mutations in a cohort of 94 patients with congenital myopathy. Neuromuscular Disorders, 2014, 24, 325-330.	0.3	17
30	Aerobic Training in Patients with Congenital Myopathy. PLoS ONE, 2016, 11, e0146036.	1.1	17
31	Permanent muscle weakness in hypokalemic periodic paralysis. Neurology, 2020, 95, e342-e352.	1.5	17
32	Consistent improvement with eculizumab across muscle groups in myasthenia gravis. Annals of Clinical and Translational Neurology, 2020, 7, 1327-1339.	1.7	16
33	Becker muscular dystrophy with widespread muscle hypertrophy and a non-sense mutation of exon 2. Neuromuscular Disorders, 2013, 23, 25-28.	0.3	15
34	Effects of Sildenafil on Cerebrovascular Reactivity in Patients with Becker Muscular Dystrophy. Neurotherapeutics, 2017, 14, 182-190.	2.1	14
35	Muscle involvement assessed by quantitative magnetic resonance imaging in patients with anoctamin 5 deficiency. European Journal of Neurology, 2021, 28, 3121-3132.	1.7	13
36	Collagen XII myopathy with rectus femoris atrophy and collagen XII retention in fibroblasts. Muscle and Nerve, 2018, 57, 1026-1030.	1.0	11

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37	Eculizumab in refractory generalized myasthenia gravis previously treated with rituximab: subgroup analysis of <scp>REGAIN</scp> and its extension study. Muscle and Nerve, 2021, 64, 662-669.	1.0	11
38	Frequency and Phenotype of Myotubular Myopathy Amongst Danish Patients with Congenital Myopathy Older than 5 Years. Journal of Neuromuscular Diseases, 2015, 2, 167-174.	1.1	10
39	Congenital myopathies are mainly associated with a mild cardiac phenotype. Journal of Neurology, 2019, 266, 1367-1375.	1.8	10
40	Quantitative Muscle MRI as Outcome Measure in Patients With Becker Muscular Dystrophyâ€"A 1-Year Follow-Up Study. Frontiers in Neurology, 2020, 11, 613489.	1.1	9
41	Intrarater reliability and validity of outcome measures in myotonic dystrophy type 1. Neurology, 2020, 94, e2508-e2520.	1.5	7
42	Physical activity in myotonic dystrophy type 1. Journal of Neurology, 2020, 267, 1679-1686.	1.8	7
43	Cardiac arrest in anti-mitochondrial antibody associated inflammatory myopathy. Oxford Medical Case Reports, 2021, 2021, omaa150.	0.2	6
44	Muscle biopsy and <scp>MRI</scp> findings in <scp>ANO5</scp>â€related myopathy. Muscle and Nerve, 2021, 64, 743-748.	1.0	6
45	Botulinum toxin treatment improves dysphagia in patients with oculopharyngeal muscular dystrophy and sporadic inclusion body myositis. Journal of Neurology, 2022, 269, 4154-4160.	1.8	6
46	Clinical and neurophysiological response to pharmacological treatment of DOK7 congenital myasthenia in an older patient. Clinical Neurology and Neurosurgery, 2015, 130, 168-170.	0.6	5
47	Deletion of exon 26 of the dystrophin gene is associated with a mild Becker muscular dystrophy phenotype. Acta Myologica, 2011, 30, 182-4.	1.5	5
48	Axial muscle involvement in patients with limb girdle muscular dystrophy type <scp>R9</scp>. Muscle and Nerve, 2022, 65, 405-414.	1.0	3
49	Responsiveness of outcome measures in myotonic dystrophy type 1. Annals of Clinical and Translational Neurology, 2020, 7, 1382-1391.	1.7	2
50	Contractile properties are impaired in congenital myopathies. Neuromuscular Disorders, 2020, 30, 649-655.	0.3	2
51	Patients With Becker Muscular Dystrophy Have Severe Paraspinal Muscle Involvement. Frontiers in Neurology, 2021, 12, 613483.	1.1	2
52	Autophagy is affected in patients with hypokalemic periodic paralysis: an involvement in vacuolar myopathy?. Acta Neuropathologica Communications, 2021, 9, 109.	2.4	2
53	Altered somatosensory neurovascular response in patients with Becker muscular dystrophy. Brain and Behavior, 2018, 8, e00985.	1.0	1
54	Progression or Not â€“ A Small Natural History Study of Genetical Confirmed Congenital Myopathies. Journal of Neuromuscular Diseases, 2021, 8, 647-655.	1.1	1

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55	Response. Neuromuscular Disorders, 2013, 23, 193.	0.3	0
56	AB0652â€…QTC INTERVAL PROLONGATION IN A SCANDINAVIAN COHORT OF PATIENTS WITH IDIOPATHIC INFLAMMATORY MYOPATHIES AND SYSTEMIC SCLEROSIS: CORRELATIONS WITH CLINICAL VARIABLES. , 2019, , .		0
57	FRI0320â€…CN-1A AUTOANTIBODIES ARE SPECIFIC FOR SPORADIC INCLUSION BODY MYOSITIS. , 2019, , .		0
58	Effects of rhythmic auditory stimulation on walking during the 6-minute walk test in patients with generalised Myasthenia Gravis. European Journal of Physiotherapy, 0, , 1-000.	0.7	0