

Jodi Warman Chardon

List of Publications by Year in descending order

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Version: 2024-02-01

56
papers

1,045
citations

535685

17
h-index

511568

30
g-index

60
all docs

60
docs citations

60
times ranked

2494
citing authors

#	ARTICLE	IF	CITATIONS
1	Laryngospasm in amyotrophic lateral sclerosis. <i>Muscle and Nerve</i> , 2022, 65, 400-404.	1.0	6
2	Temporal evolution of nerve conduction study abnormalities in anti- <i>Myelin-associated glycoprotein</i> neuropathy. <i>Muscle and Nerve</i> , 2021, 63, 401-404.	1.0	10
3	Autologous Hematopoietic Stem Cell Transplantation for Chronic Inflammatory Demyelinating Polyradiculoneuropathy. <i>Canadian Journal of Neurological Sciences</i> , 2021, , 1-7.	0.3	3
4	Whole genome sequencing reveals biallelic <i>PLA2G6</i> mutations in siblings with cerebellar atrophy and cap myopathy. <i>Clinical Genetics</i> , 2021, 99, 746-748.	1.0	3
5	A Canadian Adult Spinal Muscular Atrophy Outcome Measures Toolkit: Results of a National Consensus using a Modified Delphi Method. <i>Journal of Neuromuscular Diseases</i> , 2021, 8, 579-588.	1.1	7
6	Pseudohypertrophy of the extensor digitorum brevis in diabetic polyneuropathy. <i>Muscle and Nerve</i> , 2021, 64, E20-E22.	1.0	0
7	MuSK not MNGIE: Atypical MuSK-antibody myasthenia presenting as a genetic disorder. <i>Neuromuscular Disorders</i> , 2021, , .	0.3	0
8	Intrafamilial variability of limb-girdle muscular dystrophy, LGMD1D type. <i>European Journal of Medical Genetics</i> , 2020, 63, 103655.	0.7	10
9	Impact of disuse muscular atrophy on the compound muscle action potential. <i>Muscle and Nerve</i> , 2020, 61, 58-62.	1.0	5
10	247th ENMC International Workshop: Muscle magnetic resonance imaging - Implementing muscle MRI as a diagnostic tool for rare genetic myopathy cohorts. Hoofddorp, The Netherlands, September 2019. <i>Neuromuscular Disorders</i> , 2020, 30, 938-947.	0.3	11
11	Myofibrillar Myopathy Mimicking Polyneuropathy. <i>Case Reports in Neurology</i> , 2020, 12, 97-102.	0.3	1
12	Advances in the diagnosis of inherited neuromuscular diseases and implications for therapy development. <i>Lancet Neurology</i> , The, 2020, 19, 522-532.	4.9	36
13	A National Spinal Muscular Atrophy Registry for Real-World Evidence. <i>Canadian Journal of Neurological Sciences</i> , 2020, 47, 810-815.	0.3	6
14	Novel Recessive <i>TNNT1</i> Congenital Core-Rod Myopathy in French Canadians. <i>Annals of Neurology</i> , 2020, 87, 568-583.	2.8	19
15	Abnormal fatty acid metabolism is a core component of spinal muscular atrophy. <i>Annals of Clinical and Translational Neurology</i> , 2019, 6, 1519-1532.	1.7	72
16	Intermittent undulating tongue as an involuntary movement in early amyotrophic lateral sclerosis. <i>Parkinsonism and Related Disorders</i> , 2019, 67, 1-2.	1.1	1
17	A Survey of Cerebrospinal Fluid Total Protein Upper Limits in Canada: Time for an Update?. <i>Canadian Journal of Neurological Sciences</i> , 2019, 46, 283-286.	0.3	2
18	The Added Value of Cardiac Magnetic Resonance in Muscular Dystrophies. <i>Journal of Neuromuscular Diseases</i> , 2019, 6, 389-399.	1.1	10

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19	MYO-MRI diagnostic protocols in genetic myopathies. <i>Neuromuscular Disorders</i> , 2019, 29, 827-841.	0.3	46
20	Age matters. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2019, 6, e576.	3.1	7
21	<i>PDXK</i> mutations cause polyneuropathy responsive to pyridoxal 5-phosphate supplementation. <i>Annals of Neurology</i> , 2019, 86, 225-240.	2.8	54
22	Dataset for worldwide survey of cerebrospinal total protein upper reference values. <i>Data in Brief</i> , 2019, 23, 103760.	0.5	0
23	Neurolymphomatosis of the lumbosacral plexus and its branches: case series and literature review. <i>BMC Cancer</i> , 2019, 19, 1149.	1.1	16
24	Distal Cervical Spondylotic Amyotrophy: Case Reports Demonstrating Clinical/Imaging Segmental Discrepancy. <i>Journal of Clinical Neuromuscular Disease</i> , 2019, 21, 107-111.	0.3	2
25	Targeted exome analysis identifies the genetic basis of disease in over 50% of patients with a wide range of ataxia-related phenotypes. <i>Genetics in Medicine</i> , 2019, 21, 195-206.	1.1	65
26	Neurolymphomatosis of the Brachial Plexus and its Branches: Case Series and Literature Review. <i>Canadian Journal of Neurological Sciences</i> , 2018, 45, 137-143.	0.3	26
27	Novel <i>ELOVL4</i> mutation associated with erythrokeratoderma and spinocerebellar ataxia (SCA) Tj ETQq1 1 0.784314 rgBT /Ove	0.9	27
28	Teaching Video NeuroImages: Rippling muscle disease with caveolin myopathy. <i>Neurology</i> , 2018, 91, e1726-e1727.	1.5	1
29	Intraneural Ganglion Cysts of the Fibular Nerve: A Cause of Fluctuating Painful Foot Drop. <i>Canadian Journal of Neurological Sciences</i> , 2018, 45, 601-603.	0.3	0
30	Whole-transcriptome sequencing in blood provides a diagnosis of spinal muscular atrophy with progressive myoclonic epilepsy. <i>Human Mutation</i> , 2017, 38, 611-614.	1.1	25
31	Cardiac Amyloidosis Phenotype Associated With a Glu89Lys Transthyretin Mutation. <i>Canadian Journal of Cardiology</i> , 2017, 33, 830.e5-830.e7.	0.8	2
32	Clinical and genetic study of hereditary spastic paraplegia in Canada. <i>Neurology: Genetics</i> , 2017, 3, e122.	0.9	82
33	Survival Motor Neuron Protein is Released from Cells in Exosomes: A Potential Biomarker for Spinal Muscular Atrophy. <i>Scientific Reports</i> , 2017, 7, 13859.	1.6	13
34	Teaching Video Neuro Images : Trapezius muscle hypertrophy in multifocal motor neuropathy. <i>Neurology</i> , 2017, 89, e81-e82.	1.5	0
35	The Role of Muscle Imaging in the Diagnosis and Assessment of Children with Genetic Muscle Disease. <i>Neuropediatrics</i> , 2017, 48, 233-241.	0.3	11
36	Combined isolated trigeminal and facial neuropathies from perineural invasion by squamous cell carcinoma: A case series and review of the literature. <i>Journal of Clinical Neuroscience</i> , 2017, 35, 5-12.	0.8	7

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37	Autosomal dominant cerebellar ataxia, deafness, and narcolepsy (ADCA-DN) associated with progressive cognitive and behavioral deterioration.. <i>Neuropsychology</i> , 2017, 31, 292-303.	1.0	4
38	A crucial first randomized controlled trial of thymectomy in non-thymomatous myasthenia gravis. <i>Journal of Thoracic Disease</i> , 2016, 8, E1375-E1378.	0.6	2
39	Supramaximal Stimulus Intensity as a Diagnostic Tool in Chronic Demyelinating Neuropathy. <i>Neuroscience Journal</i> , 2016, 2016, 1-5.	2.3	5
40	Diffuse leukoencephalopathy with spheroids presenting as primary progressive aphasia. <i>Neurology</i> , 2016, 86, 1464-1465.	1.5	3
41	Identification of a methylation profile for DNMT1-associated autosomal dominant cerebellar ataxia, deafness, and narcolepsy. <i>Clinical Epigenetics</i> , 2016, 8, 91.	1.8	66
42	Marked enlargement of neck circumference from nerve hypertrophy in CIDP. <i>Neurology</i> , 2016, 87, 442-442.	1.5	1
43	Systematic analysis of clinical deficits in unilateral hypoglossal nerve palsy. <i>Muscle and Nerve</i> , 2016, 54, 1055-1058.	1.0	0
44	Syndrome disintegration: Exome sequencing reveals that Fitzsimmons syndrome is a co-occurrence of multiple events. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 1820-1825.	0.7	19
45	Amyloid Neuropathy Following Domino Liver Transplantation. <i>JAMA Neurology</i> , 2016, 73, 477.	4.5	4
46	Very late-onset Sandhoff disease presenting as Kennedy Disease. <i>Muscle and Nerve</i> , 2015, 52, 1135-1136.	1.0	4
47	LIMS2 mutations are associated with a novel muscular dystrophy, severe cardiomyopathy and triangular tongues. <i>Clinical Genetics</i> , 2015, 88, 558-564.	1.0	23
48	Autoimmune peripheral neuropathies. <i>Clinica Chimica Acta</i> , 2015, 449, 37-42.	0.5	26
49	Axons to Exons: the Molecular Diagnosis of Rare Neurological Diseases by Next-Generation Sequencing. <i>Current Neurology and Neuroscience Reports</i> , 2015, 15, 64.	2.0	29
50	Exome Sequencing as a Diagnostic Tool for Pediatric Onset Ataxia. <i>Human Mutation</i> , 2014, 35, 45-49.	1.1	91
51	The utility of exome sequencing for genetic diagnosis in a familial microcephaly epilepsy syndrome. <i>BMC Neurology</i> , 2014, 14, 22.	0.8	18
52	Deletion of <i>AFG3L2</i> associated with spinocerebellar ataxia type 28 in the context of multiple genomic anomalies. <i>American Journal of Medical Genetics, Part A</i> , 2014, 164, 3209-3212.	0.7	11
53	Recent Advances in the Genetic Etiology of Brain Malformations. <i>Current Neurology and Neuroscience Reports</i> , 2013, 13, 364.	2.0	20
54	Missense mutations in ITPR1 cause autosomal dominant congenital nonprogressive spinocerebellar ataxia. <i>Orphanet Journal of Rare Diseases</i> , 2012, 7, 67.	1.2	124

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55	Deletion of filamin A in two female patients with periventricular nodular heterotopia. American Journal of Medical Genetics, Part A, 2012, 158A, 1512-1516.	0.7	6
56	Does Diabetes Alter CSF Total Protein Levels? A Retrospective Cohort Study. Neurohospitalist, The, 0, , 194187442110393.	0.3	0