Pierre R Bourque

List of Publications by Year in descending order

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566801 610482 52 639 15 24 citations h-index g-index papers 53 53 53 1020 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Musical Auditory Hallucinosis from Listeria Rhombencephalitis. Canadian Journal of Neurological Sciences, 1997, 24, 70-72.	0.3	61
2	Cerebrospinal Fluid Total Protein Reference Intervals Derived from 20 Years of Patient Data. Clinical Chemistry, 2017, 63, 1856-1865.	1.5	61
3	<i>PDXK</i> mutations cause polyneuropathy responsive to pyridoxal 5′â€phosphate supplementation. Annals of Neurology, 2019, 86, 225-240.	2.8	54
4	Adult CSF total protein upper reference limits should be age-partitioned and significantly higher than 0.45Âg/L: a systematic review. Journal of Neurology, 2019, 266, 616-624.	1.8	41
5	Updated cerebrospinal fluid total protein reference values improve chronic inflammatory demyelinating polyneuropathy diagnosis. Muscle and Nerve, 2019, 60, 180-183.	1.0	37
6	Advances in the diagnosis of inherited neuromuscular diseases and implications for therapy development. Lancet Neurology, The, 2020, 19, 522-532.	4.9	36
7	Novel <i>ELOVL4</i> mutation associated with erythrokeratodermia and spinocerebellar ataxia (SCA) Tj ETQq1 1	0,784314	rgBT /Overla
8	Autoimmune peripheral neuropathies. Clinica Chimica Acta, 2015, 449, 37-42.	0.5	26
9	Neurolymphomatosis of the Brachial Plexus and its Branches: Case Series and Literature Review. Canadian Journal of Neurological Sciences, 2018, 45, 137-143.	0.3	26
10	Causes of albuminocytological dissociation and the impact of age-adjusted cerebrospinal fluid protein reference intervals: a retrospective chart review of 2627 samples collected at tertiary care centre. BMJ Open, 2019, 9, e025348.	0.8	26
11	Whole-transcriptome sequencing in blood provides a diagnosis of spinal muscular atrophy with progressive myoclonic epilepsy. Human Mutation, 2017, 38, 611-614.	1.1	25
12	Dying of amyotrophic lateral sclerosis. Neurology, 2019, 93, e2083-e2093.	1.5	24
13	Myopathy with hexagonally cross-linked tubular arrays: A new autosomal dominant or sporadic congenital myopathy. Annals of Neurology, 1999, 45, 512-515.	2.8	20
14	Adult CSF total protein: Higher upper reference limits should be considered worldwide. A web-based survey. Journal of the Neurological Sciences, 2019, 396, 48-51.	0.3	20
15	Phenotypic variability of CMT4C in a Frenchâ€Canadian kindred. Muscle and Nerve, 2015, 52, 444-449.	1.0	18
16	Neurolymphomatosis of the lumbosacral plexus and its branches: case series and literature review. BMC Cancer, 2019, 19, 1149.	1.1	16
17	Cerebrospinal fluid total protein in Guillain–Barré syndrome variants: correlations with clinical category, severity, and electrophysiology. Journal of Neurology, 2020, 267, 746-751.	1.8	15
18	Edaravone for amyotrophic lateral sclerosis: barriers to access and lifeboat ethics. Cmaj, 2020, 192, E319-E320.	0.9	12

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19	Paraneoplastic recurrent tumefactive demyelination in a 62-year-old man with metastatic seminoma. Neurology: Neuroimmunology and NeuroInflammation, 2019, 6, e527.	3.1	10
20	Temporal evolution of nerve conduction study abnormalities in antiâ€myelinâ€associated glycoprotein neuropathy. Muscle and Nerve, 2021, 63, 401-404.	1.0	10
21	Evaluation of a Personalized Subcutaneous Immunoglobulin Treatment Program for Neurological Patients. Canadian Journal of Neurological Sciences, 2019, 46, 38-43.	0.3	8
22	Combined isolated trigeminal and facial neuropathies from perineural invasion by squamous cell carcinoma: A case series and review of the literature. Journal of Clinical Neuroscience, 2017, 35, 5-12.	0.8	7
23	Age matters. Neurology: Neuroimmunology and NeuroInflammation, 2019, 6, e576.	3.1	7
24	Laryngospasm in amyotrophic lateral sclerosis. Muscle and Nerve, 2022, 65, 400-404.	1.0	6
25	Supramaximal Stimulus Intensity as a Diagnostic Tool in Chronic Demyelinating Neuropathy. Neuroscience Journal, 2016, 2016, 1-5.	2.3	5
26	Impact of disuse muscular atrophy on the compound muscle action potential. Muscle and Nerve, 2020, 61, 58-62.	1.0	5
27	Amyloid Neuropathy Following Domino Liver Transplantation. JAMA Neurology, 2016, 73, 477.	4. 5	4
28	Myasthenia gravis. Cmaj, 2018, 190, E1141-E1141.	0.9	4
29	Spinal Capillary Hemangioma: A Rare Benign Extradural Tumor. Canadian Journal of Neurological Sciences, 2020, 47, 549-550.	0.3	4
30	Autosomal dominant cerebellar ataxia, deafness, and narcolepsy (ADCA-DN) associated with progressive cognitive and behavioral deterioration Neuropsychology, 2017, 31, 292-303.	1.0	4
31	Autologous Hematopoietic Stem Cell Transplantation for Chronic Inflammatory Demyelinating Polyradiculoneuropathy. Canadian Journal of Neurological Sciences, 2021, , 1-7.	0.3	3
32	A crucial first randomized controlled trial of thymectomy in non-thymomatous myasthenia gravis. Journal of Thoracic Disease, 2016, 8, E1375-E1378.	0.6	2
33	Cardiac Amyloidosis Phenotype Associated With a Glu89Lys Transthyretin Mutation. Canadian Journal of Cardiology, 2017, 33, 830.e5-830.e7.	0.8	2
34	A Survey of Cerebrospinal Fluid Total Protein Upper Limits in Canada: Time for an Update?. Canadian Journal of Neurological Sciences, 2019, 46, 283-286.	0.3	2
35	Distal Cervical Spondylotic Amyotrophy: Case Reports Demonstrating Clinical/Imaging Segmental Discrepancy. Journal of Clinical Neuromuscular Disease, 2019, 21, 107-111.	0.3	2
36	Systematic prospective electrophysiological studies of the median nerve after simple distal radius fracture. PLoS ONE, 2020, 15, e0231502.	1.1	2

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37	Marked enlargement of neck circumference from nerve hypertrophy in CIDP. Neurology, 2016, 87, 442-442.	1.5	1
38	Serpiginous cranial arterial calcification in uremia. Neurology, 2017, 89, 1530-1531.	1.5	1
39	Teaching Video Neurolmages: Rippling muscle disease with caveolin myopathy. Neurology, 2018, 91, e1726-e1727.	1.5	1
40	Intermittent undulating tongue as an involuntary movement in early amyotrophic lateral sclerosis. Parkinsonism and Related Disorders, 2019, 67, 1-2.	1.1	1
41	Myofibrillar Myopathy Mimicking Polyneuropathy. Case Reports in Neurology, 2020, 12, 97-102.	0.3	1
42	Trigeminal-Abducens Pseudosynkinesis: Distinguishing Unconscious Motor Habits From True Synkinesis. Journal of Neuro-Ophthalmology, 2021, 41, e337-e338.	0.4	1
43	Systematic analysis of clinical deficits in unilateral hypoglossal nerve palsy. Muscle and Nerve, 2016, 54, 1055-1058.	1.0	O
44	Teaching Video Neuro <i>Images</i> : Trapezius muscle hypertrophy in multifocal motor neuropathy. Neurology, 2017, 89, e81-e82.	1.5	0
45	Intraneural Ganglion Cysts of the Fibular Nerve: A Cause of Fluctuating Painful Foot Drop. Canadian Journal of Neurological Sciences, 2018, 45, 601-603.	0.3	O
46	Cerebrospinal Fluid in Posterior Reversible Encephalopathy Syndrome. Neurohospitalist, The, 2019, 9, 125-125.	0.3	0
47	Dataset for worldwide survey of cerebrospinal total protein upper reference values. Data in Brief, 2019, 23, 103760.	0.5	O
48	Vertebral Ischemic Necrosis in Diabetic Lumbosacral Radiculoplexus Neuropathy. Diabetes Care, 2021, 44, e53-e54.	4.3	0
49	Does Diabetes Alter CSF Total Protein Levels? A Retrospective Cohort Study. Neurohospitalist, The, 0, , 194187442110393.	0.3	O
50	Pseudohypertrophy of the extensor digitorum brevis in diabetic polyneuropathy. Muscle and Nerve, 2021, 64, E20-E22.	1.0	0
51	MuSK not MNGIE: Atypical MuSK-antibody myasthenia presenting as a genetic disorder. Neuromuscular Disorders, 2021, , .	0.3	0
52	Multifocal acquired demyelinating sensory and motor neuropathy presenting with a unilateral radial neuropathy. Muscle and Nerve, 2022, 65, .	1.0	0