Anna Potulska-Chromik

List of Publications by Year
in descending order

27 papers	685 citations	623574 14 h-index	610775 24 g-index
27 all docs	27 docs citations	27 times ranked	1197 citing authors

#	Article	IF	Citations
1	Swallowing disorders in Parkinson's disease. Parkinsonism and Related Disorders, 2003, 9, 349-353.	1.1	179
2	Novel A18T and pA29S substitutions in Î±-synuclein may be associated with sporadic Parkinson's disease. Parkinsonism and Related Disorders, 2013, 19, 1057-1060.	1.1	63
3	BAG3-related myopathy, polyneuropathy and cardiomyopathy with long QT syndrome. Journal of Muscle Research and Cell Motility, 2015, 36, 423-432.	0.9	57
4	Detection of variants in dystroglycanopathy-associated genes through the application of targeted whole-exome sequencing analysis to a large cohort of patients with unexplained limb-girdle muscle weakness. Skeletal Muscle, 2018, 8, 23.	1.9	40
5	Whole-exome sequencing identifies novel pathogenic mutations and putative phenotype-influencing variants in Polish limb-girdle muscular dystrophy patients. Human Genomics, 2018, 12, 34.	1.4	39
6	A novel dominant D109A CRYAB mutation in a family with myofibrillar myopathy affects Î±B-crystallin structure. BBA Clinical, 2017, 7, 1-7.	4.1	36
7	Efficacy and safety of abobotulinumtoxinA liquid formulation in cervical dystonia: A randomizedâ€controlled trial. Movement Disorders, 2016, 31, 1649-1657.	2.2	35
8	Genomic instability in the PARK2 locus is associated with Parkinsonâ€™s disease. Journal of Applied Genetics, 2015, 56, 451-461.	1.0	27
9	Controlling sialorrhoea: a review of available treatment options. Expert Opinion on Pharmacotherapy, 2005, 6, 1551-1554.	0.9	26
10	Clinical, electrophysiological, and molecular findings in early onset hereditary neuropathy with liability to pressure palsy. Muscle and Nerve, 2014, 50, 914-918.	1.0	21
11	The LITAF/SIMPLE I92V sequence variant results in an earlier age of onset of CMT1A/HNPP diseases. Neurogenetics, 2015, 16, 27-32.	0.7	20
12	Carpal Tunnel Syndrome in Children. Journal of Child Neurology, 2014, 29, 227-231.	0.7	17
13	Andersenâ€“Tawil syndrome: Report of 3 novel mutations and high risk of symptomatic cardiac involvement. Muscle and Nerve, 2015, 51, 192-196.	1.0	16
14	A new missense GDAP1 mutation disturbing targeting to the mitochondrial membrane causes a severe form of AR-CMT2C disease. Neurogenetics, 2011, 12, 145-153.	0.7	15
15	Incidence of mutations in the PARK2, PINK1, PARK7 genes in Polish early-onset Parkinson disease patients. Neurologia I Neurochirurgia Polska, 2013, 47, 319-324.	0.6	13
16	Dopa-responsive dystonia or early-onset Parkinson disease â€“ Genotypeâ€“phenotype correlation. Neurologia I Neurochirurgia Polska, 2017, 51, 1-6.	0.6	13
17	Two Desmin Gene Mutations Associated with Myofibrillar Myopathies in Polish Families. PLoS ONE, 2014, 9, e115470.	1.1	12
18	Exome sequencing reveals mutations in <scp>MFN2</scp> and <scp>GDAP1</scp> in severe Charcotâ€“Marieâ€“Tooth disease. Journal of the Peripheral Nervous System, 2014, 19, 242-245.	1.4	12

#	Article	IF	Citations
19	Are electrophysiological criteria useful in distinguishing childhood demyelinating neuropathies?. Journal of the Peripheral Nervous System, 2016, 21, 22-26.	1.4	12
20	Long lasting dysautonomia due to botulinum toxin B poisoning: clinical- laboratory follow up and difficulties in initial diagnosis. BMC Research Notes, 2013, 6, 438.	0.6	10
21	Original article Clinical and neuroimaging correlation of movement disorders in multiple sclerosis: case series and review of the literature. Folia Neuropathologica, 2014, 1, 92-100.	0.5	7
22	Abnormal spontaneous activity in primary myopathic disorders. Muscle and Nerve, 2017, 56, 427-432.	1.0	6
23	Original article Charcot-Marie-Tooth type 1C disease coexisting with progressive multiple sclerosis: a study of an overlapping syndrome. Folia Neuropathologica, 2012, 4, 369-374.	0.5	4
24	Scintigraphic Evaluation of Mild to Moderate Dysphagia in Motor Neuron Disease. Clinical Nuclear Medicine, 2016, 41, e175-e180.	0.7	4
25	Parametric analysis of pilot voice signals in Parkinson?s disease diagnostics. Journal of Automation Electronics and Electrical Engineering, 2022, 4, 21-28.	0.2	1
26	A review of functional assessment scales in non-sitters with spinal muscular atrophy (SMA). , 2018, 27, 11-17.		0
27	Analysis of hand and face images for the purpose of engineering support for Parkinson's disease diagnosis. Journal of Automation Electronics and Electrical Engineering, 2022, 4, 13-20.	0.2	0