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	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
2	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
3	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
4	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
5	A review of functional assessment scales in non-sitters with spinal muscular atrophy (SMA). , 2018, 27, 11-17.		0
6	Abnormal spontaneous activity in primary myopathic disorders. Muscle and Nerve, 2017, 56, 427-432.	1.0	6
7	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
8	Dopa-responsive dystonia or early-onset Parkinson disease â€“ Genotypeâ€“phenotype correlation. Neurologia I Neurochirurgia Polska, 2017, 51, 1-6.	0.6	13
9	Efficacy and safety of abobotulinumtoxinA liquid formulation in cervical dystonia: A randomizedâ€controlled trial. Movement Disorders, 2016, 31, 1649-1657.	2.2	35
10	Are electrophysiological criteria useful in distinguishing childhood demyelinating neuropathies?. Journal of the Peripheral Nervous System, 2016, 21, 22-26.	1.4	12
11	Scintigraphic Evaluation of Mild to Moderate Dysphagia in Motor Neuron Disease. Clinical Nuclear Medicine, 2016, 41, e175-e180.	0.7	4
12	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
13	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
14	Genomic instability in the PARK2 locus is associated with Parkinsonâ€™s disease. Journal of Applied Genetics, 2015, 56, 451-461.	1.0	27
15	BAG3-related myopathy, polyneuropathy and cardiomyopathy with long QT syndrome. Journal of Muscle Research and Cell Motility, 2015, 36, 423-432.	0.9	57
16	Two Desmin Gene Mutations Associated with Myofibrillar Myopathies in Polish Families. PLoS ONE, 2014, 9, e115470.	1.1	12
17	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
18	Carpal Tunnel Syndrome in Children. Journal of Child Neurology, 2014, 29, 227-231.	0.7	17

#	Article	IF	Citations
19	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
20	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
21	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
22	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
23	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
24	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
25	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
26	Controlling sialorrhoea: a review of available treatment options. Expert Opinion on Pharmacotherapy, 2005, 6, 1551-1554.	0.9	26
27	Swallowing disorders in Parkinson's disease. Parkinsonism and Related Disorders, 2003, 9, 349-353.	1.1	179