GEORGIA KARADIMA

List of Publications by Year
in descending order

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		759233	580821
47	725	12	25
papers	citations	h-index	g-index

47	47	47	1357
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	DNA repair pathways underlie a common genetic mechanism modulating onset in polyglutamine diseases. Annals of Neurology, 2016, 79, 983-990.	5.3	183
2	Apolipoprotein E polymorphism in the Greek population. Clinical Genetics, 1997, 52, 216-218.	2.0	44
3	Hereditary spastic paraplegia in Greece: characterisation of a previously unexplored population using next-generation sequencing. European Journal of Human Genetics, 2016, 24, 857-863.	2.8	43
4	C9ORF72 hexanucleotide repeat expansions are a frequent cause of Huntington disease phenocopies in the Greek population. Neurobiology of Aging, 2015, 36, 547.e13-547.e16.	3.1	38
5	APOE Â4 is associated with impaired verbal learning in patients with MS. Neurology, 2007, 68, 546-549.	1.1	35
6	Huntington's disease in Greece: the experience of 14 years. Clinical Genetics, 2011, 80, 586-590.	2.0	35
7	Late-onset Huntington's disease: Diagnostic and prognostic considerations. Parkinsonism and Related Disorders, 2014, 20, 726-730.	2.2	27
8	Replication study of GWAS risk loci in Greek multiple sclerosis patients. Neurological Sciences, 2019, 40, 253-260.	1.9	24
9	Phenotypic discordance in a pair of monozygotic twins with Huntingtonâ€™s disease. Clinical Genetics, 2008, 74, 291-292.	2.0	22
10	Genetic screening of Greek patients with Huntington's disease phenocopies identifies an SCA8 expansion. Journal of Neurology, 2012, 259, 1874-1878.	3.6	20
11	An APOA1 promoter polymorphism is associated with cognitive performance in patients with multiple sclerosis. Multiple Sclerosis Journal, 2009, 15, 174-179.	3.0	16
12	Four novel connexin 32 mutations in X-linked Charcotâ€“Marieâ€“Tooth disease. Phenotypic variability and central nervous system involvement. Journal of the Neurological Sciences, 2014, 341, 158-161.	0.6	15
13	Friedreich's ataxia and other hereditary ataxias in Greece: An 18-year perspective. Journal of the Neurological Sciences, 2014, 336, 87-92.	0.6	13
14	The challenge of juvenile Huntington disease. Neurology, 2013, 80, 990-996.	1.1	12
15	Biallelic <scp>RFC1</scp> pentanucleotide repeat expansions in Greek patients with lateâ€onset ataxia. Clinical Genetics, 2021, 100, 90-94.	2.0	11
16	Mutational analysis of PMP22, GJB1 and MPZ in Greek Charcot-Marie-Tooth type 1 neuropathy patients. Clinical Genetics, 2011, 80, 497-499.	2.0	10
17	Age at onset in Huntington's disease: Replication study on the association of HAP1. Parkinsonism and Related Disorders, 2012, 18, 1027-1028.	2.2	10
18	The different faces of the p. A53T alpha-synuclein mutation: A screening of Greek patients with parkinsonism and/or dementia. Neuroscience Letters, 2018, 672, 136-139.	2.1	10

#	Article	IF	Citations
19	Complex distal 10q rearrangement in a girl with mild intellectual disability: Follow up of the patient and review of the literature of nonâ€acrocentric satellited chromosomes. American Journal of Medical Genetics, Part A, 2011, 155, 2841-2854.	1.2	9
20	Hereditary Neuropathy Unmasked by Levofloxacin. Annals of Pharmacotherapy, 2011, 45, 1312-1313.	1.9	9
21	Novel peripheral myelin protein 22 (PMP22) micromutations associated with variable phenotypes in Greek patients with Charcot-Marie-Tooth disease. Brain, 2012, 135, e217-e217.	7.6	9
22	Expanding the Spectrum of <scp>AP5Z1â€</scp>Related Hereditary Spastic Paraplegia (<scp>HSPâ€SPG48</scp>): A Multicenter Study on a Rare Disease. Movement Disorders, 2021, 36, 1034-1038.	3.9	9
23	New mutation of theMPZ gene in a family with the Dejerineâ€“Sottas disease phenotype. Muscle and Nerve, 2007, 35, 667-669.	2.2	8
24	APOE genotypes in Greek multiple sclerosis patients: no effect on the MS Severity Score. Journal of Neurology, 2007, 254, 394-395.	3.6	8
25	A novel ABCD1 mutation detected by next generation sequencing in presumed hereditary spastic paraplegia: A 30-year diagnostic delay caused by misleading biochemical findings. Journal of the Neurological Sciences, 2015, 355, 199-201.	0.6	8
26	Mutational screening of the SH3TC2 gene in Greek patients with suspected demyelinating recessive Charcotâ€Marieâ€Tooth disease reveals a varied and unusual phenotypic spectrum. Journal of the Peripheral Nervous System, 2019, 24, 125-130.	3.1	8
27	X linked Charcot-Marie-Tooth disease and multiple sclerosis: emerging evidence for an association. Journal of Neurology, Neurosurgery and Psychiatry, 2019, 90, 187-194.	1.9	8
28	Four novel connexin 32 mutations in Xâ€“linked Charcotâ€“Marieâ€“Tooth disease with phenotypic variability. Journal of Neurology, 2006, 253, 263-264.	3.6	7
29	The rs10492972 KIF1B polymorphism and disease progression in Greek patients with multiple sclerosis. Journal of Neurology, 2011, 258, 1726-1728.	3.6	7
30	Charcot-Marie-Tooth disease type 2C and scapuloperoneal muscular atrophy overlap syndrome in a patient with the R232C TRPV4 mutation. Journal of Neurology, 2015, 262, 1972-1975.	3.6	7
31	Three new case reports of Arteriovenous malformation-related Amyotrophic Lateral Sclerosis. Journal of the Neurological Sciences, 2018, 393, 58-62.	0.6	7
32	Elevated Serum Î±-Synuclein Levels in Huntingtonâ€™s Disease Patients. Neuroscience, 2020, 431, 34-39.	2.3	7
33	Screening for spinocerebellar ataxia type 36 (SCA36) in the Greek population. Journal of the Neurological Sciences, 2019, 402, 131-132.	0.6	6
34	Lack of genetic association between the phospholipase A2 gene and bipolar mood disorder in a European multicentre caseâ€“control study. Psychiatric Genetics, 2006, 16, 169-171.	1.1	5
35	Screening for the C9ORF72 repeat expansion in a greek frontotemporal dementia cohort. Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration, 2018, 19, 152-154.	1.7	5
36	A homozygous GDAP2 loss-of-function variant in a patient with adult-onset cerebellar ataxia. Brain, 2020, 143, e49-e49.	7.6	5

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37	Disentangling balance impairments in spinal and bulbar muscular atrophy. Neuroscience Letters, 2019, 705, 94-98.	2.1	4
38	Co-segregation of Huntington Disease and Hereditary Spastic Paraplegia in 4 Generations. Neurologist, 2011, 17, 211-212.	0.7	3
39	Mutational analysis of Greek patients with suspected hereditary neuropathy with liability to pressure palsies (<scp>HNPP</scp>): a 15â€year experience. Journal of the Peripheral Nervous System, 2015, 20, 79-85.	3.1	3
40	Symptomatic striopallidodentate calcinosis (Fahrâ€™s syndrome) in a thalassemic patient with hypoparathyroidism. Annals of Hematology, 2015, 94, 897-899.	1.8	3
41	Association of 5-HTTLPR Polymorphism with the Nursing Diagnoses and the Achievement of Nursing Outcomes in Patients with Major Depression. Issues in Mental Health Nursing, 2017, 38, 798-804.	1.2	3
42	Evidence for Cognitive Deficits in X-Linked Charcot-Marie-Tooth Disease. Journal of the International Neuropsychological Society, 2020, 26, 294-302.	1.8	3
43	Mutational analysis of <scp>PMP22</scp>, <scp>EGR2</scp>, <scp>LITAF</scp> and <scp>NEFL</scp> in Greek Charcotâ€“Marieâ€“Tooth type 1 patients. Clinical Genetics, 2013, 83, 388-391.	2.0	2
44	Spinobulbar muscular atrophy (Kennedy's disease): A rare diagnosis in the Greek population. Journal of the Neurological Sciences, 2015, 359, 450-451.	0.6	2
45	Bellâ€™s palsy and hereditary neuropathy with liability to pressure palsy (HNPP): Is there a common genetic background?. Journal of Clinical Neuroscience, 2013, 20, 1042.	1.5	1
46	Reevaluation of the <scp>CMT1A</scp> duplication frequency in Greek Charcotâ€Marieâ€Tooth type 1 patients. Clinical Genetics, 2014, 86, 603-603.	2.0	1
47	Complex phenotype in a C9ORF72â€positive patient with highâ€titer antiâ€glutamic acid decarboxylase antibodies: neuroimmunology meets neurogenetics. European Journal of Neurology, 2019, 26, e73-e74.	3.3	0