

Carolina Lemos

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

46
papers

656
citations

15
h-index

23
g-index

53
ext. papers

765
ext. citations

4.3
avg, IF

3.55
L-index

#	Paper	IF	Citations
46	First mutation in the voltage-gated Nav1.1 subunit gene SCN1A with co-occurring familial hemiplegic migraine and epilepsy. <i>Cephalalgia</i> , 2009 , 29, 308-13	6.1	68
45	Ophthalmological manifestations in hereditary transthyretin (ATTR V30M) carriers: a review of 513 cases. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2015 , 22, 117-22	2.7	51
44	Overcoming artefact: anticipation in 284 Portuguese kindreds with familial amyloid polyneuropathy (FAP) ATTRV30M. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2014 , 85, 326-30	5.5	47
43	BDNF and CGRP interaction: implications in migraine susceptibility. <i>Cephalalgia</i> , 2010 , 30, 1375-82	6.1	38
42	Variants in RBP4 and AR genes modulate age at onset in familial amyloid polyneuropathy (FAP ATTRV30M). <i>European Journal of Human Genetics</i> , 2016 , 24, 756-60	5.3	34
41	Impact of liver transplantation on the natural history of oculopathy in Portuguese patients with transthyretin (V30M) amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2015 , 22, 31-5	2.7	23
40	Identification of genetic risk factors for maxillary lateral incisor agenesis. <i>Journal of Dental Research</i> , 2014 , 93, 452-8	8.1	23
39	Familial clustering of migraine: further evidence from a Portuguese study. <i>Headache</i> , 2009 , 49, 404-11	4.2	23
38	Two novel functional mutations in the Na ⁺ ,K ⁺ -ATPase alpha2-subunit ATP1A2 gene in patients with familial hemiplegic migraine and associated neurological phenotypes. <i>Clinical Genetics</i> , 2008 , 73, 37-43	4	23
37	Familial aggregation of maxillary lateral incisor agenesis. <i>Journal of Dental Research</i> , 2010 , 89, 621-5	8.1	21
36	Role of the disease in the psychological impact of pre-symptomatic testing for SCA2 and FAP ATTRV30M: Experience with the disease, kinship and gender of the transmitting parent. <i>Journal of Genetic Counseling</i> , 2009 , 18, 483-93	2.5	20
35	Cellular clearance of circulating transthyretin decreases cell-nonautonomous proteotoxicity in. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018 , 115, E7710-E7719	11.5	19
34	The C677T polymorphism in MTHFR is not associated with migraine in Portugal. <i>Disease Markers</i> , 2008 , 25, 107-13	3.2	19
33	A Trans-acting Factor May Modify Age at Onset in Familial Amyloid Polyneuropathy ATTRV30M in Portugal. <i>Molecular Neurobiology</i> , 2018 , 55, 3676-3683	6.2	15
32	mtDNA copy number associated with age of onset in familial amyloid polyneuropathy. <i>Journal of Neurology, Neurosurgery and Psychiatry</i> , 2018 , 89, 300-304	5.5	15
31	Recurrent ATP1A2 mutations in Portuguese families with familial hemiplegic migraine. <i>Journal of Human Genetics</i> , 2007 , 52, 990-998	4.3	15
30	βTubulin detyrosination impairs mitotic error correction by suppressing MCAK centromeric activity. <i>Journal of Cell Biology</i> , 2020 , 219,	7.3	15

29	Assessing risk factors for migraine: differences in gender transmission. <i>PLoS ONE</i> , 2012 , 7, e50626	3.7	14
28	Interaction between GABA _A receptor genes: new evidence in migraine susceptibility. <i>PLoS ONE</i> , 2013 , 8, e74087	3.7	14
27	Comparison between the Visual Analog Scale and the Numerical Rating Scale in the perception of esthetics and pain. <i>International Orthodontics</i> , 2017 , 15, 543-560	0.9	13
26	Familial hemiplegic migraine due to L263V SCN1A mutation: discordance for epilepsy between two kindreds from Douro Valley. <i>Cephalalgia</i> , 2014 , 34, 1015-20	6.1	13
25	Psychological follow-up of presymptomatic genetic testing for spinocerebellar ataxia type 2 (SCA2) in Cuba. <i>Journal of Genetic Counseling</i> , 2007 , 16, 469-79	2.5	13
24	A role for endothelin receptor type A in migraine without aura susceptibility? A study in Portuguese patients. <i>European Journal of Neurology</i> , 2011 , 18, 649-55	6	12
23	Twenty Years of a Pre-Symptomatic Testing Protocol for Late-Onset Neurological Diseases in Portugal. <i>Acta Medica Portuguesa</i> , 2019 , 32, 295-304	1.4	12
22	Canine mammary tumor risk is associated with polymorphisms in RAD51 and STK11 genes. <i>Journal of Veterinary Diagnostic Investigation</i> , 2018 , 30, 733-738	1.5	10
21	Intergenerational instability in Huntington disease: extreme repeat changes among 134 transmissions. <i>Movement Disorders</i> , 2012 , 27, 583-5	7	10
20	Evidence of syntaxin 1A involvement in migraine susceptibility: a Portuguese study. <i>Archives of Neurology</i> , 2010 , 67, 422-7		10
19	Common symptoms of temporomandibular disorders do not mean same treatment plans: A case series. <i>International Orthodontics</i> , 2018 , 16, 174-214	0.9	8
18	Familial aggregation of cluster headache. <i>Arquivos De Neuro-Psiquiatria</i> , 2013 , 71, 866-70	1.6	8
17	Large normal alleles of ATXN2 decrease age at onset in transthyretin familial amyloid polyneuropathy Val30Met patients. <i>Annals of Neurology</i> , 2019 , 85, 251-258	9.4	8
16	and modify age-at-onset in familial amyloid polyneuropathy patients. <i>Annals of Clinical and Translational Neurology</i> , 2019 , 6, 748-754	5.3	7
15	Familial amyloid polyneuropathy in Portugal: New genes modulating age-at-onset. <i>Annals of Clinical and Translational Neurology</i> , 2017 , 4, 98-105	5.3	6
14	Craniofacial repercussions in maxillary lateral incisors agenesis. <i>International Orthodontics</i> , 2011 , 9, 274-859		5
13	Long-Term Follow-Up of Advanced Liver Disease after Sustained Virological Response to Treatment of Hepatitis C with Direct-Acting Antivirals: Outcomes from a Real-World Portuguese Cohort. <i>GE Portuguese Journal of Gastroenterology</i> , 2020 , 27, 149-159	1.1	4
12	Going Deep into Synaptic Vesicle Machinery Genes and Migraine Susceptibility - A Case-Control Association Study. <i>Headache</i> , 2020 , 60, 2152-2165	4.2	4

11	Monozygotic twin sisters discordant for familial hemiplegic migraine. <i>Journal of Headache and Pain</i> , 2013 , 14, 77	8.8	2
10	Dental repercussions of maxillary lateral incisor agenesis. <i>European Journal of Orthodontics</i> , 2012 , 34, 698-703	3.3	2
9	A genetic interaction of NRXN2 with GABRE, SYT1 and CASK in migraine patients: a case-control study. <i>Journal of Headache and Pain</i> , 2021 , 22, 57	8.8	2
8	CD44v6 expression is a novel predictive marker of therapy response and poor prognosis in gastric cancer patients		1
7	Female preponderance in genetic generalized epilepsies. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2021 , 91, 167-171	3.2	1
6	Beyond Val30Met transthyretin (TTR): variants associated with age-at-onset in hereditary ATTRv amyloidosis. <i>Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis</i> , 2021 , 28, 100-106	2.7	1
5	Genetic overlap between temporomandibular disorders and primary headaches: A systematic review.. <i>Japanese Dental Science Review</i> , 2022 , 58, 69-88	6.8	1
4	A review of migraine genetics: gathering genomic and transcriptomic factors. <i>Human Genetics</i> , 2021 , 141, 1	6.3	0
3	Comparison of East-Asia and West-Europe cohorts explains disparities in survival outcomes and highlights predictive biomarkers of early gastric cancer aggressiveness. <i>International Journal of Cancer</i> , 2021 , 150, 868	7.5	0
2	APOEε4-TOMM40L Haplotype Increases the Risk of Mild Cognitive Impairment Conversion to Alzheimer's Disease. <i>Journal of Alzheimer's Disease</i> , 2020 , 78, 587-601	4.3	
1	A High Methylation Level of a Novel 84 bp CpG Island in the RAMP1 Gene Promoter Is Potentially Associated with Migraine in Women. <i>Brain Sciences</i> , 2022 , 12, 526	3.4	