Carolina Lemos

List of Publications by Year in Descending Order

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	Genetic overlap between temporomandibular disorders and primary headaches: A systematic review.. *Japanese Dental Science Review*, **2022**, 58, 69-88	6.8	1
45	A High Methylation Level of a Novel 284 bp CpG Island in the RAMP1 Gene Promoter Is Potentially Associated with Migraine in Women. *Brain Sciences*, **2022**, 12, 526	3.4	
44	A review of migraine genetics: gathering genomic and transcriptomic factors. *Human Genetics*, **2021**, 141, 1	6.3	0
43	Comparison of East-Asia and West-Europe cohorts explains disparities in survival outcomes and highlights predictive biomarkers of early gastric cancer aggressiveness. *International Journal of Cancer*, **2021**, 150, 868	7.5	0
42	A genetic interaction of NRXN2 with GABRE, SYT1 and CASK in migraine patients: a case-control study. *Journal of Headache and Pain*, **2021**, 22, 57	8.8	2
41	Female preponderance in genetic generalized epilepsies. *Seizure: the Journal of the British Epilepsy Association*, **2021**, 91, 167-171	3.2	1
40	Beyond Val30Met transthyretin (TTR): variants associated with age-at-onset in hereditary ATTRv amyloidosis. *Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis*, **2021**, 28, 100-106	2.7	1
39	Tubulin detyrosination impairs mitotic error correction by suppressing MCAK centromeric activity. *Journal of Cell Biology*, **2020**, 219,	7.3	15
38	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. *GE Portuguese Journal of Gastroenterology*, **2020**, 27, 149-159	1.1	4
37	Going Deep into Synaptic Vesicle Machinery Genes and Migraine Susceptibility - A Case-Control Association Study. *Headache*, **2020**, 60, 2152-2165	4.2	4
36	APOEe4-TOMM40L Haplotype Increases the Risk of Mild Cognitive Impairment Conversion to Alzheimer's Disease. *Journal of Alzheimer's Disease*, **2020**, 78, 587-601	4.3	
35	and modify age-at-onset in familial amyloid polyneuropathy patients. *Annals of Clinical and Translational Neurology*, **2019**, 6, 748-754	5.3	7
34	Twenty Years of a Pre-Symptomatic Testing Protocol for Late-Onset Neurological Diseases in Portugal. *Acta Medica Portuguesa*, **2019**, 32, 295-304	1.4	12
33	Large normal alleles of ATXN2 decrease age at onset in transthyretin familial amyloid polyneuropathy Val30Met patients. *Annals of Neurology*, **2019**, 85, 251-258	9.4	8
32	A Trans-acting Factor May Modify Age at Onset in Familial Amyloid Polyneuropathy ATTRV30M in Portugal. *Molecular Neurobiology*, **2018**, 55, 3676-3683	6.2	15
31	Common symptoms of temporomandibular disorders do not mean same treatment plans: A case series. *International Orthodontics*, **2018**, 16, 174-214	0.9	8
30	mtDNA copy number associated with age of onset in familial amyloid polyneuropathy. *Journal of Neurology, Neurosurgery and Psychiatry*, **2018**, 89, 300-304	5.5	15

29	Cellular clearance of circulating transthyretin decreases cell-nonautonomous proteotoxicity in. *Proceedings of the National Academy of Sciences of the United States of America*, **2018**, 115, E7710-E7719	11.5	19
28	Canine mammary tumor risk is associated with polymorphisms in RAD51 and STK11 genes. *Journal of Veterinary Diagnostic Investigation*, **2018**, 30, 733-738	1.5	10
27	Familial amyloid polyneuropathy in Portugal: New genes modulating age-at-onset. *Annals of Clinical and Translational Neurology*, **2017**, 4, 98-105	5.3	6
26	Comparison between the Visual Analog Scale and the Numerical Rating Scale in the perception of esthetics and pain. *International Orthodontics*, **2017**, 15, 543-560	0.9	13
25	Variants in RBP4 and AR genes modulate age at onset in familial amyloid polyneuropathy (FAP ATTRV30M). *European Journal of Human Genetics*, **2016**, 24, 756-60	5.3	34
24	Impact of liver transplantation on the natural history of oculopathy in Portuguese patients with transthyretin (V30M) amyloidosis. *Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis*, **2015**, 22, 31-5	2.7	23
23	Ophthalmological manifestations in hereditary transthyretin (ATTR V30M) carriers: a review of 513 cases. *Amyloid: the International Journal of Experimental and Clinical Investigation: the Official Journal of the International Society of Amyloidosis*, **2015**, 22, 117-22	2.7	51
22	Overcoming artefact: anticipation in 284 Portuguese kindreds with familial amyloid polyneuropathy (FAP) ATTRV30M. *Journal of Neurology, Neurosurgery and Psychiatry*, **2014**, 85, 326-30	5.5	47
21	Identification of genetic risk factors for maxillary lateral incisor agenesis. *Journal of Dental Research*, **2014**, 93, 452-8	8.1	23
20	Familial hemiplegic migraine due to L263V SCN1A mutation: discordance for epilepsy between two kindreds from Douro Valley. *Cephalalgia*, **2014**, 34, 1015-20	6.1	13
19	Monozygotic twin sisters discordant for familial hemiplegic migraine. *Journal of Headache and Pain*, **2013**, 14, 77	8.8	2
18	Familial aggregation of cluster headache. *Arquivos De Neuro-Psiquiatria*, **2013**, 71, 866-70	1.6	8
17	Interaction between γ-aminobutyric acid A receptor genes: new evidence in migraine susceptibility. *PLoS ONE*, **2013**, 8, e74087	3.7	14
16	Intergenerational instability in Huntington disease: extreme repeat changes among 134 transmissions. *Movement Disorders*, **2012**, 27, 583-5	7	10
15	Assessing risk factors for migraine: differences in gender transmission. *PLoS ONE*, **2012**, 7, e50626	3.7	14
14	Dental repercussions of maxillary lateral incisor agenesis. *European Journal of Orthodontics*, **2012**, 34, 698-703	3.3	2
13	A role for endothelin receptor type A in migraine without aura susceptibility? A study in Portuguese patients. *European Journal of Neurology*, **2011**, 18, 649-55	6	12
12	Craniofacial repercussions in maxillary lateral incisors agenesis. *International Orthodontics*, **2011**, 9, 274-85	0.9	5

11	Familial aggregation of maxillary lateral incisor agenesis. *Journal of Dental Research*, **2010**, 89, 621-5	8.1	21
10	BDNF and CGRP interaction: implications in migraine susceptibility. *Cephalalgia*, **2010**, 30, 1375-82	6.1	38
9	Evidence of syntaxin 1A involvement in migraine susceptibility: a Portuguese study. *Archives of Neurology*, **2010**, 67, 422-7		10
8	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. *Journal of Genetic Counseling*, **2009**, 18, 483-93	2.5	20
7	Familial clustering of migraine: further evidence from a Portuguese study. *Headache*, **2009**, 49, 404-11	4.2	23
6	First mutation in the voltage-gated Nav1.1 subunit gene SCN1A with co-occurring familial hemiplegic migraine and epilepsy. *Cephalalgia*, **2009**, 29, 308-13	6.1	68
5	The C677T polymorphism in MTHFR is not associated with migraine in Portugal. *Disease Markers*, **2008**, 25, 107-13	3.2	19
4	Two novel functional mutations in the Na+,K+-ATPase alpha2-subunit ATP1A2 gene in patients with familial hemiplegic migraine and associated neurological phenotypes. *Clinical Genetics*, **2008**, 73, 37-43	4	23
3	Psychological follow-up of presymptomatic genetic testing for spinocerebellar ataxia type 2 (SCA2) in Cuba. *Journal of Genetic Counseling*, **2007**, 16, 469-79	2.5	13
2	Recurrent ATP1A2 mutations in Portuguese families with familial hemiplegic migraine. *Journal of Human Genetics*, **2007**, 52, 990-998	4.3	15
1	CD44v6 expression is a novel predictive marker of therapy response and poor prognosis in gastric cancer patients		1