Jos Maria Pereira-Monteiro

List of Publications by Year in Descending Order

20 papers	628 citations	11 h-index	21 g-index
21 ext. papers	687 ext. citations	8.9 avg, IF	2.79 L-index

#	Paper	IF	Citations
20	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
19	Going Deep into Synaptic Vesicle Machinery Genes and Migraine Susceptibility - A Case-Control Association Study. *Headache*, **2020**, 60, 2152-2165	4.2	4
18	SUNCT syndrome: A cohort of 15 Portuguese patients. *Cephalalgia*, **2016**, 36, 1002-6	6.1	13
17	CADASIL: MRI may be normal in the fourth decade of life - a case report. *Cephalalgia*, **2016**, 36, 1082-1085	6.1	5
16	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
15	Monozygotic twin sisters discordant for familial hemiplegic migraine. *Journal of Headache and Pain*, **2013**, 14, 77	8.8	2
14	Cerebellar ataxia, hemiplegic migraine, and related phenotypes due to a CACNA1A missense mutation: 12-year follow-up of a large Portuguese family. *JAMA Neurology*, **2013**, 70, 235-40	17.2	22
13	Interaction between □aminobutyric acid A receptor genes: new evidence in migraine susceptibility. *PLoS ONE*, **2013**, 8, e74087	3.7	14
12	Sporadic hemiplegic migraine with normal imaging as the initial manifestation of CADASIL. *Cephalalgia*, **2012**, 32, 255-7	6.1	7
11	Psychotic aura symptoms in familial hemiplegic migraine type 2 (ATP1A2). *Journal of Headache and Pain*, **2012**, 13, 581-5	8.8	8
10	Assessing risk factors for migraine: differences in gender transmission. *PLoS ONE*, **2012**, 7, e50626	3.7	14
9	Migraine-induced epistaxis and sporadic hemiplegic migraine: unusual features in the same patient. *Case Reports in Neurology*, **2012**, 4, 116-9	1	4
8	A dominant-negative mutation in the TRESK potassium channel is linked to familial migraine with aura. *Nature Medicine*, **2010**, 16, 1157-60	50.5	263
7	BDNF and CGRP interaction: implications in migraine susceptibility. *Cephalalgia*, **2010**, 30, 1375-82	6.1	38
6	Evidence of syntaxin 1A involvement in migraine susceptibility: a Portuguese study. *Archives of Neurology*, **2010**, 67, 422-7		10
5	Guidelines for telematic second opinion consultation on headaches in Europe: on behalf of the European Headache Federation (EHF). *Journal of Headache and Pain*, **2010**, 11, 345-8	8.8	9
4	Familial clustering of migraine: further evidence from a Portuguese study. *Headache*, **2009**, 49, 404-11	4.2	23

List of Publications

3	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
2	Divergent sodium channel defects in familial hemiplegic migraine. *Proceedings of the National Academy of Sciences of the United States of America*, **2008**, 105, 9799-804	11.5	86
1	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