

# Jos Maria Pereira-Monteiro

## 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.

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 dominant-negative mutation in the TRESK potassium channel is linked to familial migraine with aura. <i>Nature Medicine</i> , <b>2010</b> , 16, 1157-60	50.5	263
19	Divergent sodium channel defects in familial hemiplegic migraine. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2008</b> , 105, 9799-804	11.5	86
18	First mutation in the voltage-gated Nav1.1 subunit gene SCN1A with co-occurring familial hemiplegic migraine and epilepsy. <i>Cephalalgia</i> , <b>2009</b> , 29, 308-13	6.1	68
17	BDNF and CGRP interaction: implications in migraine susceptibility. <i>Cephalalgia</i> , <b>2010</b> , 30, 1375-82	6.1	38
16	Familial clustering of migraine: further evidence from a Portuguese study. <i>Headache</i> , <b>2009</b> , 49, 404-11	4.2	23
15	Two novel functional mutations in the Na <sup>+</sup> ,K <sup>+</sup> -ATPase alpha2-subunit ATP1A2 gene in patients with familial hemiplegic migraine and associated neurological phenotypes. <i>Clinical Genetics</i> , <b>2008</b> , 73, 37-43	4	23
14	Cerebellar ataxia, hemiplegic migraine, and related phenotypes due to a CACNA1A missense mutation: 12-year follow-up of a large Portuguese family. <i>JAMA Neurology</i> , <b>2013</b> , 70, 235-40	17.2	22
13	Assessing risk factors for migraine: differences in gender transmission. <i>PLoS ONE</i> , <b>2012</b> , 7, e50626	3.7	14
12	Interaction between GABA <sub>A</sub> receptor genes: new evidence in migraine susceptibility. <i>PLoS ONE</i> , <b>2013</b> , 8, e74087	3.7	14
11	SUNCT syndrome: A cohort of 15 Portuguese patients. <i>Cephalalgia</i> , <b>2016</b> , 36, 1002-6	6.1	13
10	Familial hemiplegic migraine due to L263V SCN1A mutation: discordance for epilepsy between two kindreds from Douro Valley. <i>Cephalalgia</i> , <b>2014</b> , 34, 1015-20	6.1	13
9	Evidence of syntaxin 1A involvement in migraine susceptibility: a Portuguese study. <i>Archives of Neurology</i> , <b>2010</b> , 67, 422-7		10
8	Guidelines for telematic second opinion consultation on headaches in Europe: on behalf of the European Headache Federation (EHF). <i>Journal of Headache and Pain</i> , <b>2010</b> , 11, 345-8	8.8	9
7	Psychotic aura symptoms in familial hemiplegic migraine type 2 (ATP1A2). <i>Journal of Headache and Pain</i> , <b>2012</b> , 13, 581-5	8.8	8
6	Sporadic hemiplegic migraine with normal imaging as the initial manifestation of CADASIL. <i>Cephalalgia</i> , <b>2012</b> , 32, 255-7	6.1	7
5	CADASIL: MRI may be normal in the fourth decade of life - a case report. <i>Cephalalgia</i> , <b>2016</b> , 36, 1082-1085	1	5
4	Migraine-induced epistaxis and sporadic hemiplegic migraine: unusual features in the same patient. <i>Case Reports in Neurology</i> , <b>2012</b> , 4, 116-9	1	4

3	Going Deep into Synaptic Vesicle Machinery Genes and Migraine Susceptibility - A Case-Control Association Study. <i>Headache</i> , <b>2020</b> , 60, 2152-2165	4.2	4
2	Monozygotic twin sisters discordant for familial hemiplegic migraine. <i>Journal of Headache and Pain</i> , <b>2013</b> , 14, 77	8.8	2
1	A genetic interaction of NRXN2 with GABRE, SYT1 and CASK in migraine patients: a case-control study. <i>Journal of Headache and Pain</i> , <b>2021</b> , 22, 57	8.8	2