Sofia A Oliveira

List of Publications by Year
in descending order

29	2,353	361413	477307
papers	citations	20 h-index	29 g-index

30	30	30	5414
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	<scp>IL</scp>10 lowâ€frequency variants in <scp>B</scp>ehÃ§et's disease patients. International Journal of Rheumatic Diseases, 2017, 20, 622-627.	1.9	3
2	Ulcerative Colitis Is Under Dual (Mitochondrial and Nuclear) Genetic Control. Inflammatory Bowel Diseases, 2016, 22, 774-781.	1.9	9
3	Low-frequency and common genetic variation in ischemic stroke. Neurology, 2016, 86, 1217-1226.	1.1	141
4	Mitochondrial genome association study with peripheral arterial disease and venous thromboembolism. Atherosclerosis, 2016, 252, 97-105.	0.8	5
5	Characterization of the major histocompatibility complex locus association with BehÃ§etâ€™s disease in Iran. Arthritis Research and Therapy, 2015, 17, 81.	3.5	21
6	Brief Report: Association of CCR1, KLRC4, IL12Aâ€“AS1, STAT4, and ERAP1 With BehÃ§et's Disease in Iranians. Arthritis and Rheumatology, 2015, 67, 2742-2748.	5.6	38
7	Gene expression profiling and association studies implicate the neuregulin signaling pathway in BehÃ§et's disease susceptibility. Journal of Molecular Medicine, 2013, 91, 1013-1023.	3.9	12
8	Variants within the nitric oxide synthase 1 gene are associated with stroke susceptibility. Atherosclerosis, 2012, 220, 443-448.	0.8	23
9	TTC7B Emerges as a Novel Risk Factor for Ischemic Stroke Through the Convergence of Several Genome-Wide Approaches. Journal of Cerebral Blood Flow and Metabolism, 2012, 32, 1061-1072.	4.3	86
10	Genetic risk factors for ischaemic stroke and its subtypes (the METASTROKE Collaboration): a meta-analysis of genome-wide association studies. Lancet Neurology, The, 2012, 11, 951-962.	10.2	445
11	Association study of IL10 and IL23Râ€“IL12RB2 in Iranian patients with BehÃ§et's disease. Arthritis and Rheumatism, 2012, 64, 2761-2772.	6.7	76
12	Replication of the CELSR1 association with ischemic stroke in a Portuguese case-control cohort. Atherosclerosis, 2011, 217, 260-262.	0.8	8
13	Variants in the Inflammatory IL6 and MPO Genes Modulate Stroke Susceptibility Through Main Effects and Geneâ€”Gene Interactions. Journal of Cerebral Blood Flow and Metabolism, 2011, 31, 1751-1759.	4.3	19
14	Association of mitochondrial polymorphism m.709G>A with Behcet's disease. Annals of the Rheumatic Diseases, 2011, 70, 1514-1516.	0.9	5
15	Convergence of miRNA Expression Profiling, Î±-Synuclein Interacton and GWAS in Parkinson's Disease. PLoS ONE, 2011, 6, e25443.	2.5	235
16	Kalirin: a novel genetic risk factor for ischemic stroke. Human Genetics, 2010, 127, 513-523.	3.8	51
17	Variants of the Matrix Metalloproteinase-2 but not the Matrix Metalloproteinase-9 genes significantly influence functional outcome after stroke. BMC Medical Genetics, 2010, 11, 40.	2.1	50
18	Association of a Genetic Variant in the <i>ALOX5AP</i> with Higher Risk of Ischemic Stroke: A Case-Control, Meta-Analysis and Functional Study. Cerebrovascular Diseases, 2010, 29, 528-537.	1.7	54

#	Article	IF	Citations
19	Mitochondrial haplogroup H1 is protective for ischemic stroke in Portuguese patients. BMC Medical Genetics, 2008, 9, 57.	2.1	42
20	Revealing the role of glutathione S-transferase omega in age-at-onset of Alzheimer and Parkinson diseases. Neurobiology of Aging, 2006, 27, 1087-1093.	3.1	60
21	Mutations in the pleckstrin homology domain of dynamin 2 cause dominant intermediate Charcot-Marie-Tooth disease. Nature Genetics, 2005, 37, 289-294.	21.4	324
22	Association between the neuron-specific RNA-binding protein ELAVL4 and Parkinson disease. Human Genetics, 2005, 117, 27-33.	3.8	67
23	Identification of Risk and Age-at-Onset Genes on Chromosome 1p in Parkinson Disease. American Journal of Human Genetics, 2005, 77, 252-264.	6.2	67
24	Linkage disequilibrium and haplotype tagging polymorphisms in the Tau H1 haplotype. Neurogenetics, 2004, 5, 147-155.	1.4	30
25	Parkin mutations and susceptibility alleles in lateâ€onset Parkinson's disease. Annals of Neurology, 2003, 53, 624-629.	5.3	201
26	The Q7R Saitohin gene polymorphism is not associated with Alzheimer disease. Neuroscience Letters, 2003, 347, 143-146.	2.1	12
27	Glutathione S-transferase omega-1 modifiesage-at-onset of Alzheimer disease and Parkinson disease. Human Molecular Genetics, 2003, 12, 3259-3267.	2.9	208
28	Association Study of Parkin Gene Polymorphisms With Idiopathic Parkinson Disease. Archives of Neurology, 2003, 60, 975.	4.5	51
29	Dissecting A Complex Disease Using Modern Techniques of Molecular Biology. Laboratory Medicine, 2001, 32, 594-598.	1.2	0