## **Michel Clanet**

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Diagnostic criteria for multiple sclerosis: 2010 Revisions to the McDonald criteria. Annals of Neurology, 2011, 69, 292-302.	5.3	8,001
2	Defining the clinical course of multiple sclerosis. Neurology, 2014, 83, 278-286.	1.1	2,344
3	ECTRIMS/EAN Guideline on the pharmacological treatment of people with multiple sclerosis. Multiple Sclerosis Journal, 2018, 24, 96-120.	3.0	458
4	MD1003 (high-dose biotin) for the treatment of progressive multiple sclerosis: A randomised, double-blind, placebo-controlled study. Multiple Sclerosis Journal, 2016, 22, 1719-1731.	3.0	249
5	Mutation of the <i>PDGFRB</i> gene as a cause of idiopathic basal ganglia calcification. Neurology, 2013, 80, 181-187.	1.1	239
6	Phenotypic spectrum of probable and genetically-confirmed idiopathic basal ganglia calcification. Brain, 2013, 136, 3395-3407.	7.6	183
7	Long-term Outcomes of CLIPPERS (Chronic Lymphocytic Inflammation With Pontine Perivascular) Tj ETQq1 1 0. 2012, 69, 847-55.	784314 rg 4.5	BT /Overlock 109
8	Antivertigo Medications and Drug-Induced Vertigo. Drugs, 1995, 50, 777-791.	10.9	96
9	Tyrosine kinase 2 variant influences T lymphocyte polarization and multiple sclerosis susceptibility. Brain, 2011, 134, 693-703.	7.6	96
10	Diffusion tensor imaging in multiple sclerosis: a tool for monitoring changes in normal-appearing white matter. Multiple Sclerosis Journal, 2004, 10, 188-196.	3.0	71
11	Tumor necrosis factor polymorphisms in multiple sclerosis: No additional association independent of HLA. Journal of Neuroimmunology, 1994, 51, 93-99.	2.3	61
12	A central nervous system B-cell lymphoma arising two years after initial diagnosis of CLIPPERS. Journal of the Neurological Sciences, 2014, 344, 224-226.	0.6	58
13	A Role for <i>VAV1</i> in Experimental Autoimmune Encephalomyelitis and Multiple Sclerosis. Science Translational Medicine, 2009, 1, 10ra21.	12.4	52
14	Biallelic MYORG mutation carriers exhibit primary brain calcification with a distinct phenotype. Brain, 2019, 142, 1573-1586.	7.6	49
15	Genetic interaction of <i>CTLAâ€4</i> with HLAâ€DR15 in multiple sclerosis patients. Annals of Neurology, 2003, 54, 119-122.	5.3	46
16	Cytokines in genetic susceptibility to multiple sclerosis: a candidate gene approach. Journal of Neuroimmunology, 2000, 102, 107-112.	2.3	45
17	Urinary complications and risk factors in symptomatic multiple sclerosis patients. Study of a cohort of 328 patients. Neurourology and Urodynamics, 2015, 34, 32-36.	1.5	37
18	Evidence for Linkage Disequilibrium Between HLA-DRB1 Gene and Multiple Sclerosis. Science, 1997, 276, 661g-665.	12.6	36

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19	Investigation of seven proposed regions of linkage in multiple sclerosis: an American and French collaborative study. Neurogenetics, 2004, 5, 45-48.	1.4	23
20	HLA-DPB1 gene polymorphism and multiple sclerosis: a large case-control study in the southwest of France. Journal of Neuroimmunology, 1991, 34, 215-222.	2.3	10
21	Aquaporin 4 distribution in the brain and its relevance for the radiological appearance of neuromyelitis optica spectrum disease. Journal of Neuroradiology, 2021, 48, 170-175.	1.1	4
22	Relevance of the skewness index in DTI exploration of multiple sclerosis. Magnetic Resonance Materials in Physics, Biology, and Medicine, 2009, 22, 89-100.	2.0	1
23	Assessment of a program to encourage the multidisciplinary management of urinary disorders in multiple sclerosis. Neurourology and Urodynamics, 2017, 36, 706-709.	1.5	1