

Benedicte Dubois

List of Publications by Year in descending order

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Version: 2024-02-01

61
papers

9,091
citations

236612

25
h-index

155451

55
g-index

62
all docs

62
docs citations

62
times ranked

14385
citing authors

#	ARTICLE	IF	CITATIONS
1	Effects of Vitamin D and Body Mass Index on Disease Risk and Relapse Hazard in Multiple Sclerosis. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2022, 9, .	3.1	23
2	Environmental risk factors in multiple sclerosis: bridging Mendelian randomization and observational studies. <i>Journal of Neurology</i> , 2022, 269, 4565-4574.	1.8	14
3	Body Mass Index, Interleukin-6 Signaling and Multiple Sclerosis: A Mendelian Randomization Study. <i>Frontiers in Immunology</i> , 2022, 13, 834644.	2.2	12
4	Neurocysticercosis: An Uncommon Cause of Acute Supratentorial Hydrocephalus. <i>Journal of the Belgian Society of Radiology</i> , 2022, 106, 31.	0.1	0
5	Quantitative MRI phenotypes capture biological heterogeneity in multiple sclerosis patients. <i>Scientific Reports</i> , 2021, 11, 1573.	1.6	5
6	Genetic Variation in <i>WNT9B</i> Increases Relapse Hazard in Multiple Sclerosis. <i>Annals of Neurology</i> , 2021, 89, 884-894.	2.8	12
7	Treatment-Induced BAFF Expression and B Cell Biology in Multiple Sclerosis. <i>Frontiers in Immunology</i> , 2021, 12, 676619.	2.2	6
8	The Multiple Sclerosis Data Alliance Catalogue. <i>International Journal of MS Care</i> , 2021, 23, 261-268.	0.4	3
9	Community-acquired bacterial meningitis in adults: emergency department management protocol. <i>Acta Neurologica Belgica</i> , 2020, 120, 1033-1043.	0.5	0
10	CHIT1 at Diagnosis Reflects Long-Term Multiple Sclerosis Disease Activity. <i>Annals of Neurology</i> , 2020, 87, 633-645.	2.8	22
11	Somatic Variants: New Kids on the Block in Human Immunogenetics. <i>Trends in Genetics</i> , 2019, 35, 935-947.	2.9	29
12	Multiple sclerosis genomic map implicates peripheral immune cells and microglia in susceptibility. <i>Science</i> , 2019, 365, .	6.0	710
13	Leveraging human genetics to inform intervention strategies for multiple sclerosis. <i>Neurology</i> , 2019, 92, 735-736.	1.5	2
14	Horizontal saccadic palsy as a prominent symptom of anti-NMDAR encephalitis. <i>Neurology: Clinical Practice</i> , 2019, 11, 10.1212/CPJ.0000000000000750.	0.8	0
15	A robust pipeline with high replication rate for detection of somatic variants in the adaptive immune system as a source of common genetic variation in autoimmune disease. <i>Human Molecular Genetics</i> , 2019, 28, 1369-1380.	1.4	16
16	Management of immune thrombocytopenia in multiple sclerosis patients treated with alemtuzumab: a Belgian consensus. <i>Acta Neurologica Belgica</i> , 2018, 118, 7-11.	0.5	10
17	Multiple sclerosis risk variants alter expression of co-stimulatory genes in B cells. <i>Brain</i> , 2018, 141, 786-796.	3.7	39
18	Genetic basis for relapse rate in multiple sclerosis: Association with <i>LRP2</i> genetic variation. <i>Multiple Sclerosis Journal</i> , 2018, 24, 1773-1775.	1.4	13

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19	A Belgian consensus protocol for autologous hematopoietic stem cell transplantation in multiple sclerosis. <i>Acta Neurologica Belgica</i> , 2018, 118, 161-168.	0.5	6
20	Siponimod versus placebo in secondary progressive multiple sclerosis (EXPAND): a double-blind, randomised, phase 3 study. <i>Lancet</i> , The, 2018, 391, 1263-1273.	6.3	684
21	Low-Frequency and Rare-Coding Variation Contributes to Multiple Sclerosis Risk. <i>Cell</i> , 2018, 175, 1679-1687.e7.	13.5	115
22	Genetic Architecture of Adaptive Immune System Identifies Key Immune Regulators. <i>Cell Reports</i> , 2018, 25, 798-810.e6.	2.9	36
23	LncRNAs expression profile in peripheral blood mononuclear cells from multiple sclerosis patients. <i>Journal of Neuroimmunology</i> , 2018, 324, 129-135.	1.1	37
24	Corticosteroids in the management of acute multiple sclerosis exacerbations. <i>Acta Neurologica Belgica</i> , 2017, 117, 623-633.	0.5	31
25	New insights into the burden and costs of multiple sclerosis in Europe: Results for Belgium. <i>Multiple Sclerosis Journal</i> , 2017, 23, 29-40.	1.4	12
26	Analysis of Plasminogen Genetic Variants in Multiple Sclerosis Patients. <i>G3: Genes, Genomes, Genetics</i> , 2016, 6, 2073-2079.	0.8	13
27	Immunologic profiles of multiple sclerosis treatments reveal shared early B cell alterations. <i>Neurology: Neuroimmunology and NeuroInflammation</i> , 2016, 3, e240.	3.1	35
28	Role of Genetic Factors in Pathophysiology of Multiple Sclerosis. , 2016, , 153-180.		2
29	Power estimation for non-standardized multisite studies. <i>NeuroImage</i> , 2016, 134, 281-294.	2.1	36
30	Characteristic callosal involvement in Susac's syndrome. <i>Acta Neurologica Belgica</i> , 2015, 115, 395-396.	0.5	7
31	Genetic variants are major determinants of CSF antibody levels in multiple sclerosis. <i>Brain</i> , 2015, 138, 632-643.	3.7	54
32	Burden of risk variants correlates with phenotype of multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 2015, 21, 1670-1680.	1.4	27
33	CCR2 defines in vivo development and homing of IL-23-driven GM-CSF-producing Th17 cells. <i>Nature Communications</i> , 2015, 6, 8644.	5.8	117
34	Class II HLA interactions modulate genetic risk for multiple sclerosis. <i>Nature Genetics</i> , 2015, 47, 1107-1113.	9.4	312
35	No evidence for shared genetic basis of common variants in multiple sclerosis and amyotrophic lateral sclerosis. <i>Human Molecular Genetics</i> , 2014, 23, 1916-1922.	1.4	23
36	HLA-E restricted CD8+ T cell subsets are phenotypically altered in multiple sclerosis patients. <i>Multiple Sclerosis Journal</i> , 2014, 20, 790-801.	1.4	14

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37	Analysis of immune-related loci identifies 48 new susceptibility variants for multiple sclerosis. <i>Nature Genetics</i> , 2013, 45, 1353-1360.	9.4	1,213
38	Rare <i>MEFV</i> variants are not associated with risk to develop multiple sclerosis and severity of disease. <i>Multiple Sclerosis Journal</i> , 2013, 19, 1132-1136.	1.4	6
39	Genetic variants in the immunoglobulin heavy chain locus are associated with the IgG index in multiple sclerosis. <i>Annals of Neurology</i> , 2013, 73, 86-94.	2.8	38
40	Progress in Multiple Sclerosis Genetics. <i>Current Genomics</i> , 2012, 13, 646-663.	0.7	20
41	Genetic risk and a primary role for cell-mediated immune mechanisms in multiple sclerosis. <i>Nature</i> , 2011, 476, 214-219.	13.7	2,400
42	Secondary autoimmune diseases occurring after HSCT for an autoimmune disease: a retrospective study of the EBMT Autoimmune Disease Working Party. <i>Blood</i> , 2011, 118, 1693-1698.	0.6	140
43	<i>TNFRSF1A</i> coding variants in multiple sclerosis. <i>Journal of Neuroimmunology</i> , 2011, 235, 110-112.	1.1	16
44	Replication of <i>KIF21B</i> as a susceptibility locus for multiple sclerosis. <i>Journal of Medical Genetics</i> , 2010, 47, 775-776.	1.5	38
45	The role of the <i>CD58</i> locus in multiple sclerosis. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2009, 106, 5264-5269.	3.3	185
46	Interleukin 7 receptor α chain (<i>IL7R</i>) shows allelic and functional association with multiple sclerosis. <i>Nature Genetics</i> , 2007, 39, 1083-1091.	9.4	578
47	Corrigendum to "Linkage disequilibrium screening for multiple sclerosis implicates <i>JAG1</i> and <i>POU2AF1</i> as susceptibility genes in Europeans" [<i>J. Neuroimmunol.</i> 179 (2006) 108-116]. <i>Journal of Neuroimmunology</i> , 2007, 189, 175-176.	1.1	1
48	Multiple Sclerosis Severity Score. <i>Neurology</i> , 2005, 64, 1144-1151.	1.5	836
49	Corrigendum to "Paxantrone (BBR2778): A new immunosuppressant in multiple sclerosis with a low cardiotoxicity" [<i>Neurol Sci</i> 223 (2004) 81-86]. <i>Journal of the Neurological Sciences</i> , 2005, 235, 79.	0.3	1
50	Gelatinase B, PECAM-1 and MCP-3 gene polymorphisms in Belgian multiple sclerosis. <i>Journal of the Neurological Sciences</i> , 2002, 200, 43-48.	0.3	19
51	Biochemistry and Molecular Biology of Gelatinase B or Matrix Metalloproteinase-9 (MMP-9). <i>Critical Reviews in Biochemistry and Molecular Biology</i> , 2002, 37, 375-536.	2.3	805
52	Gelatinase B deficiency protects against endotoxin shock. <i>European Journal of Immunology</i> , 2002, 32, 2163.	1.6	81
53	Mitochondrial mutations of Leber's hereditary optic neuropathy: a risk factor for multiple sclerosis. <i>Journal of Neurology</i> , 2000, 247, 535-543.	1.8	61
54	In Vivo Neutrophil Recruitment by Granulocyte Chemotactic Protein-2 Is Assisted by Gelatinase B/MMP-9 in the Mouse. <i>Journal of Interferon and Cytokine Research</i> , 2000, 20, 667-674.	0.5	69

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55	Microsatellite polymorphisms in the gene promoter of monocyte chemotactic protein-3 and analysis of the association between monocyte chemotactic protein-3 alleles and multiple sclerosis development. <i>Journal of Neuroimmunology</i> , 1999, 95, 195-201.	1.1	24
56	Regulation of gelatinases in microglia and astrocyte cell cultures by plant lectins. , 1999, 27, 53-61.		20
57	Gelatinase B in multiple sclerosis and experimental autoimmune encephalomyelitis. <i>Acta Neurologica Belgica</i> , 1999, 99, 53-6.	0.5	19
58	The immunological basis of current and novel therapies of multiple sclerosis. <i>Archivum Immunologiae Et Therapiae Experimentalis</i> , 1999, 47, 7-16.	1.0	0
59	Regulation of gelatinase B (MMP-9) in leukocytes by plant lectins. <i>FEBS Letters</i> , 1998, 427, 275-278.	1.3	22
60	Toxicity in a double-blind, placebo-controlled pilot trial with D-penicillamine and metacycline in secondary progressive multiple sclerosis. <i>Multiple Sclerosis Journal</i> , 1998, 4, 74-78.	1.4	18
61	Antiviral treatment with fluoxetine for rituximab-associated chronic echovirus 13 meningoencephalitis and myofasciitis. <i>European Journal of Neurology</i> , 0, , .	1.7	2