## Andrea Harrer

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Tumefactive MS lesions under fingolimod. Neurology, 2013, 81, 1654-1658.	1.5	72
2	Cerebrospinal fluid parameters of B cell-related activity in patients with active disease during natalizumab therapy. Multiple Sclerosis Journal, 2013, 19, 1209-1212.	1.4	69
3	Current therapies in ischemic stroke. Part A. Recent developments in acute stroke treatment and in stroke prevention. Drug Discovery Today, 2012, 17, 296-309.	3.2	59
4	Reshaping the Bet v 1 fold modulates TH polarization. Journal of Allergy and Clinical Immunology, 2011, 127, 1571-1578.e9.	1.5	53
5	Diclofenac Hypersensitivity: Antibody Responses to the Parent Drug and Relevant Metabolites. PLoS ONE, 2010, 5, e13707.	1.1	48
6	Importance of cerebrospinal fluid analysis in the era of McDonald 2010 criteria: a German–Austrian retrospective multicenter study in patients with a clinically isolated syndrome. Journal of Neurology, 2016, 263, 2499-2504.	1.8	46
7	lsoform identification and characterization of Art v 3, the lipid-transfer protein of mugwort pollen. Molecular Immunology, 2009, 46, 1919-1924.	1.0	42
8	Characterization of plant food allergens: An overview on physicochemical and immunological techniques. Molecular Nutrition and Food Research, 2010, 54, 93-112.	1.5	35
9	Glatiramer acetate attenuates the pro-migratory profile of adhesion molecules on various immune cell subsets in multiple sclerosis. Clinical and Experimental Immunology, 2013, 173, 381-389.	1.1	32
10	Adhesion molecules are promising candidates to establish surrogate markers for natalizumab treatment. Multiple Sclerosis Journal, 2011, 17, 16-23.	1.4	30
11	Natalizumab therapy decreases surface expression of both VLA-heterodimer subunits on peripheral blood mononuclear cells. Journal of Neuroimmunology, 2011, 234, 148-154.	1.1	29
12	Circadian rhythmicity of inflammatory serum parameters: a neglected issue in the search of biomarkers in multiple sclerosis. Journal of Neurology, 2013, 260, 221-227.	1.8	28
13	Molecular characterization of Api g 2, a novel allergenic member of the lipidâ€ŧransfer protein 1 family from celery stalks. Molecular Nutrition and Food Research, 2011, 55, 568-577.	1.5	26
14	Current therapies in ischemic stroke. Part B. Future candidates in stroke therapy and experimental studies. Drug Discovery Today, 2012, 17, 671-684.	3.2	25
15	Basophil Reactivity as Biomarker in Immediate Drug Hypersensitivity Reactions—Potential and Limitations. Frontiers in Pharmacology, 2016, 7, 171.	1.6	21
16	High interindividual variability in the CD4/CD8 T cell ratio and natalizumab concentration levels in the cerebrospinal fluid of patients with multiple sclerosis. Clinical and Experimental Immunology, 2015, 180, 383-392.	1.1	19
17	Lymphocyte Subsets Show Different Response Patterns to In Vivo Bound Natalizumab—A Flow Cytometric Study on Patients with Multiple Sclerosis. PLoS ONE, 2012, 7, e31784.	1.1	18
18	Natalizumab saturation: biomarker for individual treatment holiday after natalizumab withdrawal?. Acta Neurologica Scandinavica, 2014, 129, e12-e15.	1.0	18

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19	The Evolution of Human Basophil Biology from Neglect towards Understanding of Their Immune Functions. BioMed Research International, 2016, 2016, 1-16.	0.9	15
20	Cerebrospinal fluid CXLC13 indicates disease course in neuroinfection: an observational study. Journal of Neuroinflammation, 2019, 16, 13.	3.1	14
21	Chemokine CXCL13 in serum, CSF and blood–CSF barrier function: evidence of compartment restriction. Fluids and Barriers of the CNS, 2020, 17, 7.	2.4	14
22	Human Cerebrospinal Fluid Promotes Neuronal Viability and Activity of Hippocampal Neuronal Circuits In Vitro. Frontiers in Cellular Neuroscience, 2016, 10, 54.	1.8	13
23	Molecular evidence of transient therapeutic effectiveness of natalizumab despite high-titre neutralizing antibodies. Multiple Sclerosis Journal, 2012, 18, 506-509.	1.4	12
24	The CXCL13/CXCR5-chemokine axis in neuroinflammation: evidence of CXCR5+CD4 T cell recruitment to CSF. Fluids and Barriers of the CNS, 2021, 18, 40.	2.4	12
25	Basophil Activation Test for Investigation of IgE-Mediated Mechanisms in Drug Hypersensitivity. Journal of Visualized Experiments, 2011, , .	0.2	11
26	Recall response to COVID-19 antigen is preserved in people with multiple sclerosis on anti-CD20 medications – A pilot study. Multiple Sclerosis and Related Disorders, 2022, 59, 103560.	0.9	11
27	Adaptive Immune Responses in a Multiple Sclerosis Patient with Acute Varicella-Zoster Virus Reactivation during Treatment with Fingolimod. International Journal of Molecular Sciences, 2015, 16, 21832-21845.	1.8	10
28	Recent developments in approved and oral multiple sclerosis treatment and an update on future treatment options. Drug Discovery Today, 2011, 16, 8-21.	3.2	9
29	Beyond LNB: Real life data on occurrence and extent of CSF CXCL13 in neuroinflammatory diseases. Journal of Neuroimmunology, 2020, 338, 577087.	1.1	9
30	Neurological complications associated with influenza in season 2017/18 in Austria- a retrospective single center study. Journal of Clinical Virology, 2020, 127, 104340.	1.6	9
31	CD1d expression on chronic lymphocytic leukemia B cells affects disease progression and induces T cell skewing in CD8 positive and CD4CD8 double negative T cells. Oncotarget, 2016, 7, 49459-49469.	0.8	8
32	Isolated leptomeningeal infiltration of a primary CNS B-cell lymphoma diagnosed by flow cytometry and confirmed by necropsy. Acta Neurologica Scandinavica, 2012, 126, e11-e16.	1.0	6
33	From natalizumab to fingolimod in eight weeks — Immunological, clinical, and radiological data in quest of the optimal switch. Clinical Immunology, 2017, 176, 87-93.	1.4	6
34	Immune phenotyping study revealing caveats regarding a switch from fingolimod to cladribine. Multiple Sclerosis and Related Disorders, 2021, 48, 102727.	0.9	5
35	Elevated Toll-Like Receptor-Induced CXCL8 Secretion in Human Blood Basophils from Allergic Donors Is Independent of Toll-Like Receptor Expression Levels. PLoS ONE, 2016, 11, e0149275.	1.1	5
36	Role and Relevance of Cerebrospinal Fluid Cells in Diagnostics and Research: State-of-the-Art and Underutilized Opportunities. Diagnostics, 2022, 12, 79.	1.3	4

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37	High-dose intravenous interferon-beta in multiple sclerosis patients with high-titer neutralizing antibodies (HINABS II) – A pilot study. Multiple Sclerosis and Related Disorders, 2014, 3, 220-226.	0.9	3
38	Epitope competition and neutralizing antidrug antibodies: immune monitoring of antiprogrammed deathâ€I therapies and lessons learned from natalizumab. British Journal of Dermatology, 2020, 183, 404-404.	1.4	3
39	TaqManR Proximity ligation technology for the detection of heterodimeric adhesion receptors on lymphocytes. Journal of Immunological Methods, 2014, 404, 81-86.	0.6	2
40	Serial flow cytometric analyses of blood and cerebrospinal fluid in natalizumabâ€associated progressive multifocal leukencephalopathy with an excellent outcome. Clinical and Experimental Neuroimmunology, 2015, 6, 172-174.	0.5	1
41	Therapeutisches Drug-Monitoring der Natalizumab-SÃŧtigung von Immunzellen mittels Durchflusszytometrie bei Multipler Sklerose/Flow cytometry and drug monitoring of natalizumab saturation of immune cells in multiple sclerosis. Laboratoriums Medizin, 2012, 36, .	0.1	0
42	Zukünftiger Stellenwert von Liquor-Biomarker in der modernen MS-Therapie/Future relevance of CSF biomarkers in modern MS treatment. Laboratoriums Medizin, 2012, 36, .	0.1	0
43	Übersicht über Labormethoden zur Überwachung innovativer Therapieregimes bei Multipler Sklerose/Overview of laboratory methods to monitor innovative treatment regimens in multiple sclerosis. Laboratoriums Medizin, 2012, 36, .	0.1	0
44	Flow cytometry and drug-monitoring of natalizumab saturation of immune cells in multiple	0.1	0
45	A routine-qualified flow cytometric method for the identification of multiple sclerosis patients with a reduced therapeutic effectiveness of natalizumab. Laboratoriums Medizin, 2015, 38, .	0.1	0
46	Routine-taugliche durchflusszytometrische Methode zur Identifikation von Multiple Sklerose PatientInnen mit einer nicht ausreichenden Therapieeffizienz unter einer Natalizumab-Therapie. Laboratoriums Medizin, 2014, 38, .	0.1	0