

Brigitte Bader-Meunier

List of Publications by Year in Descending Order

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Version: 2024-04-27

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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.

62

papers

2,392

citations

21

h-index

48

g-index

70

ext. papers

3,245

ext. citations

6.6

avg, IF

4.28

L-index

#	Paper	IF	Citations
62	Hyper inflammatory syndrome following COVID-19 mRNA vaccine in children: A national post-authorization pharmacovigilance study.. <i>Lancet Regional Health - Europe, The</i> , 2022 , 100393		5
61	Human OTULIN haploinsufficiency impairs cell-intrinsic immunity to staphylococcal Exotoxin.. <i>Science</i> , 2022 , eabm6380	33.3	1
60	Inflammatory myopathies in childhood. <i>Neuromuscular Disorders</i> , 2021 , 31, 1051-1061	2.9	1
59	French recommendations for the management of systemic sclerosis. <i>Orphanet Journal of Rare Diseases</i> , 2021 , 16, 322	4.2	3
58	Improving the diagnostic efficiency of primary immunodeficiencies with targeted next-generation sequencing. <i>Journal of Allergy and Clinical Immunology</i> , 2021 , 147, 734-737	11.5	8
57	Rheumatoid factor positive polyarticular juvenile idiopathic arthritis associated with a novel COPA mutation. <i>Rheumatology</i> , 2021 , 60, e171-e173	3.9	1
56	Overview of STING-Associated Vasculopathy with Onset in Infancy (SAVI) Among 21 Patients. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2021 , 9, 803-818.e11	5.4	19
55	Differential Expression of Interferon-Alpha Protein Provides Clues to Tissue Specificity Across Type I Interferonopathies. <i>Journal of Clinical Immunology</i> , 2021 , 41, 603-609	5.7	7
54	JAK inhibitors are effective in a subset of patients with juvenile dermatomyositis: a monocentric retrospective study. <i>Rheumatology</i> , 2021 , 60, 5801-5808	3.9	6
53	Evaluation of Hydroxychloroquine Blood Concentrations and Effects in Childhood-Onset Systemic Lupus Erythematosus. <i>Pharmaceuticals</i> , 2021 , 14,	5.2	3
52	Onset and Relapse of Juvenile Dermatomyositis Following Asymptomatic SARS-CoV-2 Infection. <i>Journal of Clinical Immunology</i> , 2021 , 1	5.7	0
51	The association of Greig syndrome and mastocytosis reveals the involvement of the hedgehog pathway in advanced mastocytosis. <i>Blood</i> , 2021 , 138, 2396-2407	2.2	2
50	Enhanced cGAS-STING-dependent interferon signaling associated with mutations in ATAD3A. <i>Journal of Experimental Medicine</i> , 2021 , 218,	16.6	8
49	A monocyte/dendritic cell molecular signature of SARS-CoV-2-related multisystem inflammatory syndrome in children with severe myocarditis. <i>Med</i> , 2021 , 2, 1072-1092.e7	31.7	9
48	Mevalonate Kinase Deficiency: A Cause of Severe Very-Early-Onset Inflammatory Bowel Disease. <i>Inflammatory Bowel Diseases</i> , 2021 , 27, 1853-1857	4.5	1
47	Serious adverse events in children with juvenile idiopathic arthritis and other rheumatic diseases on tocilizumab - a real-world experience. <i>Seminars in Arthritis and Rheumatism</i> , 2020 , 50, 744-748	5.3	1
46	Paediatric multisystem inflammatory syndrome temporally associated with SARS-CoV-2 mimicking Kawasaki disease (Kawa-COVID-19): a multicentre cohort. <i>Annals of the Rheumatic Diseases</i> , 2020 , 79, 999-1006	2.4	256

45	Effectiveness and safety of ruxolitinib for the treatment of refractory systemic idiopathic juvenile arthritis like associated with interstitial lung disease : a case report. <i>Annals of the Rheumatic Diseases</i> , 2020 ,	2.4	12
44	Comment on: Monogenic mimics of Behçet disease in the young. <i>Rheumatology</i> , 2020 , 59, e109-e111	3.9	
43	Anti-MDA5 juvenile idiopathic inflammatory myopathy: a specific subgroup defined by differentially enhanced interferon-β signalling. <i>Rheumatology</i> , 2020 , 59, 1927-1937	3.9	12
42	Janus kinase inhibition for autoinflammation in patients with DNASE2 deficiency. <i>Journal of Allergy and Clinical Immunology</i> , 2020 , 145, 701-705.e8	11.5	3
41	Clinical Characteristics of Acne Fulminans Associated With Chronic Nonbacterial Osteomyelitis in Pediatric Patients. <i>Journal of Rheumatology</i> , 2020 , 47, 1793-1799	4.1	4
40	European consensus-based recommendations for diagnosis and treatment of immunoglobulin A vasculitis-the SHARE initiative. <i>Rheumatology</i> , 2019 , 58, 1607-1616	3.9	79
39	The IgG2 Isotype of Anti-Transcription Intermediary Factor 1 Autoantibodies Is a Biomarker of Cancer and Mortality in Adult Dermatomyositis. <i>Arthritis and Rheumatology</i> , 2019 , 71, 1360-1370	9.5	18
38	Comment on: Aberrant tRNA processing causes an autoinflammatory syndrome responsive to TNF inhibitors by Giannelou : mutations in result in a constitutive activation of type I interferon signalling. <i>Annals of the Rheumatic Diseases</i> , 2019 , 78, e86	2.4	7
37	Monogenic lupus: Dissecting heterogeneity. <i>Autoimmunity Reviews</i> , 2019 , 18, 102361	13.6	30
36	Control of TLR7-mediated type I IFN signaling in pDCs through CXCR4 engagement-A new target for lupus treatment. <i>Science Advances</i> , 2019 , 5, eaav9019	14.3	18
35	Genetic of Sporadic Hemophagocytic Lymphohistiocytosis. <i>Blood</i> , 2019 , 134, 82-82	2.2	
34	Therapeutic Plasma Exchange in Pediatrics for Immunologic Disorders; Tolerated and Safe Process for Pediatric Life-Threatening Conditions. <i>Blood</i> , 2019 , 134, 4986-4986	2.2	
33	European consensus-based recommendations for the diagnosis and treatment of rare paediatric vasculitides - the SHARE initiative. <i>Rheumatology</i> , 2019 , 58, 656-671	3.9	50
32	European consensus-based recommendations for the diagnosis and treatment of Kawasaki disease - the SHARE initiative. <i>Rheumatology</i> , 2019 , 58, 672-682	3.9	57
31	Muscle ischaemia associated with NXP2 autoantibodies: a severe subtype of juvenile dermatomyositis. <i>Rheumatology</i> , 2018 , 57, 873-879	3.9	35
30	Inherited Immunodeficiency: A New Association With Early-Onset Childhood Panniculitis. <i>Pediatrics</i> , 2018 , 141, S496-S500	7.4	14
29	Conventional radiography in juvenile idiopathic arthritis: Joint recommendations from the French societies for rheumatology, radiology and paediatric rheumatology. <i>European Radiology</i> , 2018 , 28, 3963-3976	8.9	3
28	Clinical features of children with enthesitis-related juvenile idiopathic arthritis / juvenile spondyloarthritis followed in a French tertiary care pediatric rheumatology centre. <i>Pediatric Rheumatology</i> , 2018 , 16, 21	3.5	28

27	The French version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). <i>Rheumatology International</i> , 2018 , 38, 195-201	3.6	
26	Self-healing juvenile cutaneous mucinosis: Clinical and histopathologic findings of 9 patients: The relevance of long-term follow-up. <i>Journal of the American Academy of Dermatology</i> , 2018 , 78, 1164-1170 ^{4,5}	4.5	14
25	Myogenic Progenitor Cells Exhibit Type I Interferon-Driven Proangiogenic Properties and Molecular Signature During Juvenile Dermatomyositis. <i>Arthritis and Rheumatology</i> , 2018 , 70, 134-145	9.5	25
24	Bone involvement in monogenic autoinflammatory syndromes. <i>Rheumatology</i> , 2018 , 57, 606-618	3.9	11
23	A child with severe juvenile dermatomyositis treated with ruxolitinib. <i>Brain</i> , 2018 , 141, e80	11.2	37
22	Consensus-based recommendations for the management of juvenile dermatomyositis. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 329-340	2.4	119
21	International and multidisciplinary expert recommendations for the use of biologics in systemic lupus erythematosus. <i>Autoimmunity Reviews</i> , 2017 , 16, 650-657	13.6	19
20	Detection of interferon alpha protein reveals differential levels and cellular sources in disease. <i>Journal of Experimental Medicine</i> , 2017 , 214, 1547-1555	16.6	192
19	European evidence-based recommendations for diagnosis and treatment of paediatric antiphospholipid syndrome: the SHARE initiative. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 1637-1641 ^{2,4}	2.4	59
18	European evidence-based recommendations for diagnosis and treatment of childhood-onset systemic lupus erythematosus: the SHARE initiative. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 1788-1796 ^{2,4}	2.4	90
17	Familial and syndromic lupus share the same phenotype as other early-onset forms of lupus. <i>Joint Bone Spine</i> , 2017 , 84, 589-593	2.9	4
16	Assessment of Type I Interferon Signaling in Pediatric Inflammatory Disease. <i>Journal of Clinical Immunology</i> , 2017 , 37, 123-132	5.7	94
15	Fibrous Arthropathy Associated With Morphea: A New Cause of Diffuse Acquired Joint Contractures. <i>Pediatrics</i> , 2017 , 140,	7.4	4
14	European evidence-based recommendations for the diagnosis and treatment of childhood-onset lupus nephritis: the SHARE initiative. <i>Annals of the Rheumatic Diseases</i> , 2017 , 76, 1965-1973	2.4	65
13	Childhood-onset autoimmune cytopenia as the presenting feature of biallelic ACP5 mutations. <i>Pediatric Blood and Cancer</i> , 2017 , 64, 306-310	3	2
12	Type I interferon-mediated autoinflammation due to DNase II deficiency. <i>Nature Communications</i> , 2017 , 8, 2176	17.4	111
11	Vasculopathy-related clinical and pathological features are associated with severe juvenile dermatomyositis. <i>Rheumatology</i> , 2016 , 55, 470-9	3.9	11
10	Efficacy of the Janus kinase 1/2 inhibitor ruxolitinib in the treatment of vasculopathy associated with TMEM173-activating mutations in 3 children. <i>Journal of Allergy and Clinical Immunology</i> , 2016 , 138, 1752-1755	11.5	141

9	Spondyloenchondrodysplasia Due to Mutations in ACP5: A Comprehensive Survey. <i>Journal of Clinical Immunology</i> , 2016 , 36, 220-34	5.7	48
8	Initial presentation and outcome of pediatric-onset mixed connective tissue disease: A French multicenter retrospective study. <i>Joint Bone Spine</i> , 2016 , 83, 369-71	2.9	11
7	Acute pancreatitis as a cause of mortality in pediatric systemic lupus erythematosus: Results of a multiple cause-of-death analysis in France. <i>Seminars in Arthritis and Rheumatism</i> , 2016 , 46, e6-7	5.3	3
6	Gain-of-function mutations in IFIH1 cause a spectrum of human disease phenotypes associated with upregulated type I interferon signaling. <i>Nature Genetics</i> , 2014 , 46, 503-509	36.3	376
5	Association of mastocytosis with inflammatory joint diseases: a series of 31 patients. <i>Seminars in Arthritis and Rheumatism</i> , 2014 , 44, 362-5	5.3	8
4	Clonal cytophagic histiocytic panniculitis in children may be cured by cyclosporine A. <i>Pediatrics</i> , 2013 , 132, e545-9	7.4	17
3	Thrombotic microangiopathy and Purtscher-like retinopathy as a rare presentation of juvenile dermatomyositis. <i>Pediatrics</i> , 2012 , 129, e821-4	7.4	15
2	Safety and efficacy of rituximab in severe juvenile dermatomyositis: results from 9 patients from the French Autoimmunity and Rituximab registry. <i>Journal of Rheumatology</i> , 2011 , 38, 1436-40	4.1	53
1	Mevalonate kinase deficiency: a survey of 50 patients. <i>Pediatrics</i> , 2011 , 128, e152-9	7.4	160