List of Publications by Year in descending order

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		53751	28275
196	12,941	45	105
papers	citations	h-index	g-index
251	251	251	18390
all docs	docs citations	times ranked	citing authors

ALEXANDRE RELAT

#	Article	IF	CITATIONS
1	Autoantibodies against type I IFNs in patients with life-threatening COVID-19. Science, 2020, 370, .	6.0	1,983
2	Inborn errors of type I IFN immunity in patients with life-threatening COVID-19. Science, 2020, 370, .	6.0	1,749
3	EULAR/PRINTO/PRES criteria for Henoch-Schonlein purpura, childhood polyarteritis nodosa, childhood Wegener granulomatosis and childhood Takayasu arteritis: Ankara 2008. Part II: Final classification criteria. Annals of the Rheumatic Diseases, 2010, 69, 798-806.	0.5	1,073
4	Should we stimulate or suppress immune responses in COVID-19? Cytokine and anti-cytokine interventions. Autoimmunity Reviews, 2020, 19, 102567.	2.5	521
5	Interleukin 17 acts in synergy with B cell–activating factor to influence B cell biology and the pathophysiology of systemic lupus erythematosus. Nature Immunology, 2009, 10, 778-785.	7.0	415
6	Autoantibodies neutralizing type I IFNs are present in ~4% of uninfected individuals over 70 years old and account for ~20% of COVID-19 deaths. Science Immunology, 2021, 6, .	5.6	357
7	Detection of interferon alpha protein reveals differential levels and cellular sources in disease. Journal of Experimental Medicine, 2017, 214, 1547-1555.	4.2	288
8	X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. Science Immunology, 2021, 6, .	5.6	267
9	Association of Intravenous Immunoglobulins Plus Methylprednisolone vs Immunoglobulins Alone With Course of Fever in Multisystem Inflammatory Syndrome in Children. JAMA - Journal of the American Medical Association, 2021, 325, 855.	3.8	250
10	SARS-CoV-2-related paediatric inflammatory multisystem syndrome, an epidemiological study, France, 1 March to 17 May 2020. Eurosurveillance, 2020, 25, .	3.9	246
11	Type I IFN immunoprofiling in COVID-19 patients. Journal of Allergy and Clinical Immunology, 2020, 146, 206-208.e2.	1.5	234
12	Human genetic and immunological determinants of critical COVID-19 pneumonia. Nature, 2022, 603, 587-598.	13.7	216
13	COVID-19 outcomes in patients with inflammatory rheumatic and musculoskeletal diseases treated with rituximab: a cohort study. Lancet Rheumatology, The, 2021, 3, e419-e426.	2.2	211
14	Efficacy of the Janus kinase 1/2 inhibitor ruxolitinib in the treatment of vasculopathy associated with TMEM173 -activating mutations in 3 children. Journal of Allergy and Clinical Immunology, 2016, 138, 1752-1755.	1.5	192
15	Assessment of Type I Interferon Signaling in Pediatric Inflammatory Disease. Journal of Clinical Immunology, 2017, 37, 123-132.	2.0	163
16	Severity of COVID-19 and survival in patients with rheumatic and inflammatory diseases: data from the French RMD COVID-19 cohort of 694 patients. Annals of the Rheumatic Diseases, 2021, 80, 527-538.	0.5	156
17	Protein Kinase Cl̂´ Deficiency Causes Mendelian Systemic Lupus Erythematosus With B Cellâ€Defective Apoptosis and Hyperproliferation. Arthritis and Rheumatism, 2013, 65, 2161-2171.	6.7	155
18	Pathogenesis of adult-onset Still's disease: new insights from the juvenile counterpart. Immunologic Research, 2015, 61, 53-62.	1.3	148

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19	Idiopathic inflammatory myopathies and the lung. European Respiratory Review, 2015, 24, 216-238.	3.0	125
20	Vaccination recommendations for the adult immunosuppressed patient: A systematic review and comprehensive field synopsis. Journal of Autoimmunity, 2017, 80, 10-27.	3.0	114
21	Severe Pulmonary Fibrosis as the First Manifestation of Interferonopathy (TMEM173 Mutation). Chest, 2016, 150, e65-e71.	0.4	112
22	ADJUVITE: a double-blind, randomised, placebo-controlled trial of adalimumab in early onset, chronic, juvenile idiopathic arthritis-associated anterior uveitis. Annals of the Rheumatic Diseases, 2018, 77, 1003-1011.	0.5	110
23	The risk of COVID-19 death is much greater and age dependent with type I IFN autoantibodies. Proceedings of the National Academy of Sciences of the United States of America, 2022, 119, e2200413119.	3.3	110
24	Reverse-Transcriptase Inhibitors in the Aicardi–Goutières Syndrome. New England Journal of Medicine, 2018, 379, 2275-2277.	13.9	106
25	Polyclonal expansion of TCR Vβ 21.3 ⁺ CD4 ⁺ and CD8 ⁺ T cells is a hallmark of multisystem inflammatory syndrome in children. Science Immunology, 2021, 6, .	5.6	105
26	SARS-CoV-2–related MIS-C: A key to the viral and genetic causes of Kawasaki disease?. Journal of Experimental Medicine, 2021, 218, .	4.2	100
27	Overview of STING-Associated Vasculopathy with Onset in Infancy (SAVI) Among 21 Patients. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 803-818.e11.	2.0	98
28	Mutations in CECR1 associated with a neutrophil signature in peripheral blood. Pediatric Rheumatology, 2014, 12, 44.	0.9	88
29	Early nasal type I IFN immunity against SARS-CoV-2 is compromised in patients with autoantibodies against type I IFNs. Journal of Experimental Medicine, 2021, 218, .	4.2	85
30	PRKDC mutations associated with immunodeficiency, granuloma, and autoimmune regulator–dependent autoimmunity. Journal of Allergy and Clinical Immunology, 2015, 135, 1578-1588.e5.	1.5	84
31	Antibodies against type I interferon: detection and association with severe clinical outcome in COVIDâ€19 patients. Clinical and Translational Immunology, 2021, 10, e1327.	1.7	79
32	Severe combined immunodeficiency in stimulator of interferon genes (STING) V154M/wild-type mice. Journal of Allergy and Clinical Immunology, 2019, 143, 712-725.e5.	1.5	74
33	Monogenic lupus: Dissecting heterogeneity. Autoimmunity Reviews, 2019, 18, 102361.	2.5	74
34	Early-onset autoimmunity associated with SOCS1 haploinsufficiency. Nature Communications, 2020, 11, 5341.	5.8	74
35	Factors Associated With Severe SARS-CoV-2 Infection. Pediatrics, 2021, 147, .	1.0	73
36	Studying severe long COVID to understand post-infectious disorders beyond COVID-19. Nature Medicine, 2022, 28, 879-882.	15.2	72

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37	The Relapsing Polychondritis Disease Activity Index: Development of a disease activity score for relapsing polychondritis. Autoimmunity Reviews, 2012, 12, 204-209.	2.5	71
38	DNA-PK deficiency potentiates cGAS-mediated antiviral innate immunity. Nature Communications, 2020, 11, 6182.	5.8	70
39	Pseudohypoaldosteronisms, report on a 10-patient series. Nephrology Dialysis Transplantation, 2008, 23, 1636-1641.	0.4	69
40	Familial Mediterranean fever mutations are hypermorphic mutations that specifically decrease the activation threshold of the Pyrin inflammasome. Rheumatology, 2018, 57, 100-111.	0.9	67
41	A decision tree for the genetic diagnosis of deficiency of adenosine deaminase 2 (DADA2): a French reference centres experience. European Journal of Human Genetics, 2018, 26, 960-971.	1.4	65
42	Genetic and phenotypic spectrum associated with IFIH1 gainâ€ofâ€function. Human Mutation, 2020, 41, 837-849.	1.1	63
43	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. Journal of Experimental Medicine, 2022, 219, .	4.2	59
44	Monogenic forms of systemic lupus erythematosus: new insights into SLE pathogenesis. Pediatric Rheumatology, 2012, 10, 21.	0.9	55
45	Refining "Long-COVID―by a Prospective Multimodal Evaluation of Patients with Long-Term Symptoms Attributed to SARS-CoV-2 Infection. Infectious Diseases and Therapy, 2021, 10, 1747-1763.	1.8	55
46	Pyrin dephosphorylation is sufficient to trigger inflammasome activation in familial Mediterranean fever patients. EMBO Molecular Medicine, 2019, 11, e10547.	3.3	54
47	Pediatric-Onset Relapsing Polychondritis: Case Series and Systematic Review. Journal of Pediatrics, 2010, 156, 484-489.	0.9	52
48	Comparison of RT-qPCR and Nanostring in the measurement of blood interferon response for the diagnosis of type I interferonopathies. Cytokine, 2019, 113, 446-452.	1.4	51
49	Phase II Open Label Study of Anakinra in Intravenous Immunoglobulin–Resistant Kawasaki Disease. Arthritis and Rheumatology, 2021, 73, 151-161.	2.9	51
50	Recommendations for using TNFα antagonists and French Clinical Practice Guidelines endorsed by the French National Authority for Health. Joint Bone Spine, 2013, 80, 574-581.	0.8	48
51	Tartrateâ€Resistant Acid Phosphatase Deficiency in the Predisposition to Systemic Lupus Erythematosus. Arthritis and Rheumatology, 2017, 69, 131-142.	2.9	47
52	Clinical characteristics and outcomes of childhood-onset ANCA-associated vasculitis: a French nationwide study. Nephrology Dialysis Transplantation, 2015, 30 Suppl 1, i104-12.	0.4	45
53	Subcutaneous Abatacept in Patients With Polyarticularâ€Course Juvenile Idiopathic Arthritis. Arthritis and Rheumatology, 2018, 70, 1144-1154.	2.9	45
54	Mosaicism in autoinflammatory diseases: Cryopyrin-associated periodic syndromes (CAPS) and beyond. A systematic review. Critical Reviews in Clinical Laboratory Sciences, 2018, 55, 432-442.	2.7	45

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55	Inherited IFNAR1 Deficiency in a Child with Both Critical COVID-19 Pneumonia and Multisystem Inflammatory Syndrome. Journal of Clinical Immunology, 2022, 42, 471-483.	2.0	44
56	Hyper inflammatory syndrome following COVID-19 mRNA vaccine in children: A national post-authorization pharmacovigilance study. Lancet Regional Health - Europe, The, 2022, 17, 100393.	3.0	44
57	PROMIDISα: AÂT-cell receptor α signature associated with immunodeficiencies caused by V(D)J recombination defects. Journal of Allergy and Clinical Immunology, 2019, 143, 325-334.e2.	1.5	43
58	Does type-l interferon drive systemic autoimmunity?. Autoimmunity Reviews, 2017, 16, 897-902.	2.5	40
59	The pyrin inflammasome: from sensing RhoA GTPases-inhibiting toxins to triggering autoinflammatory syndromes. Pathogens and Disease, 2018, 76, .	0.8	40
60	S1PR5 is essential for human natural killer cell migration toward sphingosine-1 phosphate. Journal of Allergy and Clinical Immunology, 2018, 141, 2265-2268.e1.	1.5	39
61	Contribution of rare and predicted pathogenic gene variants to childhood-onset lupus: a large, genetic panel analysis of British and French cohorts. Lancet Rheumatology, The, 2020, 2, e99-e109.	2.2	38
62	Varicella as a trigger of atypical haemolytic uraemic syndrome associated with complement dysfunction: two cases. Nephrology Dialysis Transplantation, 2009, 24, 2752-2754.	0.4	37
63	French recommendations for the management of systemic sclerosis. Orphanet Journal of Rare Diseases, 2021, 16, 322.	1.2	37
64	Multisystem Inflammatory Syndrome in Children in the United States. New England Journal of Medicine, 2020, 383, 1793-1796.	13.9	34
65	Pediatric Inflammatory Multisystem Syndrome and Rheumatic Diseases During SARS-CoV-2 Pandemic. Frontiers in Pediatrics, 2020, 8, 605807.	0.9	34
66	Rare diseases that mimic Systemic Lupus Erythematosus (Lupus mimickers). Joint Bone Spine, 2019, 86, 165-171.	0.8	31
67	Severe infections in patients with anti-neutrophil cytoplasmic antibody-associated vasculitides receiving rituximab: A meta-analysis. Autoimmunity Reviews, 2020, 19, 102505.	2.5	30
68	Pro-inflammatory genotype as a risk factor for aPL-associated thrombosis: Report of a family with multiple anti-phospholipid positive members. Journal of Autoimmunity, 2009, 32, 60-63.	3.0	28
69	Orbital mass in ANCA-associated vasculitides: data on clinical, biological, radiological and histological presentation, therapeutic management, and outcome from 59 patients. Rheumatology, 2019, 58, 1565-1573.	0.9	28
70	Geoepidemiology and Immunologic Features of Autoinflammatory Diseases: a Comprehensive Review. Clinical Reviews in Allergy and Immunology, 2018, 54, 454-479.	2.9	27
71	Anti-MDA5 juvenile idiopathic inflammatory myopathy: a specific subgroup defined by differentially enhanced interferon- $\hat{1}_{\pm}$ signalling. Rheumatology, 2020, 59, 1927-1937.	0.9	26
72	Efficacy and safety of TNF-α antagonists and tocilizumab in Takayasu arteritis: multicentre retrospective study of 209 patients. Rheumatology, 2022, 61, 1376-1384.	0.9	26

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73	Transplant Tolerance Induction in Newborn Infants: Mechanisms, Advantages, and Potential Strategies. Frontiers in Immunology, 2016, 7, 116.	2.2	24
74	Building a transitional care checklist in rheumatology: A Delphi-like survey. Joint Bone Spine, 2018, 85, 435-440.	0.8	24
75	Human Naive and Memory T Cells Display Opposite Migratory Responses to Sphingosine-1 Phosphate. Journal of Immunology, 2018, 200, 551-557.	0.4	23
76	Tapering Canakinumab Monotherapy in Patients With Systemic Juvenile Idiopathic Arthritis in Clinical Remission: Results From a Phase IIIb/IV Open‣abel, Randomized Study. Arthritis and Rheumatology, 2021, 73, 336-346.	2.9	23
77	Impaired respiratory burst contributes to infections in PKCδ-deficient patients. Journal of Experimental Medicine, 2021, 218, .	4.2	23
78	Chemoresistance of Human Monocyte-Derived Dendritic Cells Is Regulated by IL-17A. PLoS ONE, 2013, 8, e56865.	1.1	22
79	ANCA-associated vasculitides: Recommendations of the French Vasculitis Study Group on the use of immunosuppressants and biotherapies for remission induction and maintenance. Presse Medicale, 2020, 49, 104031.	0.8	21
80	DEF6 deficiency, a mendelian susceptibility to EBV infection, lymphoma, and autoimmunity. Journal of Allergy and Clinical Immunology, 2021, 147, 740-743.e9.	1.5	21
81	From Your Nose to Your Toes: A Review of Severe Acute Respiratory Syndrome Coronavirus 2 Pandemic‒Associated Pernio. Journal of Investigative Dermatology, 2021, 141, 2791-2796.	0.3	21
82	Trichuris suis induces human non-classical patrolling monocytes via the mannose receptor and PKC: implications for multiple sclerosis. Acta Neuropathologica Communications, 2015, 3, 45.	2.4	20
83	Protracted viral shedding and viral load are associated with ICU mortality in Covid-19 patients with acute respiratory failure. Annals of Intensive Care, 2020, 10, 167.	2.2	20
84	Further delineation of the clinical spectrum of de novo <i>TRIM8</i> truncating mutations. American Journal of Medical Genetics, Part A, 2018, 176, 2470-2478.	0.7	19
85	COPA Syndrome as a Cause of Lupus Nephritis. Kidney International Reports, 2019, 4, 1187-1189.	0.4	19
86	Massive increase in monocyte HLA-DR expression can be used to discriminate between septic shock and hemophagocytic lymphohistiocytosis-induced shock. Critical Care, 2018, 22, 213.	2.5	18
87	Initial presentation and outcome of pediatric-onset mixed connective tissue disease: A French multicenter retrospective study. Joint Bone Spine, 2016, 83, 369-371.	0.8	17
88	LACC1 deficiency links juvenile arthritis with autophagy and metabolism in macrophages. Journal of Experimental Medicine, 2021, 218, .	4.2	17
89	Fast diagnostic test for familial Mediterranean fever based on a kinase inhibitor. Annals of the Rheumatic Diseases, 2021, 80, 128-132.	0.5	16
90	INSAID Variant Classification and Eurofever Criteria Guide Optimal Treatment Strategy in Patients with TRAPS: Data from the Eurofever Registry. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 783-791.e4.	2.0	16

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91	Successful Immunotherapy in Life-threatening Parvovirus B19 Infection in a Child. Pediatric Infectious Disease Journal, 2013, 32, 789-792.	1.1	15
92	Safety of biological agents in paediatric rheumatic diseases: A real-life multicenter retrospective study using the JIRcohorte database. Joint Bone Spine, 2019, 86, 343-350.	0.8	15
93	Circulating Interferonâ€Î± Measured With a Highly Sensitive Assay as a Biomarker for Juvenile Inflammatory Myositis Activity: Comment on the Article by Mathian et al. Arthritis and Rheumatology, 2020, 72, 195-197.	2.9	15
94	Severe Acute Respiratory Syndrome Coronavirus 2 Vaccination in Children with a History of Multisystem Inflammatory Syndrome in Children: AnÂInternational Survey. Journal of Pediatrics, 2022, 248, 114-118.	0.9	15
95	Large deletion in 6q associated to A20 haploinsufficiency and thoracoabdominal heterotaxy. Annals of the Rheumatic Diseases, 2018, 77, 1697-1698.	0.5	14
96	DADA2 diagnosed in adulthood versus childhood: A comparative study on 306 patients including a systematic literature review and 12 French cases. Seminars in Arthritis and Rheumatism, 2021, 51, 1170-1179.	1.6	14
97	Inherited anomalies of innate immune receptors in pediatric-onset inflammatory diseases. Autoimmunity Reviews, 2015, 14, 1147-1153.	2.5	13
98	Tocilizumab in the treatment of mixed connective tissue disease and overlap syndrome in children. RMD Open, 2016, 2, e000271.	1.8	13
99	Reversible cerebral vasoconstriction syndrome in paediatric patients with systemic lupus erythematosus: implications for management. Developmental Medicine and Child Neurology, 2019, 61, 725-729.	1.1	13
100	Type I Interferon in Children with Viral or Bacterial Infections. Clinical Chemistry, 2020, 66, 802-808.	1.5	13
101	Therapy for Multisystem Inflammatory Syndrome in Children. New England Journal of Medicine, 2021, 385, e42.	13.9	13
102	Infectious adverse events in children with Juvenile Idiopathic Arthritis treated with Biological Agents in a real-life setting: Data from the JIRcohorte. Joint Bone Spine, 2020, 87, 49-55.	0.8	12
103	Outcomes of SARS-CoV-2 infection among children and young people with pre-existing rheumatic and musculoskeletal diseases. Annals of the Rheumatic Diseases, 2022, 81, 998-1005.	0.5	12
104	Mevalonate Kinase Deficiency: A Cause of Severe Very-Early-Onset Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2021, 27, 1853-1857.	0.9	11
105	Detection and Prediction of Macrophage Activation Syndrome in Still's Disease. Journal of Clinical Medicine, 2022, 11, 206.	1.0	11
106	ANCA-Associated Glomerulonephritis in Systemic-Onset Juvenile Idiopathic Arthritis. American Journal of Kidney Diseases, 2012, 59, 439-443.	2.1	10
107	PKCδ is dispensible for oxLDL uptake and foam cell formation by human and murine macrophages. Cardiovascular Research, 2014, 104, 467-476.	1.8	10
108	MISS questionnaire in French version: a good tool for children and parents to assess methotrexate intolerance. Clinical Rheumatology, 2017, 36, 1281-1288.	1.0	10

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109	Positive Impact of Expert Reference Center Validation on Performance of Next-Generation Sequencing for Genetic Diagnosis of Autoinflammatory Diseases. Journal of Clinical Medicine, 2019, 8, 1729.	1.0	9
110	French expert opinion for the management of juvenile dermatomyositis. Archives De Pediatrie, 2019, 26, 120-125.	0.4	9
111	Validation of the new classification criteria for hereditary recurrent fever in an independent cohort: experience from the JIR Cohort Database. Rheumatology, 2020, 59, 2947-2952.	0.9	9
112	Impaired microbial killing by neutrophils from patients with protein kinase C delta deficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 1404-1407.e10.	1.5	8
113	Clinical Profile of Methotrexate-resistant Juvenile Localised Scleroderma. Acta Dermato-Venereologica, 2019, 99, 539-543.	0.6	8
114	The relapsing polychondritis damage index (RPDAM): Development of a disease-specific damage score for relapsing polychondritis. Joint Bone Spine, 2019, 86, 363-368.	0.8	8
115	Practical management of patients on hydroxychloroquine. Joint Bone Spine, 2021, 88, 105316.	0.8	8
116	Familial and syndromic lupus share the same phenotype as other early-onset forms of lupus. Joint Bone Spine, 2017, 84, 589-593.	0.8	7
117	New classification for juvenile idiopathic arthritis: Is the Tower of Babel falling?. Joint Bone Spine, 2018, 85, 139-141.	0.8	7
118	The first case report of medulloblastoma associated with Tattonâ€Brown–Rahman syndrome. American Journal of Medical Genetics, Part A, 2019, 179, 1357-1361.	0.7	7
119	Etanercept concentration and immunogenicity do not influence the response to Etanercept in patients with juvenile idiopathic arthritis. Seminars in Arthritis and Rheumatism, 2019, 48, 1014-1018.	1.6	7
120	NLRC4 GOF Mutations, a Challenging Diagnosis from Neonatal Age to Adulthood. Journal of Clinical Medicine, 2021, 10, 4369.	1.0	7
121	Long-Term Follow-Up and Optimization of Interleukin-1 Inhibitors in the Management of Monogenic Autoinflammatory Diseases: Real-Life Data from the JIR Cohort. Frontiers in Pharmacology, 2020, 11, 568865.	1.6	7
122	DNASE1L3 deficiency, new phenotypes, and evidence for a transient type I IFN signaling. Journal of Clinical Immunology, 2022, 42, 1310-1320.	2.0	7
123	Conseils d'utilisation des traitements anti-TNF et recommandations nationales de bonne pratique labellisées par la Haute Autorité de santé française. Revue Du Rhumatisme (Edition Francaise), 2013, 80, 459-466.	0.0	6
124	Health related quality of life measure in systemic pediatric rheumatic diseases and its translation to different languages: an international collaboration. Pediatric Rheumatology, 2014, 12, 49.	0.9	6
125	A Case of Type 2 Hypersensitivity to Rasburicase Diagnosed with a Natural Killer Cell Activation Assay. Frontiers in Immunology, 2018, 9, 110.	2.2	6
126	Patients' association programs for adolescents and young adults: The JAP study. Archives De Pediatrie, 2019, 26, 205-213.	0.4	6

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127	Comparison of paediatric and adult classification criteria in juvenile idiopathic arthritis during the transition from paediatric to adult care. Joint Bone Spine, 2021, 88, 105047.	0.8	6
128	The benefit–risk balance for biological agents in juvenile idiopathic arthritis: a meta-analysis of randomized clinical trials. Rheumatology, 2020, 59, 2226-2236.	0.9	6
129	Causes of death in pediatric systemic lupus erythematosus. Clinical and Experimental Rheumatology, 2009, 27, 538-9.	0.4	6
130	Anti-C1q autoantibodies as markers of renal involvement in childhood-onset systemic lupus erythematosus. Pediatric Nephrology, 2017, 32, 1537-1545.	0.9	5
131	Deletion of Inflammasome Components Is Not Sufficient To Prevent Fatal Inflammation in Models of Familial Hemophagocytic Lymphohistiocytosis. Journal of Immunology, 2018, 200, 3769-3776.	0.4	5
132	Le lupus de l'enfant à travers les âges. Revue Du Rhumatisme Monographies, 2012, 79, 24-29.	0.0	4
133	Kimura Disease Mimicking an Aneurysm of the Radial Artery. Journal of Pediatrics, 2015, 167, 1166-1166.e2.	0.9	4
134	Earlyâ€onset hypoparathyroidism and chronic keratitis revealing <scp>APECED</scp> . Clinical Case Reports (discontinued), 2015, 3, 809-813.	0.2	4
135	Typeâ€i Interferon assessment in 45 minutes using the FilmArray [®] PCR platform in SARSâ€CoVâ€2 and other viral infections. European Journal of Immunology, 2021, 51, 989-994.	1.6	4
136	Impact of hydroxychloroquine used as DMARD on SARS-CoV-2 tests and infection evolution in a population of 871 patients with inflammatory rheumatic and musculoskeletal diseases. Joint Bone Spine, 2021, 88, 105226.	0.8	4
137	An Immunological Axis Involving Interleukin 1β and Leucine-Rich-α2-Glycoprotein Reflects Therapeutic Response of Children with Kawasaki Disease: Implications from the KAWAKINRA Trial. Journal of Clinical Immunology, 2022, 42, 1330-1341.	2.0	4
138	Hypertensive crisis, hepatitis B virus and polyarteritis nodosa in a child. Pediatric Nephrology, 2007, 22, 97-100.	0.9	3
139	It sounds like a relapsing polychondritis. Lancet Infectious Diseases, The, 2013, 13, 638.	4.6	3
140	French Amyloidosis CAPS study: AA Amyloidosis complicating cryopyrin-associated periodic syndrome: a study on 14 cases and review of 53 cases from literature. Pediatric Rheumatology, 2015, 13, .	0.9	3
141	Acute pancreatitis as a cause of mortality in pediatric systemic lupus erythematosus: Results of a multiple cause-of-death analysis in France. Seminars in Arthritis and Rheumatism, 2016, 46, e6-e7.	1.6	3
142	Lateâ€onset hemophagocytic lymphohistiocytosis with neurological presentation. Clinical Case Reports (discontinued), 2017, 5, 1743-1749.	0.2	3
143	Educational Setting and SARS-CoV-2 Transmission Among Children With Multisystem Inflammatory Syndrome: A French National Surveillance System. Frontiers in Pediatrics, 2021, 9, 745364.	0.9	3
144	Chronic non-bacterial osteomyelitis: a retrospective international study on clinical manifestations and response to treatment. Clinical and Experimental Rheumatology, 2020, 38, 1255-1262.	0.4	3

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145	THU0235â€Adjuvite: A Double-Blind, Randomized, Placebo-Controlled Trial of Adalimumab in Juvenile Idiopathic Arthritis Associated Uveitis: Table 1 Annals of the Rheumatic Diseases, 2016, 75, 273.1-273.	0.5	2
146	lsolated positive anti-SS-B autoantibodies are not related to clinical features of systemic autoimmune diseases: Results from a routine population survey. PLoS ONE, 2017, 12, e0185104.	1.1	2
147	POS1183â€OUTCOMES OF COVID-19 INFECTION AMONG CHILDREN AND YOUNG PEOPLE WITH PRE-EXISTING RHEUMATIC AND MUSCULOSKELETAL DISEASES. Annals of the Rheumatic Diseases, 2021, 80, 872.2-873.	0.5	2
148	Real-Life Indications of Interleukin-1 Blocking Agents in Hereditary Recurrent Fevers: Data From the JIRcohort and a Literature Review. Frontiers in Immunology, 2021, 12, 744780.	2.2	2
149	Human Papilloma Virus Vaccination in Patients with Rheumatic Diseases in France: A Study of Vaccination Coverage and Drivers of Vaccination. Journal of Clinical Medicine, 2022, 11, 4137.	1.0	2
150	Tocilizumab for the treatment of refractory pediatric mixed connective tissue disease (MCTD), in two patients. Pediatric Rheumatology, 2014, 12, .	0.9	1
151	Prescribed but not approved: biologic agents used without approval in juvenile idiopathic arthritis in Switzerland, France and Belgium. Pediatric Rheumatology, 2014, 12, .	0.9	1
152	PRKDC mutations associated with immunodeficiency, granuloma and aire-dependent autoimmunity. Pediatric Rheumatology, 2014, 12, .	0.9	1
153	SAT0263â€Cohort Study of 80 Patients with Juvenile Idiopathic Arthritis during Transition from Pediatric To Adult Care. Annals of the Rheumatic Diseases, 2016, 75, 763.3-764.	0.5	1
154	Particularités pédiatriques des vascularites. Revue Du Rhumatisme Monographies, 2017, 84, 290-296.	0.0	1
155	Polymorphic gastric lesions and hemorrhage after first dose of chemotherapy in a child with diffuse large B-cell lymphoma. Clinics and Research in Hepatology and Gastroenterology, 2018, 42, 175-177.	0.7	1
156	A quest for Q fever. Lancet, The, 2019, 394, 419.	6.3	1
157	AB0947â€TREATMENT WITH SIMULTANEOUS BIOLOGICAL AGENTS IN JUVENILE IDIOPATHIC ARTHRITIS: SINGLE-CENTER CASE SERIES AND REVIEW OF LITERATURE. , 2019, , .		1
158	Immunomodulatory treatment and surgical management of idiopathic uveitis and juvenile idiopathic arthritis-associated uveitis in children: a French survey practice. Pediatric Rheumatology, 2021, 19, 139.	0.9	1
159	Neutralizing Anti-IL-17A Antibody Demonstrates Preclinical Activity Enhanced by Vinblastine in Langerhans Cell Histiocytosis. Frontiers in Oncology, 2021, 11, 780191.	1.3	1
160	"P2RY8-son―break of tolerance promotes SLE. Journal of Experimental Medicine, 2022, 219, .	4.2	1
161	Relapsing polychondritis: a pediatric series of ten patients. Pediatric Rheumatology, 2008, 6, .	0.9	Ο
162	Systemic-onset juvenile rheumatoid arthritis and ANCA-associated glomerulonephritis. Pediatric Rheumatology, 2008, 6, .	0.9	0

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163	IL-17 in B Cell Biology and Systemic Lupus Erythematosus. , 2011, , 401-410.		0
164	Pathogénie de la maladie de KawasakiÂ: quoi de neufÂ?. Revue Du Rhumatisme Monographies, 2012, 79, 20-23.	0.0	0
165	PReS-FINAL-2311: Rituximab in paediatric ANCA-associated vasculitis. Pediatric Rheumatology, 2013, 11, .	0.9	0
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