Paul Brogan

List of Publications by Year in descending order

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304743 345221 2,824 36 22 36 h-index citations g-index papers 36 36 36 3298 times ranked docs citations citing authors all docs

#	Article	IF	CITATIONS
1	Phase <scp>IIa</scp> Global Study Evaluating Rituximab for the Treatment of Pediatric Patients With Granulomatosis With Polyangiitis or Microscopic Polyangiitis. Arthritis and Rheumatology, 2022, 74, 124-133.	5.6	12
2	The impact of the Eurofever criteria and the new InFevers MEFV classification in real life: Results from a large international FMF cohort. Seminars in Arthritis and Rheumatism, 2022, 52, 151957.	3.4	7
3	Evaluation of Serious Infection in Pediatric Patients with Low Immunoglobulin Levels Receiving Rituximab for Granulomatosis with Polyangiitis or Microscopic Polyangiitis. Rheumatology and Therapy, 2022, 9, 721-734.	2.3	2
4	Management of severe hyperinflammation in the COVID-19 era: the role of the rheumatologist. Rheumatology, 2021, 60, 911-917.	1.9	7
5	Ofatumumab use in juvenile systemic lupus erythematosus: A single centre experience. Lupus, 2021, 30, 527-530.	1.6	10
6	Development and Validation of a Targeted Next-Generation Sequencing Gene Panel for Children With Neuroinflammation. JAMA Network Open, 2019, 2, e1914274.	5.9	14
7	Efficacy and safety of anakinra for undifferentiated autoinflammatory diseases in children: a retrospective case review. Rheumatology Advances in Practice, 2019, 3, rkz004.	0.7	14
8	European consensus-based recommendations for diagnosis and treatment of immunoglobulin A vasculitisâ€"the SHARE initiative. Rheumatology, 2019, 58, 1607-1616.	1.9	165
9	Secondary C1q Deficiency in Activated PI3Kδ Syndrome Type 2. Frontiers in Immunology, 2019, 10, 2589.	4.8	7
10	Mycophenolate mofetil versus cyclophosphamide for remission induction in ANCA-associated vasculitis: a randomised, non-inferiority trial. Annals of the Rheumatic Diseases, 2019, 78, 399-405.	0.9	165
11	European consensus-based recommendations for the diagnosis and treatment of rare paediatric vasculitides $\hat{a} \in \text{HARE}$ initiative. Rheumatology, 2019, 58, 656-671.	1.9	77
12	European consensus-based recommendations for the diagnosis and treatment of Kawasaki disease – the SHARE initiative. Rheumatology, 2019, 58, 672-682.	1.9	103
13	Moyamoya-like cerebrovascular disease in a child with a novel mutation in myosin heavy chain 11. Neurology, 2018, 90, 136-138.	1.1	18
14	Molecular genetic investigation, clinical features, and response to treatment in 21 patients with Schnitzler syndrome. Blood, 2018, 131, 974-981.	1.4	83
15	Vasculitis update: pathogenesis and biomarkers. Pediatric Nephrology, 2018, 33, 187-198.	1.7	45
16	Successful use of ofatumumab in two cases of early-onset juvenile SLE with thrombocytopenia caused by a mutation in protein kinase C Î'. Pediatric Rheumatology, 2018, 16, 61.	2.1	23
17	Allogeneic hematopoietic stem cell transplantation for severe, refractory juvenile idiopathic arthritis. Blood Advances, 2018, 2, 777-786.	5.2	37
18	Canakinumab for the Treatment of Autoinflammatory Recurrent Fever Syndromes. New England Journal of Medicine, 2018, 378, 1908-1919.	27.0	327

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19	European evidence-based recommendations for diagnosis and treatment of paediatric antiphospholipid syndrome: the SHARE initiative. Annals of the Rheumatic Diseases, 2017, 76, 1637-1641.	0.9	75
20	European evidence-based recommendations for diagnosis and treatment of childhood-onset systemic lupus erythematosus: the SHARE initiative. Annals of the Rheumatic Diseases, 2017, 76, 1788-1796.	0.9	139
21	European evidence-based recommendations for the diagnosis and treatment of childhood-onset lupus nephritis: the SHARE initiative. Annals of the Rheumatic Diseases, 2017, 76, 1965-1973.	0.9	105
22	Late-Onset Cryopyrin-Associated Periodic Syndromes Caused by Somatic NLRP3 Mosaicism—UK Single Center Experience. Frontiers in Immunology, 2017, 8, 1410.	4.8	109
23	De Novo <i>PTEN</i> Mutation in a Young Boy with Cutaneous Vasculitis. Case Reports in Pediatrics, 2017, 2017, 1-4.	0.4	12
24	Brief Report: Association of Tumor Necrosis Factor Receptor–Associated Periodic Syndrome With Gonosomal Mosaicism of a Novel 24â€Nucleotide <i>TNFRSF1A</i> Deletion. Arthritis and Rheumatology, 2016, 68, 2044-2049.	5.6	49
25	Progressive neurologic disorder: Initial manifestation of hemophagocytic lymphohistiocytosis. Neurology, 2016, 86, 2109-2111.	1.1	14
26	Allogeneic Haematopoietic Stem Cell Transplantation for Systemic Onset Juvenile Idiopathic Arthritis. Biology of Blood and Marrow Transplantation, 2015, 21, S46.	2.0	2
27	Evidence-based recommendations for genetic diagnosis of familial Mediterranean fever. Annals of the Rheumatic Diseases, 2015, 74, 635-641.	0.9	145
28	Additive loss-of-function proteasome subunit mutations in CANDLE/PRAAS patients promote type I IFN production. Journal of Clinical Investigation, 2015, 125, 4196-4211.	8.2	258
29	Bayesian methods for the design and interpretation of clinical trials in very rare diseases. Statistics in Medicine, 2014, 33, 4186-4201.	1.6	74
30	Clinical features of childhood granulomatosis with polyangiitis (wegener's granulomatosis). Pediatric Rheumatology, 2014, 12, 18.	2.1	85
31	Long-term outcome of paediatric patients with ANCA vasculitis. Pediatric Rheumatology, 2011, 9, 12.	2.1	42
32	Small vessel vasculitis. Pediatric Nephrology, 2010, 25, 1025-1035.	1.7	53
33	EULAR points to consider in the development of classification and diagnostic criteria in systemic vasculitis. Annals of the Rheumatic Diseases, 2010, 69, 1744-1750.	0.9	139
34	EULAR/PRINTO/PRES criteria for Henoch-Schonlein purpura, childhood polyarteritis nodosa, childhood Wegener granulomatosis and childhood Takayasu arteritis: Ankara 2008. Part I: Overall methodology and clinical characterisation. Annals of the Rheumatic Diseases, 2010, 69, 790-797.	0.9	187
35	Cardiovascular involvement in primary systemic vasculitis. Best Practice and Research in Clinical Rheumatology, 2009, 23, 419-428.	3.3	24
36	Juvenile polyarteritis: Results of a multicenter survey of 110 children. Journal of Pediatrics, 2004, 145, 517-522.	1.8	196