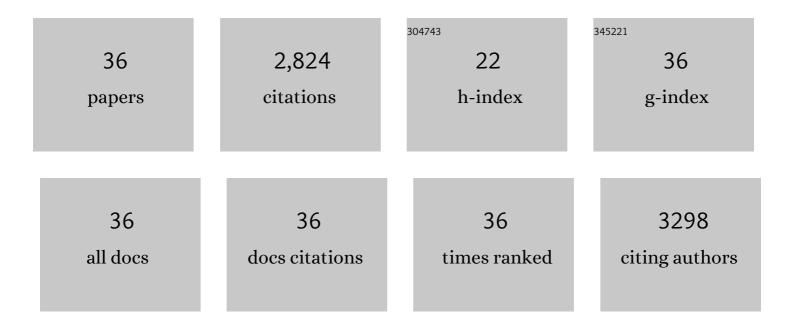
Paul Brogan

List of Publications by Year in descending order

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DALLI RROCAN

#	Article	IF	CITATIONS
1	Canakinumab for the Treatment of Autoinflammatory Recurrent Fever Syndromes. New England Journal of Medicine, 2018, 378, 1908-1919.	27.0	327
2	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
3	Juvenile polyarteritis: Results of a multicenter survey of 110 children. Journal of Pediatrics, 2004, 145, 517-522.	1.8	196
4	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
5	European consensus-based recommendations for diagnosis and treatment of immunoglobulin A vasculitis—the SHARE initiative. Rheumatology, 2019, 58, 1607-1616.	1.9	165
6	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
7	Evidence-based recommendations for genetic diagnosis of familial Mediterranean fever. Annals of the Rheumatic Diseases, 2015, 74, 635-641.	0.9	145
8	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
9	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
10	Late-Onset Cryopyrin-Associated Periodic Syndromes Caused by Somatic NLRP3 Mosaicism—UK Single Center Experience. Frontiers in Immunology, 2017, 8, 1410.	4.8	109
11	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
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	Clinical features of childhood granulomatosis with polyangiitis (wegener's granulomatosis). Pediatric Rheumatology, 2014, 12, 18.	2.1	85
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	European consensus-based recommendations for the diagnosis and treatment of rare paediatric vasculitides – the SHARE initiative. Rheumatology, 2019, 58, 656-671.	1.9	77
16	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
17	Bayesian methods for the design and interpretation of clinical trials in very rare diseases. Statistics in Medicine, 2014, 33, 4186-4201.	1.6	74
18	Small vessel vasculitis. Pediatric Nephrology, 2010, 25, 1025-1035.	1.7	53

PAUL BROGAN

#	Article	IF	CITATIONS
19	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
20	Vasculitis update: pathogenesis and biomarkers. Pediatric Nephrology, 2018, 33, 187-198.	1.7	45
21	Long- term outcome of paediatric patients with ANCA vasculitis. Pediatric Rheumatology, 2011, 9, 12.	2.1	42
22	Allogeneic hematopoietic stem cell transplantation for severe, refractory juvenile idiopathic arthritis. Blood Advances, 2018, 2, 777-786.	5.2	37
23	Cardiovascular involvement in primary systemic vasculitis. Best Practice and Research in Clinical Rheumatology, 2009, 23, 419-428.	3.3	24
24	Successful use of ofatumumab in two cases of early-onset juvenile SLE with thrombocytopenia caused by a mutation in protein kinase C Ĩ. Pediatric Rheumatology, 2018, 16, 61.	2.1	23
25	Moyamoya-like cerebrovascular disease in a child with a novel mutation in myosin heavy chain 11. Neurology, 2018, 90, 136-138.	1.1	18
26	Progressive neurologic disorder: Initial manifestation of hemophagocytic lymphohistiocytosis. Neurology, 2016, 86, 2109-2111.	1.1	14
27	Development and Validation of a Targeted Next-Generation Sequencing Gene Panel for Children With Neuroinflammation. JAMA Network Open, 2019, 2, e1914274.	5.9	14
28	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
29	De Novo <i> PTEN</i> Mutation in a Young Boy with Cutaneous Vasculitis. Case Reports in Pediatrics, 2017, 2017, 1-4.	0.4	12
30	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
31	Ofatumumab use in juvenile systemic lupus erythematosus: A single centre experience. Lupus, 2021, 30, 527-530.	1.6	10
32	Secondary C1q Deficiency in Activated PI3Kδ Syndrome Type 2. Frontiers in Immunology, 2019, 10, 2589.	4.8	7
33	Management of severe hyperinflammation in the COVID-19 era: the role of the rheumatologist. Rheumatology, 2021, 60, 911-917.	1.9	7
34	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
35	Allogeneic Haematopoietic Stem Cell Transplantation for Systemic Onset Juvenile Idiopathic Arthritis. Biology of Blood and Marrow Transplantation, 2015, 21, S46.	2.0	2
36	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