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List of Publications by Year in descending order

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55 5,015 31 51
papers citations h-index g-index

90 90 90 3689 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Metabolomics analysis identifies a lipidomic profile in treatment-na \tilde{A} -ve juvenile dermatomyositis patients <i>vs</i> healthy control subjects. Rheumatology, 2022, 61, 1699-1708.	1.9	4
2	Gene Expression Profiles of Treatment Response and <scp>Nonâ€Response</scp> in Children With Juvenile Dermatomyositis. ACR Open Rheumatology, 2022, 4, 671-681.	2.1	4
3	Pilot Study of the Juvenile Dermatomyositis Consensus Treatment Plans: A CARRA Registry Study. Journal of Rheumatology, 2021, 48, 114-122.	2.0	9
4	Juvenile dermatomyositis: advances in clinical presentation, myositis-specific antibodies and treatment. World Journal of Pediatrics, 2020, 16, 31-43.	1.8	22
5	A Clinically and Biologically Based Subclassification of the Idiopathic Inflammatory Myopathies Using Machine Learning. ACR Open Rheumatology, 2020, 2, 158-166.	2.1	12
6	Management Considerations: Juvenile Dermatomyositis., 2020,, 285-298.		0
7	Interferon Chemokine Score and Other Cytokine Measures Track With Changes in Disease Activity in Patients With Juvenile and Adult Dermatomyositis. ACR Open Rheumatology, 2019, 1, 83-89.	2.1	10
8	Update on outcome assessment in myositis. Nature Reviews Rheumatology, 2018, 14, 303-318.	8.0	100
9	Brief Report: The Genetic Profile of Rheumatoid Factor–Positive Polyarticular Juvenile Idiopathic Arthritis Resembles That of Adult Rheumatoid Arthritis. Arthritis and Rheumatology, 2018, 70, 957-962.	5.6	53
10	T-cell transcriptomics from peripheral blood highlights differences between polymyositis and dermatomyositis patients. Arthritis Research and Therapy, 2018, 20, 188.	3.5	21
11	Predictors of changes in disease activity among children with juvenile dermatomyositis enrolled in the Childhood Arthritis and Rheumatology Research Alliance (CARRA) Legacy Registry. Clinical Rheumatology, 2018, 37, 1011-1015.	2.2	5
12	The Role of a Division of Quantitative Sciences Division in Enhancing Academic Productivity of a Department of Pediatrics. Journal of Pediatrics, 2017, 180, 4-5.	1.8	0
13	2016 American College of Rheumatology/European League Against Rheumatism Criteria for Minimal, Moderate, and Major Clinical Response in Juvenile Dermatomyositis. Annals of the Rheumatic Diseases, 2017, 76, 782-791.	0.9	51
14	2017 European League Against Rheumatism/American College of Rheumatology classification criteria for adult and juvenile idiopathic inflammatory myopathies and their major subgroups. Annals of the Rheumatic Diseases, 2017, 76, 1955-1964.	0.9	754
15	EULAR/ACR classification criteria for adult and juvenile idiopathic inflammatory myopathies and their major subgroups: a methodology report. RMD Open, 2017, 3, e000507.	3.8	115
16	2017 European League Against Rheumatism/American College of Rheumatology Classification Criteria for Adult and Juvenile Idiopathic Inflammatory Myopathies and Their Major Subgroups. Arthritis and Rheumatology, 2017, 69, 2271-2282.	5.6	391
17	Childhood Arthritis and Rheumatology Research Alliance Consensus Clinical Treatment Plans for Juvenile Dermatomyositis with Persistent Skin Rash. Journal of Rheumatology, 2017, 44, 110-116.	2.0	43
18	Cutaneous improvement in refractory adult and juvenile dermatomyositis after treatment with rituximab. Rheumatology, 2017, 56, 247-254.	1.9	82

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19	Gene Expression Profiling in Blood and Affected Muscle Tissues Reveals Differential Activation Pathways in Patients with New-onset Juvenile and Adult Dermatomyositis. Journal of Rheumatology, 2017, 44, 117-124.	2.0	25
20	2016 ACR-EULAR adult dermatomyositis and polymyositis and juvenile dermatomyositis response criteriaâ€"methodological aspects. Rheumatology, 2017, 56, 1884-1893.	1.9	33
21	Rilonacept maintains long-term inflammatory remission in patients with deficiency of the IL-1 receptor antagonist. JCI Insight, $2017, 2, .$	5.0	35
22	Brief Report: HLA–DRB1, DQA1, and DQB1 in Juvenileâ€Onset Systemic Sclerosis. Arthritis and Rheumatology, 2016, 68, 2772-2777.	5.6	15
23	Autoantibody levels in myositis patients correlate with clinical response during B cell depletion with rituximab. Rheumatology, 2016, 55, 991-999.	1.9	75
24	Dense genotyping of immune-related loci in idiopathic inflammatory myopathies confirms HLA alleles as the strongest genetic risk factor and suggests different genetic background for major clinical subgroups. Annals of the Rheumatic Diseases, 2016, 75, 1558-1566.	0.9	127
25	Adipokine gene expression in peripheral blood of adult and juvenile dermatomyositis patients and their relation to clinical parameters and disease activity measures. Journal of Inflammation, 2015, 12, 29.	3.4	16
26	Mineral Oil Aspiration Related Juvenile Idiopathic Arthritis. Case Reports in Pulmonology, 2015, 2015, 1-3.	0.3	0
27	Biologic predictors of clinical improvement in rituximab-treated refractory myositis. BMC Musculoskeletal Disorders, 2015, 16, 257.	1.9	42
28	Immunological Biomarkers in Dermatomyositis. Current Rheumatology Reports, 2015, 17, 68.	4.7	13
29	Autoantibodies to Dense Fine Speckles in Pediatric Diseases and Controls. Journal of Rheumatology, 2015, 42, 2419-2426.	2.0	34
30	Dermatologic Features of ADA2 Deficiency in Cutaneous Polyarteritis Nodosa. JAMA Dermatology, 2015, 151, 1230.	4.1	75
31	Interferon-regulated chemokine score associated with improvement in disease activity in refractory myositis patients treated with rituximab. Clinical and Experimental Rheumatology, 2015, 33, 655-63.	0.8	9
32	Clinical Characteristics of Children With Juvenile Dermatomyositis: The Childhood Arthritis and Rheumatology Research Alliance Registry. Arthritis Care and Research, 2014, 66, 404-410.	3.4	82
33	Pediatrics Practice at Mayo Clinic—A Historical Vignette. Mayo Clinic Proceedings, 2014, 89, e23-e25.	3.0	0
34	Predictors of Clinical Improvement in Rituximabâ€Treated Refractory Adult and Juvenile Dermatomyositis and Adult Polymyositis. Arthritis and Rheumatology, 2014, 66, 740-749.	5.6	210
35	Idiopathic Inflammatory Myopathies: Current Trends in Pathogenesis, Clinical Features, and Up-to-Date Treatment Recommendations. Mayo Clinic Proceedings, 2013, 88, 83-105.	3.0	145
36	Rituximab in the treatment of refractory adult and juvenile dermatomyositis and adult polymyositis: A randomized, placeboâ€phase trial. Arthritis and Rheumatism, 2013, 65, 314-324.	6.7	514

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37	BAFF Expression Correlates with Idiopathic Inflammatory Myopathy Disease Activity Measures and Autoantibodies. Journal of Rheumatology, 2013, 40, 294-302.	2.0	33
38	Increased expression of ADAMTS13 mRNA correlates with ischemic cerebrovascular disease in systemic lupus erythematosus patients. SAGE Open Medicine, 2013, 1, 205031211351440.	1.8	1
39	Changes in novel biomarkers of disease activity in juvenile and adult dermatomyositis are sensitive biomarkers of disease course. Arthritis and Rheumatism, 2012, 64, 4078-4086.	6.7	76
40	Consensus treatments for moderate juvenile dermatomyositis: Beyond the first two months. Results of the Second Childhood Arthritis and Rheumatology Research Alliance Consensus Conference. Arthritis Care and Research, 2012, 64, 546-553.	3.4	101
41	Protocols for the initial treatment of moderately severe juvenile dermatomyositis: Results of a Children's Arthritis and Rheumatology Research Alliance Consensus Conference. Arthritis Care and Research, 2010, 62, 219-225.	3.4	77
42	Treatment Approaches to Juvenile Dermatomyositis (JDM) Across North America: The Childhood Arthritis and Rheumatology Research Alliance (CARRA) JDM Treatment Survey. Journal of Rheumatology, 2010, 37, 1953-1961.	2.0	90
43	Interleukinâ€6 and type I interferon–regulated genes and chemokines mark disease activity in dermatomyositis. Arthritis and Rheumatism, 2009, 60, 3436-3446.	6.7	198
44	The inflammatory milieu in idiopathic inflammatory myositis. Current Rheumatology Reports, 2009, 11, 295-301.	4.7	20
45	Juvenile dermatomyositis and other idiopathic inflammatory myopathies of childhood. Lancet, The, 2008, 371, 2201-2212.	13.7	383
46	An Interferon Signature in the Peripheral Blood of Dermatomyositis Patients is Associated with Disease Activity. Molecular Medicine, 2007, 13, 59-68.	4.4	262
47	History of infection before the onset of juvenile dermatomyositis: Results from the National Institute of Arthritis and Musculoskeletal and Skin Diseases Research Registry. Arthritis and Rheumatism, 2005, 53, 166-172.	6.7	130
48	Recent advances in juvenile dermatomyositis. Current Rheumatology Reports, 2005, 7, 94-98.	4.7	6
49	Validation and clinical significance of the Childhood Myositis Assessment Scale for assessment of muscle function in the juvenile idiopathic inflammatory myopathies. Arthritis and Rheumatism, 2004, 50, 1595-1603.	6.7	195
50	Chimerism in myositis. Current Rheumatology Reports, 2003, 5, 421-424.	4.7	4
51	Microchimerism in children with rheumatic disorders: What does it mean?. Current Rheumatology Reports, 2003, 5, 458-462.	4.7	8
52	Genetic and environmental risk factors for idiopathic inflammatory myopathies. Rheumatic Disease Clinics of North America, 2002, 28, 891-916.	1.9	63
53	Juvenile Dermatomyositis. Paediatric Drugs, 2002, 4, 315-321.	3.1	32
54	Juvenile Dermatomyositis. Paediatric Drugs, 2002, 4, 315-321.	3.1	6

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55	Development of validated disease activity and damage indices for the juvenile idiopathic inflammatory myopathies: II. The childhood myositis assessment scale (CMAS): a quantitative tool for the evaluation of muscle function. Arthritis and Rheumatism, 1999, 42, 2213-2219.	6.7	194