

Sulaiman M Al-Mayouf

List of Publications by Year in Descending Order

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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.

81
papers

2,256
citations

19
h-index

46
g-index

85
ext. papers

2,834
ext. citations

4
avg, IF

4.21
L-index

#	Paper	IF	Citations
81	Epidemiology and demographics of juvenile idiopathic arthritis in Africa and Middle East. <i>Pediatric Rheumatology</i> , 2021 , 19, 166	3.5	1
80	Autosomal Recessive ISG15 Deficiency Underlies Type I Interferonopathy with Systemic Lupus Erythematosus and Inflammatory Myositis. <i>Journal of Clinical Immunology</i> , 2021 , 41, 1361-1364	5.7	1
79	Familial aggregation of juvenile idiopathic arthritis with other autoimmune diseases: Impact on clinical characteristics, disease activity status and disease damage. <i>International Journal of Rheumatic Diseases</i> , 2021 , 24, 1080-1085	2.3	0
78	International Consensus For The Dosing Of Corticosteroids In Childhood-Onset Systemic Lupus Erythematosus With Proliferative Lupus Nephritis. <i>Arthritis and Rheumatology</i> , 2021 ,	9.5	3
77	New or vanishing frontiers: associated juvenile arthritis. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2021 , 8, 44-47	1.6	1
76	Cumulative Damage in Juvenile Idiopathic Arthritis: A Multicenter Study From the Pediatric Rheumatology Arab Group. <i>Arthritis Care and Research</i> , 2021 , 73, 586-592	4.7	4
75	Lupus manifestations in children with primary immunodeficiency diseases: Comprehensive phenotypic and genetic features and outcome. <i>Modern Rheumatology</i> , 2021 , 31, 1171-1178	3.3	0
74	Familial Clustering of Juvenile Psoriatic Arthritis Associated with a Hemizygous FOXP3 Mutation. <i>Current Rheumatology Reports</i> , 2021 , 23, 64	4.9	0
73	Consensus clinical approach for a newly diagnosed systemic juvenile idiopathic arthritis among members of the pediatric rheumatology Arab group. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2021 , 8, 129-133	1.6	0
72	Systemic lupus erythematosus in a girl with PTEN variant and transaldolase deficiency: a novel phenotype. <i>Clinical Rheumatology</i> , 2020 , 39, 3511-3515	3.9	5
71	A Case with Purine Nucleoside Phosphorylase Deficiency Suffering from Late-Onset Systemic Lupus Erythematosus and Lymphoma. <i>Journal of Clinical Immunology</i> , 2020 , 40, 833-839	5.7	5
70	Efficacy of a sequential treatment by belimumab in monogenic systemic lupus erythematosus. <i>European Journal of Rheumatology</i> , 2020 , 7, 184-189	1.7	0
69	Stand-alone renal SLICC criterion with full house glomerular deposits: is it enough for childhood lupus nephritis?. <i>Clinical Rheumatology</i> , 2020 , 39, 401-405	3.9	1
68	Pattern and diagnostic evaluation of systemic autoinflammatory diseases other than familial Mediterranean fever among Arab children: a multicenter study from the Pediatric Rheumatology Arab Group (PRAG). <i>Rheumatology International</i> , 2020 , 40, 49-56	3.6	5
67	The Value of Screening for Celiac Disease in Systemic Lupus Erythematosus: A Single Experience of a Tertiary Medical Center. <i>Rheumatology and Therapy</i> , 2020 , 7, 649-656	4.4	1
66	Development and initial validation of a composite disease activity score for systemic juvenile idiopathic arthritis. <i>Rheumatology</i> , 2020 , 59, 3505-3514	3.9	16
65	Utility of serum ferritin and soluble interleukin-2 receptor as markers of disease activity in childhood systemic lupus erythematosus. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2020 , 7, 112-115	1.6	1

64	Outcome of children with systemic rheumatic diseases admitted to pediatric intensive care unit: An experience of a tertiary hospital. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2019 , 6, 142-145	1.6	1
63	Lessons Learned from Large-Scale, First-Tier Clinical Exome Sequencing in a Highly Consanguineous Population. <i>American Journal of Human Genetics</i> , 2019 , 104, 1182-1201	11	95
62	Phenotypic variability and disparities in treatment and outcomes of childhood arthritis throughout the world: an observational cohort study. <i>The Lancet Child and Adolescent Health</i> , 2019 , 3, 255-263	14.5	58
61	Early and late effects of therapeutic plasma exchange in patients with systemic lupus erythematosus and antineutrophil cytoplasmic antibody-associated vasculitis: A single-center experience. <i>Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia</i> , 2019 , 30, 775-780	0.6	
60	Noninflammatory disorders mimic juvenile idiopathic arthritis. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2018 , 5, 1-4	1.6	5
59	The Algerian Arabic version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). <i>Rheumatology International</i> , 2018 , 38, 27-33	3.6	3
58	Treating juvenile idiopathic arthritis to target: recommendations of an international task force. <i>Annals of the Rheumatic Diseases</i> , 2018 , 77, 819-828	2.4	99
57	The Arabic version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). <i>Rheumatology International</i> , 2018 , 38, 43-49	3.6	5
56	In silico validation of the Autoinflammatory Disease Damage Index. <i>Annals of the Rheumatic Diseases</i> , 2018 , 77, 1599-1605	2.4	17
55	The Libyan Arabic version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). <i>Rheumatology International</i> , 2018 , 38, 267-274	3.6	5
54	The Omani Arabic version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). <i>Rheumatology International</i> , 2018 , 38, 299-306	3.6	3
53	Combination of tacrolimus and mycophenolate mofetil in persistent proteinuria due to refractory childhood lupus nephritis. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2018 , 5, 99-102	1.6	3
52	Monogenic interferonopathies: Phenotypic and genotypic findings of CANDLE syndrome and its overlap with C1q deficient SLE. <i>International Journal of Rheumatic Diseases</i> , 2018 , 21, 208-213	2.3	18
51	Electrocardiographic disturbances in children with systemic lupus erythematosus. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2018 , 5, 127-130	1.6	5
50	Validation of the Cutaneous Lupus Erythematosus Disease Area and Severity Index and pSkindex27 for use in childhood-onset systemic lupus erythematosus. <i>Lupus Science and Medicine</i> , 2018 , 5, e000275	4.6	5
49	Juvenile idiopathic arthritis in multiplex families: longitudinal follow-up. <i>International Journal of Rheumatic Diseases</i> , 2017 , 20, 898-902	2.3	4
48	The landscape of genetic diseases in Saudi Arabia based on the first 1000 diagnostic panels and exomes. <i>Human Genetics</i> , 2017 , 136, 921-939	6.3	132
47	Phenotypic characteristics and outcome of juvenile dermatomyositis in Arab children. <i>Rheumatology International</i> , 2017 , 37, 1513-1517	3.6	4

46	Evolving spectrum of LRBA deficiency-associated chronic arthritis: is there a causative role in juvenile idiopathic arthritis?. <i>Clinical and Experimental Rheumatology</i> , 2017 , 35, 327-329	2.2	6
45	Biologic agents therapy for Saudi children with rheumatic diseases: indications and safety. <i>International Journal of Rheumatic Diseases</i> , 2016 , 19, 600-5	2.3	3
44	Coexistence of endocrinopathies in children with rheumatic diseases. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2016 , 3, 119-122	1.6	3
43	Marked variability in clinical presentation and outcome of patients with C1q immunodeficiency. <i>Journal of Autoimmunity</i> , 2015 , 62, 39-44	15.5	27
42	Dissecting the heterogeneity of macrophage activation syndrome complicating systemic juvenile idiopathic arthritis. <i>Journal of Rheumatology</i> , 2015 , 42, 994-1001	4.1	47
41	Vitamin D status in children with systemic lupus erythematosus and its association with clinical and laboratory parameters. <i>Clinical Rheumatology</i> , 2015 , 34, 81-4	3.9	20
40	The impact of antiphospholipid antibodies in children with lupus nephritis. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2015 , 2, 147-151	1.6	3
39	Association of a mutation in LACC1 with a monogenic form of systemic juvenile idiopathic arthritis. <i>Arthritis and Rheumatology</i> , 2015 , 67, 288-95	9.5	85
38	Familial aggregation of Crohn's disease and necrotizing sarcoid-like granulomatous disease. <i>European Journal of Rheumatology</i> , 2015 , 2, 122-124	1.7	2
37	Childhood systemic sarcoid-like necrotizing granulomatous disease: Another piece of the puzzle. <i>International Journal of Pediatrics and Adolescent Medicine</i> , 2014 , 1, 31-34	1.6	
36	Grave aortic aneurysmal dilatation in DOCK8 deficiency. <i>Modern Rheumatology</i> , 2014 , 24, 690-3	3.3	7
35	Health related quality of life measure in systemic pediatric rheumatic diseases and its translation to different languages: an international collaboration. <i>Pediatric Rheumatology</i> , 2014 , 12, 49	3.5	5
34	Study of Mendelian forms of Crohn's disease in Saudi Arabia reveals novel risk loci and alleles. <i>Gut</i> , 2014 , 63, 1831-2	19.2	23
33	Safety and efficacy of combined cyclophosphamide and rituximab treatment in recalcitrant childhood lupus. <i>Rheumatology International</i> , 2014 , 34, 529-33	3.6	17
32	Systemic lupus erythematosus in Saudi children: long-term outcomes. <i>International Journal of Rheumatic Diseases</i> , 2013 , 16, 56-60	2.3	6
31	Camptodactyly-arthropathy-coxavara-pericarditis syndrome in Saudi Arabia: clinical and molecular genetic findings in 22 patients. <i>Seminars in Arthritis and Rheumatism</i> , 2013 , 43, 292-6	5.3	18
30	Granulomatous disease in a child treated with etanercept. <i>International Journal of Rheumatic Diseases</i> , 2013 , 16, 472-4	2.3	5
29	Grave aortic aneurysmal dilatation in DOCK8 deficiency. <i>Modern Rheumatology</i> , 2013 , 1	3.3	3

28	A retrospective review of autoinflammatory diseases in Saudi children at a rheumatology clinic. <i>Annals of Saudi Medicine</i> , 2012 , 32, 43-8	1.6	9
27	Familial juvenile systemic lupus erythematosus in Arab children. <i>Rheumatology International</i> , 2012 , 32, 1939-43	3.6	6
26	Impact of C1q deficiency on the severity and outcome of childhood systemic lupus erythematosus. <i>International Journal of Rheumatic Diseases</i> , 2011 , 14, 81-5	2.3	22
25	Clinical and laboratory variables of childhood systemic lupus erythematosus in western province of Saudi Arabia. <i>Rheumatology International</i> , 2011 , 31, 23-6	3.6	5
24	Subcutaneous nodules, arthropathy, coarse face, cataract and glaucoma: a newly recognized syndrome. <i>Clinical Dysmorphology</i> , 2011 , 20, 50-52	0.9	
23	Loss-of-function variant in DNASE1L3 causes a familial form of systemic lupus erythematosus. <i>Nature Genetics</i> , 2011 , 43, 1186-8	36.3	259
22	Localized calcinosis in juvenile dermatomyositis: successful treatment with intralesional corticosteroids injection. <i>International Journal of Rheumatic Diseases</i> , 2010 , 13, e26-8	2.3	14
21	EULAR/PRINTO/PRES criteria for Henoch-Schönlein purpura, childhood polyarteritis nodosa, childhood Wegener granulomatosis and childhood Takayasu arteritis: Ankara 2008. Part II: Final classification criteria. <i>Annals of the Rheumatic Diseases</i> , 2010 , 69, 798-806	2.4	769
20	Tufting enteropathy and chronic arthritis: a newly recognized association with a novel EpCAM gene mutation. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2009 , 49, 642-4	2.8	36
19	Extensive ulcerations due to pyoderma gangrenosum in a child with juvenile systemic lupus erythematosus and C1q deficiency. <i>Annals of Saudi Medicine</i> , 2008 , 28, 466-8	1.6	8
18	Influence of gender and age of onset on the outcome in children with systemic lupus erythematosus. <i>Clinical Rheumatology</i> , 2008 , 27, 1159-62	3.9	31
17	Extensive ulcerations due to pyoderma gangrenosum in a child with juvenile systemic lupus erythematosus and C1q deficiency. <i>Annals of Saudi Medicine</i> , 2008 , 28, 466	1.6	4
16	Familial arthropathy in Saudi Arabian children: demographic, clinical, and biochemical features. <i>Seminars in Arthritis and Rheumatism</i> , 2007 , 36, 256-61	5.3	11
15	Cutaneous leukocytoclastic vasculitis associated with mycobacterial and salmonella infection. <i>Clinical Rheumatology</i> , 2007 , 26, 1563-4	3.9	5
14	Novel PRG4 mutations underlie CACP in Saudi families. <i>Human Mutation</i> , 2006 , 27, 213	4.7	24
13	Progressive sclerodermatous skin changes in a child with phenylketonuria. <i>Pediatric Dermatology</i> , 2006 , 23, 136-8	1.9	6
12	Sarcoidosis: a delayed or missed diagnosis in children. <i>Annals of Saudi Medicine</i> , 2006 , 26, 220-3	1.6	3
11	Comparison of clinical and laboratory variables in familial versus sporadic systemic onset juvenile idiopathic arthritis. <i>Journal of Rheumatology</i> , 2006 , 33, 597-600	4.1	6

10	Consequences of disease-causing mutations on lubricin protein synthesis, secretion, and post-translational processing. <i>Journal of Biological Chemistry</i> , 2005 , 280, 31325-32	5.4	50
9	Periodic fever, rash and arthropathy in a child. <i>Annals of Saudi Medicine</i> , 2005 , 25, 433, 441-2	1.6	3
8	The use of corticosteroid therapy in refractory Kawasaki patients. <i>Clinical Rheumatology</i> , 2004 , 23, 11-3	3.9	10
7	Pattern of neuropsychiatric manifestations and outcome in juvenile systemic lupus erythematosus. <i>Clinical Rheumatology</i> , 2004 , 23, 395-9	3.9	66
6	The Arabic version of childhood health assessment questionnaire modified for Arabic children. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2004 , 25, 83-7	1.1	11
5	Ocular manifestations of systemic lupus erythematosus in children. <i>Journal of King Abdulaziz University, Islamic Economics</i> , 2003 , 24, 964-6	1.1	16
4	New form of idiopathic osteolysis: nodulosis, arthropathy and osteolysis (NAO) syndrome. <i>American Journal of Medical Genetics Part A</i> , 2000 , 93, 5-10		46
3	Coexistent linear scleroderma and juvenile systemic lupus erythematosus. <i>Pediatric Dermatology</i> , 2000 , 17, 456-9	1.9	10
2	Hemophagocytosis complicating Kawasaki disease. <i>Pediatric Hematology and Oncology</i> , 2000 , 17, 323-9	1.7	27
1	Juvenile systemic lupus erythematosus in 60 Saudi children. <i>Annals of Saudi Medicine</i> , 1997 , 17, 612-5	1.6	13