Sulaiman M Al-Mayouf

List of Publications by Year
in descending order

84 papers	3,325 citations	304602 22 h-index	155592 55 g-index
85 all docs	85 docs citations	85 times ranked	4729 citing authors

#	Article	IF	Citations
1	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
2	Loss-of-function variant in DNASE1L3 causes a familial form of systemic lupus erythematosus. Nature Genetics, 2011, 43, 1186-1188.	9.4	366
3	The landscape of genetic diseases in Saudi Arabia based on the first 1000 diagnostic panels and exomes. Human Genetics, 2017, 136, 921-939.	1.8	209
4	Lessons Learned from Large-Scale, First-Tier Clinical Exome Sequencing in a Highly Consanguineous Population. American Journal of Human Genetics, 2019, 104, 1182-1201.	2.6	184
5	Treating juvenile idiopathic arthritis to target: recommendations of an international task force. Annals of the Rheumatic Diseases, 2018, 77, annrheumdis-2018-213030.	0.5	183
6	Phenotypic variability and disparities in treatment and outcomes of childhood arthritis throughout the world: an observational cohort study. The Lancet Child and Adolescent Health, 2019, 3, 255-263.	2.7	120
7	Association of a Mutation in LACC1 With a Monogenic Form of Systemic Juvenile Idiopathic Arthritis. Arthritis and Rheumatology, 2015, 67, 288-295.	2.9	111
8	Pattern of neuropsychiatric manifestations and outcome in juvenile systemic lupus erythematosus. Clinical Rheumatology, 2004, 23, 395-399.	1.0	82
9	Consequences of Disease-causing Mutations on Lubricin Protein Synthesis, Secretion, and Post-translational Processing. Journal of Biological Chemistry, 2005, 280, 31325-31332.	1.6	60
10	Dissecting the Heterogeneity of Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Journal of Rheumatology, 2015, 42, 994-1001.	1.0	59
11	New form of idiopathic osteolysis: Nodulosis, arthropathy and osteolysis (NAO) syndrome. American Journal of Medical Genetics Part A, 2000, 93, 5-10.	2.4	50
12	Influence of gender and age of onset on the outcome in children with systemic lupus erythematosus. Clinical Rheumatology, 2008, 27, 1159-1162.	1.0	39
13	Development and initial validation of a composite disease activity score for systemic juvenile idiopathic arthritis. Rheumatology, 2020, 59, 3505-3514.	0.9	39
14	Tufting Enteropathy and Chronic Arthritis: A Newly Recognized Association With a Novel EpCAM Gene Mutation. Journal of Pediatric Gastroenterology and Nutrition, 2009, 49, 642-644.	0.9	37
15	HEMOPHAGOCYTOSIS COMPLICATING KAWASAKI DISEASE. Pediatric Hematology and Oncology, 2000, 17, 323-329.	0.3	36
16	Marked variability in clinical presentation and outcome of patients with C1q immunodeficiency. Journal of Autoimmunity, 2015, 62, 39-44.	3.0	33
17	Novel PRG4 mutations underlie CACP in Saudi families. Human Mutation, 2006, 27, 213-213.	1.1	28
18	Study of Mendelian forms of Crohn's disease in Saudi Arabia reveals novel risk loci and alleles. Gut, 2014, 63, 1831-1832.	6.1	28

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19	In silico validation of the Autoinflammatory Disease Damage Index. Annals of the Rheumatic Diseases, 2018, 77, 1599-1605.	0.5	27
20	Impact of C1q deficiency on the severity and outcome of childhood systemic lupus erythematosus. International Journal of Rheumatic Diseases, 2011, 14, 81-85.	0.9	25
21	Monogenic interferonopathies: Phenotypic and genotypic findings of CANDLE syndrome and its overlap with C1q deficient SLE.. International Journal of Rheumatic Diseases, 2018, 21, 208-213.	0.9	25
22	Vitamin D status in children with systemic lupus erythematosus and its association with clinical and laboratory parameters. Clinical Rheumatology, 2015, 34, 81-84.	1.0	24
23	Camptodactyly-arthropathy-coxavara-pericarditis syndrome in Saudi Arabia: Clinical and molecular genetic findings in 22 patients. Seminars in Arthritis and Rheumatism, 2013, 43, 292-296.	1.6	22
24	Safety and efficacy of combined cyclophosphamide and rituximab treatment in recalcitrant childhood lupus. Rheumatology International, 2014, 34, 529-533.	1.5	22
25	Localized calcinosis in juvenile dermatomyositis: successful treatment with intralesional corticosteroids injection. International Journal of Rheumatic Diseases, 2010, 13, e26-8.	0.9	20
26	Juvenile Systemic Lupus Erythematosus in 60 Saudi Children. Annals of Saudi Medicine, 1997, 17, 612-615.	0.5	20
27	Ocular manifestations of systemic lupus erythematosus in children. Journal of King Abdulaziz University, Islamic Economics, 2003, 24, 964-6.	0.5	18
28	A Case with Purine Nucleoside Phosphorylase Deficiency Suffering from Late-Onset Systemic Lupus Erythematosus and Lymphoma. Journal of Clinical Immunology, 2020, 40, 833-839.	2.0	16
29	The use of corticosteroid therapy in refractory Kawasaki patients. Clinical Rheumatology, 2004, 23, 11-13.	1.0	14
30	International Consensus for the Dosing of Corticosteroids in <scp>Childhoodâ€Onset</scp> Systemic Lupus Erythematosus With Proliferative Lupus Nephritis. Arthritis and Rheumatology, 2022, 74, 263-273.	2.9	14
31	Epidemiology and demographics of juvenile idiopathic arthritis in Africa and Middle East. Pediatric Rheumatology, 2021, 19, 166.	0.9	14
32	Familial Arthropathy in Saudi Arabian Children: Demographic, Clinical, and Biochemical Features. Seminars in Arthritis and Rheumatism, 2007, 36, 256-261.	1.6	13
33	Coexistent Linear Scleroderma and Juvenile Systemic Lupusâ€ƒErythematosus. Pediatric Dermatology, 2000, 17, 456-459.	0.5	12
34	Coexistence of endocrinopathies in children with rheumatic diseases. International Journal of Pediatrics and Adolescent Medicine, 2016, 3, 119-122.	0.5	12
35	The Arabic version of childhood health assessment questionnaire modified for Arabic children. Journal of King Abdulaziz University, Islamic Economics, 2004, 25, 83-7.	0.5	12
36	Progressive Sclerodermatous Skin Changes in a Child with Phenylketonuria. Pediatric Dermatology, 2006, 23, 136-138.	0.5	11

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37	Noninflammatory disorders mimic juvenile idiopathic arthritis. International Journal of Pediatrics and Adolescent Medicine, 2018, 5, 1-4.	0.5	11
38	Clinical and laboratory variables of childhood systemic lupus erythematosus in western province of Saudi Arabia. Rheumatology International, 2011, 31, 23-26.	1.5	10
39	A retrospective review of autoinflammatory diseases in Saudi children at a rheumatology clinic. Annals of Saudi Medicine, 2012, 32, 43-48.	0.5	10
40	Phenotypic characteristics and outcome of juvenile dermatomyositis in Arab children. Rheumatology International, 2017, 37, 1513-1517.	1.5	10
41	Validation of the Cutaneous Lupus Erythematosus Disease Area and Severity Index and pSkindex27 for use in childhood-onset systemic lupus erythematosus. Lupus Science and Medicine, 2018, 5, e000275.	1.1	10
42	Systemic lupus erythematosus in a girl with PTEN variant and transaldolase deficiency: a novel phenotype. Clinical Rheumatology, 2020, 39, 3511-3515.	1.0	10
43	Cumulative Damage in Juvenile Idiopathic Arthritis: AÂMulticenter Study From the Pediatric Rheumatology ArabÂGroup. Arthritis Care and Research, 2021, 73, 586-592.	1.5	10
44	Extensive ulcerations due to pyoderma gangrenosum in a child with juvenile systemic lupus erythematosus and C1q deficiency. Annals of Saudi Medicine, 2008, 28, 466-468.	0.5	9
45	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). Rheumatology International, 2020, 40, 49-56.	1.5	9
46	Familial juvenile systemic lupus erythematosus in Arab children. Rheumatology International, 2012, 32, 1939-1943.	1.5	8
47	Grave aortic aneurysmal dilatation in DOCK8 deficiency. Modern Rheumatology, 2014, 24, 690-693.	0.9	8
48	The Arabic version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). Rheumatology International, 2018, 38, 43-49.	1.5	8
49	The Libyan Arabic version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). Rheumatology International, 2018, 38, 267-274.	1.5	8
50	Lupus manifestations in children with primary immunodeficiency diseases: Comprehensive phenotypic and genetic features and outcome. Modern Rheumatology, 2021, 31, 1171-1178.	0.9	8
51	Comparison of clinical and laboratory variables in familial versus sporadic systemic onset juvenile idiopathic arthritis. Journal of Rheumatology, 2006, 33, 597-600.	1.0	8
52	Systemic lupus erythematosus in Saudi children: longâ€term outcomes. International Journal of Rheumatic Diseases, 2013, 16, 56-60.	0.9	7
53	Granulomatous disease in a child treated with etanercept. International Journal of Rheumatic Diseases, 2013, 16, 472-474.	0.9	7
54	Electrocardiographic disturbances in children with systemic lupus erythematosus. International Journal of Pediatrics and Adolescent Medicine, 2018, 5, 127-130.	0.5	7

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55	New or vanishing frontiers: LACC1-associated juvenile arthritis. International Journal of Pediatrics and Adolescent Medicine, 2021, 8, 44-47.	0.5	7
56	Efficacy of a sequential treatment by belimumab in monogenic systemic lupus erythematosus. European Journal of Rheumatology, 2020, 7, 184-189.	1.3	7
57	Evolving spectrum of LRBA deficiency-associated chronic arthritis: is there a causative role in juvenile idiopathic arthritis?. Clinical and Experimental Rheumatology, 2017, 35, 327-329.	0.4	7
58	Cutaneous leukocytoclastic vasculitis associated with mycobacterial and salmonella infection. Clinical Rheumatology, 2007, 26, 1563-1564.	1.0	6
59	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
60	Biologic agents therapy for Saudi children with rheumatic diseases: indications and safety. International Journal of Rheumatic Diseases, 2016, 19, 600-605.	0.9	6
61	Combination of tacrolimus and mycophenolate mofetil in persistent proteinuria due to refractory childhood lupus nephritis. International Journal of Pediatrics and Adolescent Medicine, 2018, 5, 99-102.	0.5	6
62	The impact of antiphospholipid antibodies in children with lupus nephritis. International Journal of Pediatrics and Adolescent Medicine, 2015, 2, 147-151.	0.5	5
63	Juvenile idiopathic arthritis in multiplex families: longitudinal follow-up. International Journal of Rheumatic Diseases, 2017, 20, 898-902.	0.9	5
64	The Omani Arabic version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). Rheumatology International, 2018, 38, 299-306.	1.5	5
65	Sarcoidosis: a delayed or missed diagnosis in children. Annals of Saudi Medicine, 2006, 26, 220-223.	0.5	5
66	The Algerian Arabic version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). Rheumatology International, 2018, 38, 27-33.	1.5	4
67	The Value of Screening for Celiac Disease in Systemic Lupus Erythematosus: A Single Experience of a Tertiary Medical Center. Rheumatology and Therapy, 2020, 7, 649-656.	1.1	4
68	Autosomal Recessive ISG15 Deficiency Underlies Type I Interferonopathy with Systemic Lupus Erythematosus and Inflammatory Myositis. Journal of Clinical Immunology, 2021, 41, 1361-1364.	2.0	4
69	Familial aggregation of juvenile idiopathic arthritis with other autoimmune diseases: Impact on clinical characteristics, disease activity status and disease damage. International Journal of Rheumatic Diseases, 2021, 24, 1080-1085.	0.9	4
70	Extensive ulcerations due to pyoderma gangrenosum in a child with juvenile systemic lupus erythematosus and C1q deficiency. Annals of Saudi Medicine, 2008, 28, 466.	0.5	4
71	Performance of the EULAR/ACR 2019 classification criteria for systemic lupus erythematous in monogenic lupus. Clinical Rheumatology, 2022, 41, 2721-2727.	1.0	4
72	Grave aortic aneurysmal dilatation in DOCK8 deficiency. Modern Rheumatology, 2013, , 1.	0.9	3

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73	Utility of serum ferritin and soluble interleukin-2 receptor as markers of disease activity in childhood systemic lupus erythematosus. International Journal of Pediatrics and Adolescent Medicine, 2020, 7, 112-115.	0.5	3
74	Periodic fever, rash and arthropathy in a child. Annals of Saudi Medicine, 2005, 25, 433-433.	0.5	3
75	Childhood systemic sarcoid-like necrotizing granulomatous disease. International Journal of Pediatrics and Adolescent Medicine, 2014, 1, 31-34.	0.5	2
76	Outcome of children with systemic rheumatic diseases admitted to pediatric intensive care unit: An experience of a tertiary hospital. International Journal of Pediatrics and Adolescent Medicine, 2019, 6, 142-145.	0.5	2
77	Familial Clustering of Juvenile Psoriatic Arthritis Associated with a Hemizygous FOXP3 Mutation. Current Rheumatology Reports, 2021, 23, 64.	2.1	2
78	Consensus clinical approach for a newly diagnosed systemic juvenile idiopathic arthritis among members of the pediatric rheumatology Arab group. International Journal of Pediatrics and Adolescent Medicine, 2021, 8, 129-133.	0.5	2
79	Familial aggregation of Crohnâ€™s disease and necrotizing sarcoid-like granulomatous disease. European Journal of Rheumatology, 2015, 2, 122-124.	1.3	2
80	Stand-alone renal SLICC criterion with full house glomerular deposits: is it enough for childhood lupus nephritis?. Clinical Rheumatology, 2020, 39, 401-405.	1.0	1
81	Subcutaneous nodules, arthropathy, coarse face, cataract and glaucoma. Clinical Dysmorphology, 2011, 20, 50-52.	0.1	0
82	Early and late effects of therapeutic plasma exchange in patients with systemic lupus erythematosus and antineutrophil cytoplasmic antibody-associated vasculitis: A single-center experience. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2019, 30, 775.	0.4	0
83	Efficacy of a sequential treatment by belimumab in monogenic systemic lupus erythematosus. European Journal of Rheumatology, 2020, , .	1.3	0
84	A novel plasminogen mutation in a child with hereditary periodic syndrome: A case report. Rheumatology & Autoimmunity, 2022, 2, 237-243.	0.3	0