Åerife GÃ¼l KaradaÄ

List of Publications by Year
in descending order

43 papers	432 citations	840776 11 h-index	940533 16 g-index
44 all docs	44 docs citations	44 times ranked	515 citing authors

#	Article	IF	Citations
1	Differences and similarities of multisystem inflammatory syndrome in children, Kawasaki disease and macrophage activating syndrome due to systemic juvenile idiopathic arthritis: a comparative study. Rheumatology International, 2022, 42, 879-889.	3.0	35
2	Toward the integration of biosimilars into pediatric rheumatology: adalimumab ABP 501 experience of PeRA research group. Expert Opinion on Biological Therapy, 2022, 22, 197-202.	3.1	5
3	Is it all about age? Clinical characteristics of Kawasaki disease in the extremely young: PeRA research group experience. Postgraduate Medicine, 2022, 134, 429-434.	2.0	2
4	Comorbidities and phenotypeâ€“genotype correlation in children with familial Mediterranean fever. Rheumatology International, 2021, 41, 113-120.	3.0	30
5	Differential diagnosis portfolio of a pediatric rheumatologist: eight cases, eight stories. Clinical Rheumatology, 2021, 40, 769-774.	2.2	1
6	Comparison of the clinical diagnostic criteria and the results of the next-generation sequence gene panel in patients with monogenic systemic autoinflammatory diseases. Clinical Rheumatology, 2021, 40, 2327-2337.	2.2	9
7	The relevance of practical laboratory markers in predicting gastrointestinal and renal involvement in children with Henochâ€“SchÃ¶nlein Purpura. Postgraduate Medicine, 2021, 133, 272-277.	2.0	16
8	Response to â€˜How to define disease severity accurately in patients with familial Mediterranean feverâ€™. Rheumatology International, 2021, 41, 239-240.	3.0	0
9	The influence of carrying MEFV gene variants on juvenile systemic lupus erythematosus. Rheumatology International, 2021, 41, 157-161.	3.0	4
10	The Value of Serum Amyloid A Levels in Familial Mediterranean Fever to Identify Occult Inflammation During Asymptomatic Periods. Journal of Clinical Rheumatology, 2021, 27, 1-4.	0.9	9
11	Adherence to best practice consensus guidelines for familial Mediterranean fever: a modified Delphi study among paediatric rheumatologists in Turkey. Rheumatology International, 2021, , 1.	3.0	4
12	We might have the same mutation but my inflammasome beats your inflammasome: CINCA versus FCAS. ReumatologÃa ClÃnica, 2021, 17, 118-119.	0.5	0
13	Hepatitis B vaccination response of treatment-naive patients with juvenile idiopathic arthritis. Rheumatology International, 2021, , 1.	3.0	1
14	Age of onset as an influencing factor for disease severity in children with familial Mediterranean fever. Modern Rheumatology, 2021, 31, 219-222.	1.8	12
15	Comparison of Pediatric Familial Mediterranean Fever Patients Carrying Only E148Q Variant With the Ones Carrying Homozygous Pathogenic Mutations. Journal of Clinical Rheumatology, 2021, 27, 182-186.	0.9	7
16	Low disease activity state in juvenile-onset systemic lupus erythematosus. Lupus, 2021, 30, 2144-2150.	1.6	9
17	Real-Life Data From the Largest Pediatric Familial Mediterranean Fever Cohort. Frontiers in Pediatrics, 2021, 9, 805919.	1.9	22
18	The frequency of macrophage activation syndrome and disease course in systemic juvenile idiopathic arthritis. Modern Rheumatology, 2020, 30, 900-904.	1.8	12

#	Article	IF	Citations
19	Profile of new referrals to a single pediatric rheumatology center in Turkey. Rheumatology International, 2020, 40, 313-321.	3.0	9
20	Serum amyloid A as a biomarker in differentiating attacks of familial Mediterranean fever from acute febrile infections. Clinical Rheumatology, 2020, 39, 249-253.	2.2	6
21	Performance of Tel-Hashomer, Livneh, pediatric and new Eurofever/PRINTO classification criteria for familial Mediterranean fever in a referral center. Rheumatology International, 2020, 40, 21-27.	3.0	17
22	Drug reactions in children with rheumatic diseases receiving parenteral therapies: 9 yearsâ€™ experience of a tertiary pediatric rheumatology center. Rheumatology International, 2020, 40, 771-776.	3.0	5
23	ADA2 Deficiency: Case Series of Five Patients with Varying Phenotypes. Journal of Clinical Immunology, 2020, 40, 253-258.	3.8	17
24	Isotretinoinâ€induced sacroiliitis: Case series of four patients and a systematic review of the literature. Pediatric Dermatology, 2020, 37, 171-175.	0.9	5
25	Coexistence of Juvenile Systemic Lupus Erythematosus and Juvenile Spondyloarthropathy: A Case Report and Review of the Literature. Archives of Rheumatology, 2020, 35, 132-136.	0.9	0
26	Does immunosuppressive treatment entail an additional risk for children with rheumatic diseases? A survey-based study in the era of COVID-19. Rheumatology International, 2020, 40, 1613-1623.	3.0	32
27	Patient satisfaction and clinical effectiveness of switching from intravenous tocilizumab to subcutaneous tocilizumab in patients with juvenile idiopathic arthritis: an observational study. Rheumatology International, 2020, 40, 1111-1116.	3.0	8
28	Genetic panel screening in patients with clinically unclassified systemic autoinflammatory diseases. Clinical Rheumatology, 2020, 39, 3733-3745.	2.2	9
29	Rheumatic diseases in Syrian refugee children: a retrospective multicentric study in Turkey. Rheumatology International, 2020, 40, 583-589.	3.0	7
30	Characteristics of pediatric BehÃ§et's disease in Turkey and Israel: A cross-sectional cohort comparison. Seminars in Arthritis and Rheumatism, 2020, 50, 515-520.	3.4	18
31	Canakinumab in colchicine resistant familial mediterranean fever and other pediatric rheumatic diseases. Turkish Journal of Pediatrics, 2020, 62, 167.	0.6	10
32	Time to collaborate: Objectives, Design, and Methodology of PeRA-Research Group. Ä°stanbul Kuzey Klinikleri, 2020, 8, 200-202.	0.3	6
33	Leflunomide treatment in juvenile idiopathic arthritis. Rheumatology International, 2019, 39, 1615-1619.	3.0	16
34	Etiologic Spectrum and Follow-Up Results of Noninfectious Uveitis in Children: A Single Referral Center Experience. Archives of Rheumatology, 2019, 34, 294-300.	0.9	15
35	Why is the frequency of uveitis low in Turkish children with juvenile idiopathic arthritis?. Rheumatology, 2019, 59, 679-680.	1.9	2
36	The clinical spectrum of Henochâ€“SchÃ¶nlein purpura in children: a single-center study. Clinical Rheumatology, 2019, 38, 1707-1714.	2.2	30

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37	Atypical phenotype of an old disease or typical phenotype of a new disease: deficiency of adenosine deaminase 2. Turkish Journal of Pediatrics, 2019, 61, 413.	0.6	5
38	Clinical experiences in turkish paediatric patients with chronic recurrent multifocal osteomyelitis. Turkish Journal of Pediatrics, 2019, 61, 879.	0.6	7
39	Complete and sustained resolution of calcinosis universalis in a juvenile dermatomyositis case with mycophenolate mofetil. Turkish Journal of Pediatrics, 2019, 61, 771.	0.6	4
40	The necessity, efficacy and safety of biologics in juvenile idiopathic arthritis. Ä°stanbul Kuzey Klinikleri, 2019, 7, 118-123.	0.3	2
41	An extreme entity in differential diagnosis of musculoskeletal involvement-fibrodysplasia ossificans progressiva: a case based review. Turkish Journal of Pediatrics, 2018, 60, 593.	0.6	0
42	Two cases of periodic fever syndrome with coexistent mevalonate kinase and Mediterranean fever gene mutations. Turkish Journal of Pediatrics, 2017, 59, 467-470.	0.6	6
43	Subtype frequencies, demographic features, and remission rates in juvenile idiopathic arthritis - 265 cases from a Turkish center. Turkish Journal of Pediatrics, 2017, 59, 548-554.	0.6	18