

Nuray Aktay Ayaz

List of Publications by Year in descending order

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Version: 2024-02-01

123
papers

1,518
citations

430843

18
h-index

395678

33
g-index

127
all docs

127
docs citations

127
times ranked

1853
citing authors

#	ARTICLE	IF	CITATIONS
1	Performance of recent PRINTO criteria versus current ILAR criteria for systemic juvenile idiopathic arthritis: A single-centre experience. <i>Modern Rheumatology</i> , 2023, 33, 187-193.	1.8	4
2	Autoimmune and autoinflammatory diseases with mucocutaneous manifestations: A pediatric rheumatology perspective. <i>International Journal of Dermatology</i> , 2023, 62, 723-736.	1.0	1
3	Differences and similarities of multisystem inflammatory syndrome in children, Kawasaki disease and macrophage activating syndrome due to systemic juvenile idiopathic arthritis: a comparative study. <i>Rheumatology International</i> , 2022, 42, 879-889.	3.0	35
4	The clinical course of SARS-CoV-2 infection among children with rheumatic disease under biologic therapy: a retrospective and multicenter study. <i>Rheumatology International</i> , 2022, 42, 469-475.	3.0	16
5	Toward the integration of biosimilars into pediatric rheumatology: adalimumab ABP 501 experience of PeRA research group. <i>Expert Opinion on Biological Therapy</i> , 2022, 22, 197-202.	3.1	5
6	Validity and reliability of four parent/patient reported outcome measures for juvenile idiopathic arthritis remote monitoring. <i>Arthritis Care and Research</i> , 2022, , .	3.4	2
7	Exploring the attitudes, concerns, and knowledge regarding COVID-19 vaccine by the parents of children with rheumatic disease: Cross-sectional online survey. <i>Vaccine</i> , 2022, 40, 1829-1836.	3.8	7
8	Editorial: Hereditary Periodic Fevers and Autoinflammatory Diseases. <i>Frontiers in Pediatrics</i> , 2022, 10, 855738.	1.9	1
9	What is the Role of Mucocutaneous Manifestations in the Clinical Presentation of Monogenic Autoinflammatory Diseases? A Singlecenter Experience. <i>Bagcilar Medical Bulletin</i> , 2022, 7, 70-76.	0.1	0
10	The Multifaceted Presentation of the Multisystem Inflammatory Syndrome in Children: Data from a Cluster Analysis. <i>Journal of Clinical Medicine</i> , 2022, 11, 1742.	2.4	6
11	Is There an Association Between Initial Clinical Manifestations and the Development of Macrophage Activation Syndrome in Patients with Systemic Juvenile Idiopathic Arthritis?. <i>Medical Journal of Bakirkoy</i> , 2022, 18, 31-36.	0.1	0
12	Humoral response and safety of BNT162b2 mRNA vaccine in children with rheumatic diseases. <i>Rheumatology</i> , 2022, 61, 4482-4490.	1.9	14
13	Is it all about age? Clinical characteristics of Kawasaki disease in the extremely young: PeRA research group experience. <i>Postgraduate Medicine</i> , 2022, 134, 429-434.	2.0	2
14	Embracing Change: An International Survey Study on the Beliefs and Attitudes of Pediatric Rheumatologists Towards Biosimilars. <i>BioDrugs</i> , 2022, , 1.	4.6	0
15	Cluster Analysis of Pediatric Behçet's Disease: Data from The Pediatric Rheumatology Academy (PeRA)-Research Group (RG). <i>Modern Rheumatology</i> , 2022, , .	1.8	3
16	Comorbidities and phenotypeâ€“genotype correlation in children with familial Mediterranean fever. <i>Rheumatology International</i> , 2021, 41, 113-120.	3.0	30
17	Differential diagnosis portfolio of a pediatric rheumatologist: eight cases, eight stories. <i>Clinical Rheumatology</i> , 2021, 40, 769-774.	2.2	1
18	Comparison of the clinical diagnostic criteria and the results of the next-generation sequence gene panel in patients with monogenic systemic autoinflammatory diseases. <i>Clinical Rheumatology</i> , 2021, 40, 2327-2337.	2.2	9

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19	Systemic lupus erythematosus complicated with Castleman disease: a case-based review. Rheumatology International, 2021, 41, 475-479.	3.0	11
20	The relevance of practical laboratory markers in predicting gastrointestinal and renal involvement in children with Henoch-Schönlein Purpura. Postgraduate Medicine, 2021, 133, 272-277.	2.0	16
21	Response to “How to define disease severity accurately in patients with familial Mediterranean fever”. Rheumatology International, 2021, 41, 239-240.	3.0	0
22	The influence of carrying MEFV gene variants on juvenile systemic lupus erythematosus. Rheumatology International, 2021, 41, 157-161.	3.0	4
23	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
24	The readiness of pediatric rheumatology patients and their parents to transition to adult-oriented treatment. International Journal of Rheumatic Diseases, 2021, 24, 397-401.	1.9	7
25	Evaluation of Children Referred to Pediatric Rheumatology Outpatient Clinic with Suspicious Laboratory Test Results. İstanbul Kanuni Sultan Süleyman Tıp Dergisi, 2021, , .	0.0	0
26	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
27	Neuroimaging of Children With Takayasu Arteritis. Journal of Child Neurology, 2021, 36, 642-647.	1.4	2
28	We might have the same mutation but my inflammasome beats your inflammasome: CINCA versus FCAS. Reumatologia Clínica, 2021, 17, 118-119.	0.5	0
29	Hepatitis B vaccination response of treatment-naïve patients with juvenile idiopathic arthritis. Rheumatology International, 2021, , 1.	3.0	1
30	Comment on: Clinical significance of E148Q heterozygous variant in paediatric Familial Mediterranean Fever. Rheumatology, 2021, 60, e294-e295.	1.9	1
31	Approach to switching biologics in juvenile idiopathic arthritis: a real-life experience. Rheumatology International, 2021, , 1.	3.0	4
32	Sacroiliitis in children and adolescents with familial Mediterranean fever. Advances in Rheumatology, 2021, 61, 29.	1.7	4
33	Hematological involvement in pediatric systemic lupus erythematosus: A multi-center study. Lupus, 2021, 30, 1983-1990.	1.6	9
34	Nailfold capillaroscopy: A sensitive method for evaluating microvascular involvement in children with SARS-CoV-2 infection. Microvascular Research, 2021, 138, 104196.	2.5	14
35	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
36	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

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37	Low disease activity state in juvenile-onset systemic lupus erythematosus. <i>Lupus</i> , 2021, 30, 2144-2150.	1.6	9
38	Real-Life Data From the Largest Pediatric Familial Mediterranean Fever Cohort. <i>Frontiers in Pediatrics</i> , 2021, 9, 805919.	1.9	22
39	The feasibility of withdrawing canakinumab in paediatric colchicine-resistant familial Mediterranean fever patients. <i>Clinical and Experimental Rheumatology</i> , 2021, 39 Suppl 132, 118-123.	0.8	0
40	The feasibility of withdrawing canakinumab in paediatric colchicine-resistant familial Mediterranean fever patients. <i>Clinical and Experimental Rheumatology</i> , 2021, 39, 118-123.	0.8	4
41	The frequency of macrophage activation syndrome and disease course in systemic juvenile idiopathic arthritis. <i>Modern Rheumatology</i> , 2020, 30, 900-904.	1.8	12
42	Profile of new referrals to a single pediatric rheumatology center in Turkey. <i>Rheumatology International</i> , 2020, 40, 313-321.	3.0	9
43	Serum amyloid A as a biomarker in differentiating attacks of familial Mediterranean fever from acute febrile infections. <i>Clinical Rheumatology</i> , 2020, 39, 249-253.	2.2	6
44	Performance of Tel-Hashomer, Livneh, pediatric and new Eurofever/PRINTO classification criteria for familial Mediterranean fever in a referral center. <i>Rheumatology International</i> , 2020, 40, 21-27.	3.0	17
45	Drug reactions in children with rheumatic diseases receiving parenteral therapies: 9 years' experience of a tertiary pediatric rheumatology center. <i>Rheumatology International</i> , 2020, 40, 771-776.	3.0	5
46	ADA2 Deficiency: Case Series of Five Patients with Varying Phenotypes. <i>Journal of Clinical Immunology</i> , 2020, 40, 253-258.	3.8	17
47	Isotretinoin-induced sacroiliitis: Case series of four patients and a systematic review of the literature. <i>Pediatric Dermatology</i> , 2020, 37, 171-175.	0.9	5
48	Coexistence of Juvenile Systemic Lupus Erythematosus and Juvenile Spondyloarthritis: A Case Report and Review of the Literature. <i>Archives of Rheumatology</i> , 2020, 35, 132-136.	0.9	0
49	Does immunosuppressive treatment entail an additional risk for children with rheumatic diseases? A survey-based study in the era of COVID-19. <i>Rheumatology International</i> , 2020, 40, 1613-1623.	3.0	32
50	Patient satisfaction and clinical effectiveness of switching from intravenous tocilizumab to subcutaneous tocilizumab in patients with juvenile idiopathic arthritis: an observational study. <i>Rheumatology International</i> , 2020, 40, 1111-1116.	3.0	8
51	Comorbidities of antiphospholipid syndrome and systemic lupus erythematosus in children. <i>Current Rheumatology Reports</i> , 2020, 22, 21.	4.7	4
52	Genetic panel screening in patients with clinically unclassified systemic autoinflammatory diseases. <i>Clinical Rheumatology</i> , 2020, 39, 3733-3745.	2.2	9
53	Rheumatic diseases in Syrian refugee children: a retrospective multicentric study in Turkey. <i>Rheumatology International</i> , 2020, 40, 583-589.	3.0	7
54	Characteristics of pediatric Behçet's disease in Turkey and Israel: A cross-sectional cohort comparison. <i>Seminars in Arthritis and Rheumatism</i> , 2020, 50, 515-520.	3.4	18

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55	The evaluation of anxiety, depression and quality of life scores of children and adolescents with familial Mediterranean fever. <i>Rheumatology International</i> , 2020, 40, 757-763.	3.0	18
56	Comment on "Age dependent safety and efficacy of colchicine treatment for familial Mediterranean fever in children". <i>Seminars in Arthritis and Rheumatism</i> , 2020, 50, 1552.	3.4	0
57	Canakinumab in colchicine resistant familial mediterranean fever and other pediatric rheumatic diseases. <i>Turkish Journal of Pediatrics</i> , 2020, 62, 167.	0.6	10
58	How useful are Kawasaki disease risk scoring systems to the Turkish population?. <i>Anatolian Journal of Cardiology</i> , 2020, 24, 97-106.	0.9	10
59	Towards a combined pediatric rheumatology-dermatology clinic: One-year experience. <i>İstanbul Kuzey Klinikleri</i> , 2020, 8, 37-41.	0.3	0
60	Immunodeficiency-Like Phenotype, Recurrent Pulmonary Manifestations, and Persistent Polyarthritis: Mevalonate Kinase Deficiency Successfully Treated With Adalimumab. <i>Archives of Rheumatology</i> , 2020, 35, 627-628.	0.9	1
61	Time to collaborate: Objectives, Design, and Methodology of PeRA-Research Group. <i>İstanbul Kuzey Klinikleri</i> , 2020, 8, 200-202.	0.3	6
62	Like "North Americans", "Europeans", or "Others": Where do Turkish children with juvenile idiopathic arthritis stand in the new classification system?. <i>İstanbul Kuzey Klinikleri</i> , 2020, 8, 421-422.	0.3	0
63	Anterior Segment Analysis and Evaluation of Corneal Biomechanical Properties in Children with Joint Hypermobility. <i>Türk Oftalmoloji Dergisi</i> , 2020, 50, 71-74.	0.9	3
64	Otoinflamatuvar Periyodik Ateş Sendromları. <i>The Journal of Child</i> , 2020, 20, .	0.2	0
65	Leflunomide treatment in juvenile idiopathic arthritis. <i>Rheumatology International</i> , 2019, 39, 1615-1619.	3.0	16
66	Abatacept as a Long-Term Targeted Therapy for LRBA Deficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 2790-2800.e15.	3.8	112
67	Comment on: Short-term follow-up results of children with familial Mediterranean fever after cessation of colchicine: is it possible to quit?: reply. <i>Rheumatology</i> , 2019, 58, 1886-1887.	1.9	0
68	Short-term follow-up results of children with familial Mediterranean fever after cessation of colchicine: is it possible to quit?. <i>Rheumatology</i> , 2019, 58, 1818-1821.	1.9	12
69	Etiologic Spectrum and Follow-Up Results of Noninfectious Uveitis in Children: A Single Referral Center Experience. <i>Archives of Rheumatology</i> , 2019, 34, 294-300.	0.9	15
70	Why is the frequency of uveitis low in Turkish children with juvenile idiopathic arthritis?. <i>Rheumatology</i> , 2019, 59, 679-680.	1.9	2
71	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.	5.6	120
72	The clinical spectrum of Henoch-Schönlein purpura in children: a single-center study. <i>Clinical Rheumatology</i> , 2019, 38, 1707-1714.	2.2	30

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73	Diagnostic utility of a targeted next-generation sequencing gene panel in the clinical suspicion of systemic autoinflammatory diseases: a multi-center study. Rheumatology International, 2019, 39, 911-919.	3.0	37
74	THU0291â€¦THE CHARACTERISTICS OF PEDIATRIC BEHÇETâ€™S DISEASE IN TURKEY VERSUS ISRAEL. , 2019, , .		0
75	SAT0522â€¦COMPARISON OF CHILDREN CARRYING E148Q VARIANT WITH CHILDREN CARRYING HOMOZYGOUS PATHOGENIC VARIANTS. , 2019, , .		0
76	AB0991â€¦PRELIMINARY RESULTS OF REFERRALS TO A TERTIARY PEDIATRIC RHEUMATOLOGY OUTPATIENT CLINIC: A YEAR IN REVIEW. , 2019, , .		0
77	AB0990â€¦FINAL DIAGNOSIS OF THE PATIENTS WITH MUSCULOSKELETAL COMPLAINTS: PRELIMINARY RESULTS OF ONE-YEAR STUDY. , 2019, , .		0
78	AB1055â€¦FINAL DIAGNOSES OF THE PATIENTS WHO WERE REFERRED TO A TERTIARY PEDIATRIC RHEUMATOLOGY OUTPATIENT CLINIC FOR LABORATORY ABNORMALITIES. , 2019, , .		0
79	AB0594â€¦THE CLINICAL SPECTRUM OF HENOCH-SCHÖNLEIN PURPURA IN CHILDREN: A PROSPECTIVE SINGLE-CENTER STUDY. , 2019, , .		0
80	AB1061â€¦SHORT TERM FOLLOW-UP RESULTS OF CHILDREN WITH FAMILIAL MEDITERRANEAN FEVER AFTER CESSATION OF COLCHICINE: IS IT POSSIBLE TO QUIT?. , 2019, , .		0
81	FRI0536â€¦FAMILIAL MEDITERRANEAN FEVER (FMF): A SINGLE CENTER EXPERIENCE FROM TURKEY. , 2019, , .		1
82	THU0524â€¦ARE CHILDREN AND ADULTS HAVING DIFFERENT PHENOTYPE AND GENOTYPE OF FMF?. , 2019, , .		1
83	SAT0483â€¦COMPARISON OF THE CLINICAL DIAGNOSTIC CRITERIA AND RESULTS OF THE NEXT GENERATION SEQUENCE GENE PANEL IN PATIENTS WITH PERIODIC FEVER. , 2019, , .		0
84	FRI0556â€¦GENETIC SCREENING IN PATIENTS WITH UNDIFFERENTIATED PERIODIC FEVER SYNDROME. , 2019, , .		0
85	Corticosteroid-resistant anakinra-responsive protracted febrile myalgia syndrome as the first manifestation of familial Mediterranean fever. İstanbul Kuzey Klinikleri, 2019, 7, 78-80.	0.3	5
86	Clinical experiences in turkish paediatric patients with chronic recurrent multifocal osteomyelitis. Turkish Journal of Pediatrics, 2019, 61, 879.	0.6	7
87	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
88	The necessity, efficacy and safety of biologics in juvenile idiopathic arthritis. İstanbul Kuzey Klinikleri, 2019, 7, 118-123.	0.3	2
89	The Turkish version of the Juvenile Arthritis Multidimensional Assessment Report (JAMAR). Rheumatology International, 2018, 38, 395-402.	3.0	4
90	Does familial Mediterranean fever affect cognitive function in children? Electrophysiological preliminary study. International Journal of Neuroscience, 2018, 128, 10-14.	1.6	4

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91	Sustained hyperferritinemia in a child with macrophage activation syndrome secondary to systemic juvenile idiopathic arthritis - perforinopathy: case based review. Turkish Journal of Pediatrics, 2018, 60, 598.	0.6	2
92	A Case of Kawasaki Disease With Severe Lip and Oral Mucosa Involvement Complicated With Microstomia and Corrected With Surgery. Archives of Rheumatology, 2018, 33, 238-240.	0.9	5
93	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
94	Is there any difference regarding atopy between children with familial Mediterranean fever and healthy controls?. Allergologia Et Immunopathologia, 2017, 45, 549-552.	1.7	7
95	Development and Initial Validation of the Macrophage Activation Syndrome/Primary Hemophagocytic Lymphohistiocytosis Score, a Diagnostic Tool that Differentiates Primary Hemophagocytic Lymphohistiocytosis from Macrophage Activation Syndrome. Journal of Pediatrics, 2017, 189, 72-78.e3.	1.8	50
96	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
97	Leptospirosis in a child with acute respiratory distress syndrome. Turkish Journal of Pediatrics, 2017, 59, 688.	0.6	0
98	Comparison of the efficacy of once- and twice-daily colchicine dosage in pediatric patients with familial Mediterranean fever â€” a randomized controlled noninferiority trial. Arthritis Research and Therapy, 2016, 18, 85.	3.5	18
99	Kawasaki disease shock syndrome: a rare and severe complication of Kawasaki disease. Turkish Journal of Pediatrics, 2016, 58, 415-418.	0.6	6
100	Coexistence of early onset sarcoidosis and partial interferon-Î³ receptor 1 deficiency. Turkish Journal of Pediatrics, 2016, 58, 545-549.	0.6	7
101	A case of chickenpox complicated with subacute osteomyelitis. Marmara Medical Journal, 2016, 29, 110.	0.8	1
102	Orbital muscle involvement in a child with familial Mediterranean fever. Marmara Medical Journal, 2016, 29, 124.	0.8	1
103	A novel assessment tool for clinical care of patients with autoinflammatory disease: juvenile autoinflammatory disease multidimensional assessment report. Clinical and Experimental Rheumatology, 2016, 34, 129-135.	0.8	22
104	Paravertebral and Retroperitoneal Vascular Tumour Presenting with Kasabach-Merritt Phenomenon in Childhood, Diagnosed with Magnetic Resonance Imaging. Case Reports in Pediatrics, 2015, 2015, 1-4.	0.4	4
105	How Pricing And Reimbursement Policies Affect The Budget Impact of The Treatment of Systemic Juvenile Idiopathic Arthritis In Turkey. Value in Health, 2015, 18, A643.	0.3	0
106	Dissecting the Heterogeneity of Macrophage Activation Syndrome Complicating Systemic Juvenile Idiopathic Arthritis. Journal of Rheumatology, 2015, 42, 994-1001.	2.0	59
107	How do we encounter rare factor deficiencies in children? Single-centre results from Turkey. Blood Coagulation and Fibrinolysis, 2015, 26, 145-151.	1.0	4
108	Cochlear functions in children with familial Mediterranean fever: Any role of the severity of the disease?. International Journal of Pediatric Otorhinolaryngology, 2015, 79, 1566-1570.	1.0	8

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109	Granulocyte Transfusion in Febrile Neutropenia. Blood, 2015, 126, 4612-4612.	1.4	0
110	Dissecting the heterogeneity of macrophage activation syndrome. Pediatric Rheumatology, 2014, 12, .	2.1	0
111	Familial Mediterranean Fever: Diagnosing as Early as 3 Months of Age. Case Reports in Pediatrics, 2014, 2014, 1-3.	0.4	5
112	Cardiac T2* MRI assessment in patients with thalassaemia major and its effect on the preference of chelation therapy. International Journal of Hematology, 2014, 99, 706-713.	1.6	6
113	Surgical interventions in childhood rare factor deficiencies. Blood Coagulation and Fibrinolysis, 2013, 24, 854-861.	1.0	4
114	Factor VII Deficiency. Clinical and Applied Thrombosis/Hemostasis, 2012, 18, 588-593.	1.7	19
115	Time to focus on outcome assessment tools for childhood vasculitis. Pediatric Rheumatology, 2011, 9, 29.	2.1	4
116	Anti-Interleukin 1 Treatment for Patients with Familial Mediterranean Fever Resistant to Colchicine: Table 1.. Journal of Rheumatology, 2011, 38, 516-518.	2.0	132
117	The distribution of juvenile idiopathic arthritis in the eastern Mediterranean: results from the registry of the Turkish Paediatric Rheumatology Association. Clinical and Experimental Rheumatology, 2011, 29, 111-6.	0.8	35
118	Anti-IL-1 treatment for secondary amyloidosis in an adolescent with FMF and Behçet's disease. Clinical Rheumatology, 2010, 29, 209-210.	2.2	94
119	Preventing tuberculosis in children receiving anti-tnf treatment. Clinical Rheumatology, 2010, 29, 389-392.	2.2	20
120	Behçet disease: treatment of vascular involvement in children. European Journal of Pediatrics, 2010, 169, 427-430.	2.7	33
121	Genotoxicity of anti-tumor necrosis factor therapy in patients with juvenile idiopathic arthritis. Arthritis Care and Research, 2010, 62, 73-77.	3.4	12
122	Musculoskeletal sonography in juvenile systemic lupus erythematosus. Arthritis and Rheumatism, 2009, 61, 58-60.	6.7	27
123	Hyperimmunoglobulinemia D and periodic fever syndrome; treatment with etanercept and follow-up. Clinical Rheumatology, 2008, 27, 1317-1320.	2.2	55