# Rabia Miray Kisla Ekinci

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/590960/publications.pdf

Version: 2024-02-01

| 53 | 364 | 933410 10 | 996954 15 |
|---|---|---|---|
| papers | citations | h-index | g-index |

| 53 | 53 | 53 | 596 |
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| all docs | docs citations | times ranked | citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Cold-induced Vascular Changes in the Macula, Optic Disc and Choroid in Children with Raynaudâ€™s Phenomena. Ocular Immunology and Inflammation, 2022, , 1-9. | 1.8 | 0 |
| 2 | Serum vitamin B12 and D levels in children with Primary Raynaud Phenomenon: a retrospective cohort study. European Journal of Clinical Nutrition, 2022, 76, 1615-1617. | 2.9 | 1 |
| 3 | Clinical presentation of children with Deficiency of Adenosine deaminase 2: A case series. European Journal of Medical Genetics, 2022, 65, 104555. | 1.3 | 0 |
| 4 | Altered expression of apoptosis-related, circulating cell-free miRNAs in children with familial Mediterranean fever: a cross-sectional study. Rheumatology International, 2021, 41, 103-111. | 3.0 | 8 |
| 5 | Twenty-Year Experience of a Single Referral Center on Pediatric Familial Mediterranean Fever. Journal of Clinical Rheumatology, 2021, 27, 18-24. | 0.9 | 10 |
| 6 | Camptodactyly-Arthropathy-Coxa Vara-Pericarditis Syndrome Resembling Juvenile Idiopathic Arthritis: A Single-Center Experience from Southern Turkey. Molecular Syndromology, 2021, 12, 112-117. | 0.8 | 5 |
| 7 | Clinical features of childhood uveitis at a tertiary referral center in Southern Turkey. International Ophthalmology, 2021, 41, 2073-2081. | 1.4 | 6 |
| 8 | Monogenic lupus due to DNASE1L3 deficiency in a pediatric patient with urticarial rash, hypocomplementemia, pulmonary hemorrhage, and immune-complex glomerulonephritis. European Journal of Medical Genetics, 2021, 64, 104262. | 1.3 | 8 |
| 9 | Experience of autoimmune and autoinflammatory diseases in a Turkish pediatric cohort with primary immunodeficiencies. Allergologia Et Immunopathologia, 2021, 49, 1-7. | 1.7 | 2 |
| 10 | Serum lubricin levels in patients with juvenile idiopathic arthritis. Reumatologia, 2021, 59, 373-377. | 1.1 | 3 |
| 11 | Differentiating children with familial Mediterranean fever from other recurrent fever syndromes: The utility of new Eurofever/PRINTO classification criteria. Archives of Rheumatology, 2021, 36, 493-498. | 0.9 | 2 |
| 12 | Deficiency of adenosine deaminase 2: a case series revealing clinical manifestations, genotypes and treatment outcomes from Turkey. Rheumatology, 2020, 59, 254-256. | 1.9 | 4 |
| 13 | Phenotypic variability in two patients with tumor necrosis factor receptor associated periodic fever syndrome emphasizes a rare manifestation: Immunoglobulin A nephropathy. European Journal of Medical Genetics, 2020, 63, 103780. | 1.3 | 4 |
| 14 | Baricitinib experience on STING-associated vasculopathy with onset in infancy: A representative case from Turkey. Clinical Immunology, 2020, 212, 108273. | 3.2 | 38 |
| 15 | Clinical manifestations and outcomes of 420 children with Henoch SchÃ¶nlein Purpura from a single referral center from Turkey: A three-year experience. Modern Rheumatology, 2020, 30, 1039-1046. | 1.8 | 25 |
| 16 | Unicentric Castleman Disease Mimicking an Autoinflammatory Disorder: A Diagnostic Challenge in a Pediatric Patient With Recurrent Fever. Journal of Pediatric Hematology/Oncology, 2020, 42, 204-207. | 0.6 | 4 |
| 17 | Evaluation of different classification criteria in children with Behcet disease: results from a single referral center. Expert Review of Clinical Immunology, 2020, 16, 1093-1097. | 3.0 | 4 |
| 18 | Pre-Pulseless Takayasu Arteritis in a Child Represented With Prolonged Fever of Unknown Origin and Successful Management With Concomitant Mycophenolate Mofetil and Infliximab. Archives of Rheumatology, 2020, 35, 278-282. | 0.9 | 3 |

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|---|---|---|---|
| 19 | Growth Parameters of Turkish Children With an Autoinflammatory Disease Before and After Canakinumab Treatment. Indian Pediatrics, 2020, 57, 637-640. | 0.4 | 3 |
| 20 | Recurrent macroscopic hematuria in a pediatric patient: is it early to diagnose as having type I hereditary C2 deficiency?. CEN Case Reports, 2020, 9, 344-346. | 0.9 | 0 |
| 21 | The contribution of diet preference to the disease course in children with familial Mediterranean fever: a cross-sectional study. Reumatologia, 2020, 58, 81-86. | 1.1 | 7 |
| 22 | Retrospective analyzes of adverse events during biologic agents in children with juvenile idiopathic arthritis from aÂsingle center in Turkey. Reumatologia, 2020, 58, 367-374. | 1.1 | 2 |
| 23 | Different clinical manifestations of three prime repair exonuclease 1 mutation: A case series. Annals of Indian Academy of Neurology, 2020, 23, 699. | 0.5 | 0 |
| 24 | Juvenil idiopatik artritli TÃ¼rk Ã§ocuklarÄ±n bÃ¼yÃ¼me parametreleri. Cukurova Medical Journal, 2020, 45, 495-501. | 0.2 | 0 |
| 25 | Retrospective Analysis of the Factors Affecting Growth Parameters in Turkish Children With Systemic Lupus Erythematosus. Archives of Rheumatology, 2020, 35, 357-365. | 0.9 | 0 |
| 26 | Ã‡ocukluk Ã§aÄŸÄ± baÅŸlangÄ±Ã§lÄ± sistemik lupus eritematozus hastalarÄ±nda avaskÃ¼ler nekrozis sÄ±klÄ±ÄŸÄ±, Cukurova Medical Journal, 2020, 45, 200-207. | 0.2 | 1 |
| 27 | Henoch SchÃ¶nlein PurpurasÄ± tanÄ±lÄ± Ã§ocuklarda Kompleman C2 gen polimorfizmleri. Cukurova Medical Journal, 2020, 45, 89-95. | 0.2 | 0 |
| 28 | Is Henochâ€“SchÃ¶nlein purpura a susceptibility factor for functional gastrointestinal disorders in children?. Rheumatology International, 2019, 39, 317-322. | 3.0 | 8 |
| 29 | Symptomatic multifocal avascular necrosis in an adolescent with neuropsychiatric systemic lupus erythematosus. Reumatologia, 2019, 57, 182-187. | 1.1 | 7 |
| 30 | Disease Severity and Genotype Affect Physical Growth in Children With Familial Mediterranean Fever. Archives of Rheumatology, 2019, 34, 288-293. | 0.9 | 7 |
| 31 | Etanercept for the Treatment of Chronic Arthritis Related to Chronic Granulomatous Disease: A Case. Pediatric, Allergy, Immunology, and Pulmonology, 2019, 32, 131-134. | 0.8 | 4 |
| 32 | Canakinumab in Children with Familial Mediterranean Fever: A Single-Center, Retrospective Analysis. Paediatric Drugs, 2019, 21, 389-395. | 3.1 | 15 |
| 33 | A homozygote novel L451W mutation in CECR1 gene causes deficiency of adenosine deaminase 2 in a pediatric patient representing with chronic lymphoproliferation and cytopenia. Pediatric Hematology and Oncology, 2019, 36, 376-381. | 0.8 | 7 |
| 34 | Frequency of functional gastrointestinal disorders in children with familial Mediterranean fever. Clinical Rheumatology, 2019, 38, 921-926. | 2.2 | 9 |
| 35 | Hyperphosphatemic Familial Tumoral Calcinosis in Two Siblings with a Novel Mutation in <i>GALNT3</i> Gene: Experience from Southern Turkey. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 94-99. | 0.9 | 8 |
| 36 | Autoimmune Manifestations in Heterozygote Type I Complement 2 Deficiency: A Child Eventually Diagnosed With Systemic Lupus Erythematosus. Archives of Rheumatology, 2019, 34, 96-99. | 0.9 | 4 |

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|---|---|---|---|
| 37 | Do practical laboratory indices predict the outcomes of children with Henoch-SchÃ¶nlein purpura?. Postgraduate Medicine, 2019, 131, 295-298. | 2.0 | 24 |
| 38 | Hyperimmunoglobulinemia D syndrome with recurrent perianal abscess successfully treated with canakinumab. Scottish Medical Journal, 2019, 64, 103-107. | 1.3 | 1 |
| 39 | Recurrent Henoch SchÃ¶nlein purpura without renal involvement successfully treated with methotrexate. Scottish Medical Journal, 2019, 64, 74-77. | 1.3 | 9 |
| 40 | Juvenile systemic lupus erythematosus: a single-center experience from southern Turkey. Clinical Rheumatology, 2019, 38, 1459-1468. | 2.2 | 22 |
| 41 | MEFV gene variants in children with Henoch-SchÃ¶nlein purpura and association with clinical manifestations: a single-center Mediterranean experience. Postgraduate Medicine, 2019, 131, 68-72. | 2.0 | 14 |
| 42 | Ailevi Akdeniz AteÅŸi Olan Ã‡ocuklarda Ortalama Trombosit Hacminin Klinik Ã–nemi. Acibadem Universitesi Saglik Bilimleri Dergisi, 2019, , 0-0. | 0.1 | 0 |
| 43 | IgA eksikliÄŸinin juvenil dermatomiyozit prognozuna olumlu etkisi var mÄ±dÄ±r? olgu sunumu.. Cukurova Medical Journal, 2019, 44, 1511-1514. | 0.2 | 0 |
| 44 | Factors Affecting Physical Growth in Children with Primary Vesicoureteral Reflux: A Single Center Experience. Journal of Pediatric Research, 2019, 6, 186-191. | 0.2 | 0 |
| 45 | Decreased serum vitamin B12 and vitamin D levels affect sleep quality in children with familial Mediterranean fever. Rheumatology International, 2018, 38, 83-87. | 3.0 | 10 |
| 46 | The Influence of Concomitant Disorders on Disease Severity of Familial Mediterranean Fever in Children. Archives of Rheumatology, 2018, 33, 282-287. | 0.9 | 15 |
| 47 | Renal Amyloidosis in Deficiency of Adenosine Deaminase 2: Successful Experience With Canakinumab. Pediatrics, 2018, 142, . | 2.1 | 23 |
| 48 | Ã‡ocukluk Ã§aÄŸÄ± miks konnektif doku hastalÄ±ÄŸÄ±: olgu serisi. Cukurova Medical Journal, 2018, 43, 235-240. | 0.2 | 0 |
| 49 | Juvenil idiopatik inflamatuar miyopati: tek merkez verileri. Cukurova Medical Journal, 2018, 43, 685-691. | 0.2 | 1 |
| 50 | A homozygote TREX1 mutation in two siblings with different phenotypes: Chilblains and cerebral vasculitis. European Journal of Medical Genetics, 2017, 60, 690-694. | 1.3 | 13 |
| 51 | Risk Factors for Candidemia in Pediatric Intensive Care Unit Patients. Indian Journal of Pediatrics, 2014, 81, 1158-1162. | 0.8 | 10 |
| 52 | Alpha lipoic acid intoxication, treatment and outcome. Clinical Toxicology, 2013, 51, 522-522. | 1.9 | 13 |
| 53 | Multisystem Inflammatory Syndrome Occurring Simultaneously in Two Siblings. Indian Journal of Pediatrics, 0, , . | 0.8 | 0 |