# Ekrem Unal

List of Publications by Year
in descending order

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|---|---|---|---|
| 156 papers | 2,570 citations | 236612 25 h-index | 243296 44 g-index |
| 159 all docs | 159 docs citations | 159 times ranked | 4493 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | The extended clinical phenotype of 64 patients with dedicator of cytokinesis 8 deficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 402-412. | 1.5 | 163 |
| 2 | Hypomorphic homozygous mutations in phosphoglucomutase 3 (PGM3) impair immunity and increase serum IgE levels. Journal of Allergy and Clinical Immunology, 2014, 133, 1410-1419.e13. | 1.5 | 160 |
| 3 | Hematologically important mutations: Leukocyte adhesion deficiency (first update). Blood Cells, Molecules, and Diseases, 2012, 48, 53-61. | 0.6 | 147 |
| 4 | Genetic, immunological, and clinical features of patients with bacterial and fungal infections due to inherited IL-17RA deficiency. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, E8277-E8285. | 3.3 | 137 |
| 5 | Comparison of primary human cytotoxic T-cell and natural killer cell responses reveal similar molecular requirements for lytic granule exocytosis but differences in cytokine production. Blood, 2013, 121, 1345-1356. | 0.6 | 122 |
| 6 | Functional Selectivity in Cytokine Signaling Revealed Through a Pathogenic EPO Mutation. Cell, 2017, 168, 1053-1064.e15. | 13.5 | 98 |
| 7 | Inherited biallelic CSF3R mutations in severe congenital neutropenia. Blood, 2014, 123, 3811-3817. | 0.6 | 79 |
| 8 | Defective glycosylation and multisystem abnormalities characterize the primary immunodeficiency XMEN disease. Journal of Clinical Investigation, 2019, 130, 507-522. | 3.9 | 74 |
| 9 | Outcomes and Treatment Strategies for Autoimmunity and Hyperinflammation in Patients with RAG Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1970-1985.e4. | 2.0 | 64 |
| 10 | Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. Blood, 2020, 136, 2638-2655. | 0.6 | 64 |
| 11 | CD137 deficiency causes immune dysregulation with predisposition to lymphomagenesis. Blood, 2019, 134, 1510-1516. | 0.6 | 52 |
| 12 | Proteome Analysis of Human Neutrophil Granulocytes From Patients With Monogenic Disease Using Data-independent Acquisition. Molecular and Cellular Proteomics, 2019, 18, 760-772. | 2.5 | 52 |
| 13 | A case of XMEN syndrome presented with severe auto-immune disorders mimicking autoimmune lymphoproliferative disease. Clinical Immunology, 2015, 159, 58-62. | 1.4 | 41 |
| 14 | Childhood Stroke: Results of 130 Children From a Reference Center in Central Anatolia, Turkey. Pediatric Neurology, 2014, 50, 595-600. | 1.0 | 38 |
| 15 | Rapamycin has a beneficial effect on controlling epilepsy in children with tuberous sclerosis complex: results of 7 children from a cohort of 86. Child's Nervous System, 2014, 30, 227-240. | 0.6 | 37 |
| 16 | Targeted high-throughput sequencing for genetic diagnostics of hemophagocytic lymphohistiocytosis. Genome Medicine, 2015, 7, 130. | 3.6 | 37 |
| 17 | Novel Mutations in RASGRP1 are Associated with Immunodeficiency, Immune Dysregulation, and EBV-Induced Lymphoma. Journal of Clinical Immunology, 2018, 38, 699-710. | 2.0 | 37 |
| 18 | Blood repellent superhydrophobic surfaces constructed from nanoparticle-free and biocompatible materials. Colloids and Surfaces B: Biointerfaces, 2021, 205, 111864. | 2.5 | 35 |

| # | Article | IF | Citations |
|---|---|---|---|
| 19 | Genetic Deficiency and Biochemical Inhibition of ITK Affect Human Th17, Treg, and Innate Lymphoid Cells. Journal of Clinical Immunology, 2019, 39, 391-400. | 2.0 | 34 |
| 20 | Apparent diffusion coefficient in differentiation of pediatric posterior fossa tumors. Japanese Journal of Radiology, 2017, 35, 448-453. | 1.0 | 32 |
| 21 | Stem cell mobilization and collection from pediatric patients and healthy children. Transfusion and Apheresis Science, 2015, 53, 17-22. | 0.5 | 31 |
| 22 | Atypical Severe Combined Immunodeficiency Caused by a Novel Homozygous Mutation In Rag1 Gene in a Girl who Presented with Pyoderma Gangrenosum: A Case Report and Literature Review. Journal of Clinical Immunology, 2014, 34, 792-795. | 2.0 | 30 |
| 23 | Torticollis in children: an alert symptom not to be turned away. Child's Nervous System, 2015, 31, 1461-1470. | 0.6 | 30 |
| 24 | Late-type vitamin K deficiency bleeding: experience from 120 patients. Child's Nervous System, 2012, 28, 247-251. | 0.6 | 28 |
| 25 | Intracranial hemorrhage in children with congenital factor deficiencies. Child's Nervous System, 2011, 27, 1963-1966. | 0.6 | 27 |
| 26 | Atypical presentation of chronic granulomatous disease in an adolescent boy with frontal lobe located Aspergillus abscess mimicking intracranial tumor. Child's Nervous System, 2010, 26, 149-154. | 0.6 | 26 |
| 27 | CARMIL2 Deficiency Presenting as Very Early Onset Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2019, 25, 1788-1795. | 0.9 | 26 |
| 28 | Astroblastoma in a child. Child's Nervous System, 2008, 24, 165-168. | 0.6 | 25 |
| 29 | Hemophagocytic syndrome in a 4â€monthâ€old infant with biotinidase deficiency. Pediatric Blood and Cancer, 2012, 59, 191-193. | 0.8 | 23 |
| 30 | Human FCHO1 deficiency reveals role for clathrin-mediated endocytosis in development and function of T cells. Nature Communications, 2020, 11, 1031. | 5.8 | 23 |
| 31 | Autoimmune diseases detected in children with primary immunodeficiency diseases: results from a reference centre at middle anatolia. Acta Microbiologica Et Immunologica Hungarica, 2012, 59, 343-353. | 0.4 | 22 |
| 32 | Whole exome sequencing (WES) approach for diagnosing primary immunodeficiencies (PIDs) in a highly consanguineous community. Clinical Immunology, 2020, 214, 108376. | 1.4 | 22 |
| 33 | Web-based survey of resources for treatment and long-term follow-up for children with brain tumors in developing countries. Child's Nervous System, 2011, 27, 1957-1961. | 0.6 | 21 |
| 34 | Mesenchymal Hamartoma of the Liver Mimicking Hepatoblastoma. Journal of Pediatric Hematology/Oncology, 2008, 30, 458-460. | 0.3 | 20 |
| 35 | The efficacy of Pelargonium sidoides in the treatment of upper respiratory tract infections in children with transient hypogammaglobulinemia of infancy. Phytomedicine, 2012, 19, 958-961. | 2.3 | 20 |
| 36 | Expanded circulating hematopoietic stem/progenitor cells as novel cell source for the treatment of TCIRG1 osteopetrosis. Haematologica, 2020, 106, 74-86. | 1.7 | 20 |

| # | Article | IF | Citations |
|---|---|---|---|
| 37 | Evolution and longâ€term outcomes of combined immunodeficiency due to CARMIL2 deficiency. Allergy: European Journal of Allergy and Clinical Immunology, 2022, 77, 1004-1019. | 2.7 | 19 |
| 38 | Diffusion weighted imaging in differentiating malignant and benign neuroblastic tumors. Japanese Journal of Radiology, 2016, 34, 620-624. | 1.0 | 17 |
| 39 | ILC3 deficiency and generalized ILC abnormalities in DOCK8â€deficient patients. Allergy: European Journal of Allergy and Clinical Immunology, 2020, 75, 921-932. | 2.7 | 17 |
| 40 | Merkel Cell Carcinoma in a Child. Journal of Pediatric Hematology/Oncology, 2009, 31, 359-361. | 0.3 | 16 |
| 41 | Diagnostic and prognostic significance of glypicanÂ5 and glypicanÂ6 gene expression levels in gastric adenocarcinoma. Molecular and Clinical Oncology, 2015, 3, 584-590. | 0.4 | 16 |
| 42 | Malignant glioblastomatous transformation of a low-grade glioma in a child. Child's Nervous System, 2008, 24, 1385-1389. | 0.6 | 15 |
| 43 | CARDIAC FUNCTIONS EVALUATED WITH TISSUE DOPPLER IMAGING IN CHILDHOOD CANCERS TREATED WITH ANTHRACYCLINES. Pediatric Hematology and Oncology, 2010, 27, 13-23. | 0.3 | 15 |
| 44 | Intracranial hemorrhage in infants as a serious, and preventable consequence of late form of vitamin K deficiency: a selfie picture of Turkey, strategies for tomorrow. Child's Nervous System, 2014, 30, 1375-1382. | 0.6 | 15 |
| 45 | Brain Abscesses in Children. Journal of Child Neurology, 2015, 30, 458-467. | 0.7 | 15 |
| 46 | A case of congenital afibrinogenemia complicated with thromboembolic events that required repeated amputations. Blood Coagulation and Fibrinolysis, 2015, 26, 354-356. | 0.5 | 14 |
| 47 | The relationship between hematological parameters and prognosis of children with acute ischemic stroke. Child's Nervous System, 2018, 34, 655-661. | 0.6 | 14 |
| 48 | Cerebral venous sinus thrombosis in an adolescent with Ewing sarcoma. Child's Nervous System, 2008, 24, 983-986. | 0.6 | 13 |
| 49 | The Efficacy of Vitamin K2 and Calcitriol Combination on Thalassemic Osteopathy. Journal of Pediatric Hematology/Oncology, 2013, 35, 623-627. | 0.3 | 13 |
| 50 | Multiple Fungal Brain Abscesses in a Child with Acute Lymphoblastic Leukemia. Mycopathologia, 2012, 174, 505-509. | 1.3 | 12 |
| 51 | Gastric Signet Ring Carcinoma in a Patient With Ataxia-Telangiectasia. Journal of Pediatric Hematology/Oncology, 2013, 35, e341-e343. | 0.3 | 12 |
| 52 | Is there relation between COL4A1/A2 mutations and antenatally detected fetal intraventricular hemorrhage?. Child's Nervous System, 2014, 30, 419-424. | 0.6 | 12 |
| 53 | Acute Colchicine Intoxication Complicated With Extramedullary Hematopoiesis Due to Filgrastim in a Child. Journal of Pediatric Hematology/Oncology, 2014, 36, e460-e462. | 0.3 | 12 |
| 54 | Genetic Analysis of a Cohort of 275 Patients with Hyper-IgE Syndromes and/or Chronic Mucocutaneous Candidiasis. Journal of Clinical Immunology, 2021, 41, 1804-1838. | 2.0 | 12 |

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|---|---|---|---|
| 55 | Propranolol treatment for chylothorax due to diffuse lymphangiomatosis. Pediatric Blood and Cancer, 2019, 66, e27592. | 0.8 | 11 |
| 56 | Juvenile Myelomonocytic Leukemia in Turkey: A Retrospective Analysis of Sixty-five PatientsÃ–zlem TÃ¼fekÃ§i1. Turkish Journal of Haematology, 2018, 35, 27-34. | 0.2 | 11 |
| 57 | Neuroblastoma Arising From an Unresected Sacrococcygeal Teratoma in a Child. Journal of Pediatric Hematology/Oncology, 2010, 32, 233-235. | 0.3 | 10 |
| 58 | Chronic granulomatous disease with markedly elevated IgE levels mimicking hyperimmunoglobulin E syndrome. Acta Microbiologica Et Immunologica Hungarica, 2013, 60, 155-162. | 0.4 | 10 |
| 59 | X-linked severe combined immunodeficiency due to a novel mutation complicated with hemophagocytic lymphohistiocytosis and presented with invagination: A case report. European Journal of Microbiology and Immunology, 2014, 4, 174-176. | 1.5 | 10 |
| 60 | Cerebral Sinus Venous Thrombosis and Prothrombotic Risk Factors in Children: A Single-Center Experience From Turkey. Journal of Pediatric Hematology/Oncology, 2018, 40, e369-e372. | 0.3 | 10 |
| 61 | Quartzâ€crystal Microbalance Measurements of CD19 Antibody Immobilization on Gold Surface and Capturing B Lymphoblast Cells: Effect of Surface Functionalization. Electroanalysis, 2018, 30, 834-841. | 1.5 | 10 |
| 62 | A Rare Case of Activated Phosphoinositide 3-Kinase Delta Syndrome (APDS) Presenting With Hemophagocytosis Complicated With Hodgkin Lymphoma. Journal of Pediatric Hematology/Oncology, 2020, 42, 156-159. | 0.3 | 10 |
| 63 | Effectiveness of Ankaferd BloodStopper in Prophylaxis and Treatment of Oral Mucositis in Childhood Cancers Evaluated with Plasma Citrulline Levels. Turkish Journal of Haematology, 2018, 35, 85-86. | 0.2 | 10 |
| 64 | Fatal valproate overdose in a newborn baby. Human and Experimental Toxicology, 2007, 26, 453-456. | 1.1 | 9 |
| 65 | Thyroid Medullary Carcinoma in a Teenager With Cowden Syndrome. Laryngoscope, 2007, 117, 1180-1182. | 1.1 | 9 |
| 66 | Pituitary?adrenal axis suppression due to topical steroid administration in an infant. Pediatrics International, 2007, 49, 242-244. | 0.2 | 9 |
| 67 | An uncommon extrapulmonary sequestration located in the upper posterior mediastinum associated with the azygos lobe in a child. Journal of Thoracic and Cardiovascular Surgery, 2007, 133, 1110-1111. | 0.4 | 9 |
| 68 | Release of N-terminal pro-brain natriuretic peptide in children with acute rheumatic carditis. Cardiology in the Young, 2010, 20, 297-301. | 0.4 | 9 |
| 69 | Cerebellar hemangioblastoma associated with diffuse neonatal hemangiomatosis in an infant. Child's Nervous System, 2012, 28, 1801-1805. | 0.6 | 9 |
| 70 | Inflammatory Myofibroblastic Tumor of the Kidney and Bilateral Lung Nodules in a Child Mimicking Wilms Tumor With Lung Metastases. Journal of Pediatric Hematology/Oncology, 2015, 37, e390-e393. | 0.3 | 9 |
| 71 | Neuroblastoma in a Child With Wolf-Hirschhorn Syndrome. Journal of Pediatric Hematology/Oncology, 2017, 39, e224-e226. | 0.3 | 9 |
| 72 | Microfluidic Chip based direct triple antibody immunoassay for monitoring patient comparative response to leukemia treatment. Biomedical Microdevices, 2020, 22, 48. | 1.4 | 9 |

| # | Article | IF | Citations |
|---|---|---|---|
| 73 | Factor 8 Gene Mutation Spectrum of 270 Patients with Haemophilia A: Identification of 36 Novel Mutations. Turkish Journal of Haematology, 2020, 37, 145-153. | 0.2 | 9 |
| 74 | Preclinical Studies on Convalescent Human Immune Plasma-Derived Exosome: Omics and Antiviral Properties to SARS-CoV-2. Frontiers in Immunology, 2022, 13, 824378. | 2.2 | 9 |
| 75 | Mycophenolate mofetil-induced pseudotumor cerebri in a boy with autoimmune lymphoproliferative disease. Child's Nervous System, 2011, 27, 853-855. | 0.6 | 8 |
| 76 | The importance of MTHFR polymorphisms in pediatric cerebral stroke. Child's Nervous System, 2012, 28, 13-13. | 0.6 | 8 |
| 77 | Fetal intracranial hemorrhage related to maternal autoimmune thrombocytopenic purpura. Child's Nervous System, 2014, 30, 2147-2150. | 0.6 | 8 |
| 78 | Hepatosplenic Fungal Infections in Children With Leukemiaâ€"Risk Factors and Outcome: A Multicentric Study. Journal of Pediatric Hematology/Oncology, 2019, 41, 256-260. | 0.3 | 8 |
| 79 | Refractory and Fatal Presentation of Severe Autoimmune Hemolytic Anemia in a Child With the DNASE1L3 Mutation Complicated With an Additional DOCK8 Variant. Journal of Pediatric Hematology/Oncology, 2021, 43, e452-e456. | 0.3 | 7 |
| 80 | A Newborn With Familial Hemophagocytic Lymphohistiocytosis Complicated With Transfusion Associated Graft Versus Host Disease. Journal of Pediatric Hematology/Oncology, 2017, 39, e309-e311. | 0.3 | 6 |
| 81 | Common Variable Immunodeficiency, Autoimmune Hemolytic Anemia, and Pancytopenia Associated With a Defect in IKAROS. Journal of Pediatric Hematology/Oncology, 2021, 43, e351-e357. | 0.3 | 6 |
| 82 | Discovery of the First Pathogenic Human EPO Mutation Provides Mechanistic Insight into Cytokine Signaling. Blood, 2016, 128, 331-331. | 0.6 | 6 |
| 83 | Hashimoto thyroiditis associated with ataxia telangiectasia. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, . | 0.4 | 5 |
| 84 | Orbital myositis associated with focal active colitis in a teenage girl. Child's Nervous System, 2012, 28, 641-643. | 0.6 | 5 |
| 85 | Infantile tremor syndrome associated with cobalamin therapy: A case report. Clinical Neurology and Neurosurgery, 2013, 115, 1903-1905. | 0.6 | 5 |
| 86 | Outcome of autologous hematopoietic stem cell transplantation in children and adolescents with relapsed or refractory Hodgkin's lymphoma. Pediatric Transplantation, 2015, 19, 745-752. | 0.5 | 5 |
| 87 | Pediatric central nervous system tumors in the first 3Âyears of life: pre-operative mean platelet volume, neutrophil/lymphocyte count ratio, and white blood cell count correlate with the presence of a central nervous system tumor. Child's Nervous System, 2017, 33, 233-238. | 0.6 | 5 |
| 88 | The relationship between the prognosis of children with acute arterial stroke and polymorphisms of CDKN2B, HDAC9, NINJ2, NAA25 genes. Journal of Thrombosis and Thrombolysis, 2019, 47, 578-584. | 1.0 | 5 |
| 89 | Different Clinical Presentation of 3 Children With Familial Hemophagocytic Lymphohistiocytosis With 2 Novel Mutations. Journal of Pediatric Hematology/Oncology, 2020, 42, e627-e629. | 0.3 | 5 |
| 90 | Hepatitis-associated aplastic anemia in pediatric patients: single center experience. Transfusion and Apheresis Science, 2020, 59, 102900. | 0.5 | 5 |

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|---|---|---|---|
| 91 | Immunomagnetic separation of B type acute lymphoblastic leukemia cells from bone marrow with flow cytometry validation and microfluidic chip measurements. Separation Science and Technology, 2021, 56, 2659-2666. | 1.3 | 5 |
| 92 | Effect of vitamin K2 and vitamin D3 on bone mineral density in children with acute lymphoblastic leukemia: a prospective cohort study. Journal of Pediatric Endocrinology and Metabolism, 2021, 34, 441-447. | 0.4 | 5 |
| 93 | Vena Cava Superior Syndrome in Children With Mediastinal Tumors: Single Center Experience. Ä°stanbul Kuzey Klinikleri, 2019, 7, 255-259. | 0.1 | 5 |
| 94 | Pulmonary arterial pressure in infants with laryngomalacia. International Journal of Pediatric Otorhinolaryngology, 2006, 70, 2067-2071. | 0.4 | 4 |
| 95 | Hypothermia in a Child With Hodgkin Disease. Journal of Pediatric Hematology/Oncology, 2009, 31, 136-138. | 0.3 | 4 |
| 96 | A Case of Familial Hemophagocytic Lymphohistiocytosis Type 4 With Involvement of the Central Nervous System Complicated With Infarct. Journal of Pediatric Hematology/Oncology, 2017, 39, e321-e324. | 0.3 | 4 |
| 97 | Capturing B type acute lymphoblastic leukemia cells using two types of antibodies. Biotechnology Progress, 2019, 35, e2737. | 1.3 | 4 |
| 98 | Role of a second transplantation for children with acute leukemia following posttransplantation relapse: a study by the Turkish Bone Marrow Transplantation Study Group. Leukemia and Lymphoma, 2020, 61, 1465-1474. | 0.6 | 4 |
| 99 | A novel missense mutation outside the <scp>DNAJ</scp> domain of <scp>DNAJC21</scp> is associated with <scp>Shwachmanâ€“Diamond</scp> syndrome. British Journal of Haematology, 2022, 197, . | 1.2 | 4 |
| 100 | Letter to the Editor. Inhalation Toxicology, 2007, 19, 587-587. | 0.8 | 3 |
| 101 | KLUVER-BUCY SYNDROME IN A BOY WITH NON-HODGKIN LYMPHOMA. Pediatric Hematology and Oncology, 2007, 24, 149-152. | 0.3 | 3 |
| 102 | Pilocytic astrocytoma developing at the site of a previously treated medulloblastoma in a child. Child's Nervous System, 2008, 24, 289-292. | 0.6 | 3 |
| 103 | Accidental intramuscular overdose administration of vincristine. Drug and Chemical Toxicology, 2012, 35, 232-234. | 1.2 | 3 |
| 104 | Atrial fibrillation as an uncommon presentation in a large pleomorphic xanthoastrocytoma. Child's Nervous System, 2012, 28, 475-479. | 0.6 | 3 |
| 105 | Myeloperoxidase Deficiency: The Secret Under the Flag of Unstained Cell. Turkish Journal of Haematology, 2013, 30, 232-233. | 0.2 | 3 |
| 106 | Neurological complication of non Hodgkin lymphoma in childhood: experience from a single center in Turkey. Child's Nervous System, 2014, 30, 639-645. | 0.6 | 3 |
| 107 | Cranial metastatic alveolar rhabdomyosarcoma mimicking hematological malignancy in an adolescent boy. Child's Nervous System, 2014, 30, 1737-1741. | 0.6 | 3 |
| 108 | Autosomal recessive hyper IgM syndrome associated with activation-induced cytidine deaminase gene in three Turkish siblings presented with tuberculosis lymphadenitis â€" Case report. Acta Microbiologica Et Immunologica Hungarica, 2015, 62, 267-274. | 0.4 | 3 |

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|---|---|---|---|
| 109 | Malignancies in Primary Immunodeficiencies: A Single Center Experience. Pediatric, Allergy, Immunology, and Pulmonology, 2015, 28, 47-54. | 0.3 | 3 |
| 110 | Fibromuscular Dysplasia Complicated With Cerebral Stroke in a Child With Congenital Dyserythropoietic Anemia Type II. Journal of Pediatric Hematology/Oncology, 2016, 38, e333-e335. | 0.3 | 3 |
| 111 | Familial moyamoya disease in two Turkish siblings with same polymorphism in RNF213 gene but different clinical features. Child's Nervous System, 2016, 32, 569-573. | 0.6 | 3 |
| 112 | Evaluation of childhood pancreas solid pseudopapillary tumors. Ä°stanbul Kuzey Klinikleri, 2017, 5, 207-210. | 0.1 | 3 |
| 113 | Genetic Polymorphism of VKORC1-1639 in Children With Intracranial Hemorrhage Due to Vitamin K Deficiency. Clinical and Applied Thrombosis/Hemostasis, 2018, 24, 89S-93S. | 0.7 | 3 |
| 114 | Twenty children with non-Wilms renal tumors from a reference center in Central Anatolia, Turkey. Turkish Journal of Medical Sciences, 2019, 50, 18-24. | 0.4 | 3 |
| 115 | The number and activity of CD3+TCR VÎ±7.2+CD161+ cells are increased in children with acute rheumatic fever. International Journal of Cardiology, 2021, 333, 174-183. | 0.8 | 3 |
| 116 | Hematopoietic Stem Cell Transplant for Primary Immunodeficiency Diseases: A Single-Center Experience. Experimental and Clinical Transplantation, 2017, 15, 337-343. | 0.2 | 3 |
| 117 | A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia. Clinica Chimica Acta, 2022, 529, 61-66. | 0.5 | 3 |
| 118 | The Mutation of CD27 Deficiency Presented With Familial Hodgkin Lymphoma and a Review of the Literature. Journal of Pediatric Hematology/Oncology, 2022, 44, e833-e843. | 0.3 | 3 |
| 119 | Characterization of cord blood CD3<sup>+</sup>TCRVÎ±7.2<sup>+</sup>CD161<sup>high</sup> T and innate lymphoid cells in the pregnancies with gestational diabetes, morbidly adherent placenta, and pregnancy hypertension diseases. American Journal of Reproductive Immunology, 2022, 88, . | 1.2 | 3 |
| 120 | Coenzyme Q<sub>10</sub>Levels in Î²-Thalassemia and its Association with Ferritin Levels and Chelation Therapy. Hemoglobin, 2012, 36, 219-229. | 0.4 | 2 |
| 121 | The Importance of Nucleated Red Blood Cells in Patients with Beta Thalassemia Major and Comparison of Two Automated Systems with. Clinical Laboratory, 2015, 61, 1289-95. | 0.2 | 2 |
| 122 | Type 1 Plasminogen Deficiency With Pulmonary Involvement: Novel Treatment and Novel Mutation. Journal of Pediatric Hematology/Oncology, 2021, 43, e558-e560. | 0.3 | 2 |
| 123 | Hashimoto thyroiditis associated with ataxia telangiectasia. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 349-52. | 0.4 | 2 |
| 124 | Utility of the Aspergillus galactomannan antigen testing for neutropenic paediatric patients. Infezioni in Medicina, 2017, 25, 38-44. | 0.7 | 2 |
| 125 | Refractory cutaneous leishmaniasis in an adolescent: Initial manifestation of type 1 diabetes. Journal of Infection, 2006, 53, 290-291. | 1.7 | 1 |
| 126 | Reply from the authors of the article entitled â€œAtypical presentation of chronic granulomatous disease in an adolescent boy with frontal lobe located Aspergillus abscess mimicking intracranial tumorâ€• Child's Nervous System, 2010, 26, 735-735. | 0.6 | 1 |

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|---|---|---|---|
| 127 | Phenylketonuria With Acute Myeloblastic Leukemia in a 9-year-old Boy. Journal of Pediatric Hematology/Oncology, 2011, 33, e256-e257. | 0.3 | 1 |
| 128 | A mummy emerges from the grave: Scurvy confounding the clinical presentation of a child with Fanconi anemia. American Journal of Hematology, 2019, 94, 506-507. | 2.0 | 1 |
| 129 | Biliary Rhabdomyosarcoma in an Infant Male With Neurofibromatosis Type 1. Journal of Pediatric Hematology/Oncology, 2019, 41, e24-e26. | 0.3 | 1 |
| 130 | A Novel Intronic Mutation Reduces HAX1 Level and is Associated With Severe Congenital Neutropenia. Journal of Pediatric Hematology/Oncology, 2021, Publish Ahead of Print, . | 0.3 | 1 |
| 131 | A rare cause of vomiting in an adolescent: gastric burkitt`s lymphoma. Turkish Journal of Pediatrics, 2019, 61, 431. | 0.3 | 1 |
| 132 | Prognostic Factors and a New Prognostic Index Model for Children and Adolescents with Hodgkinâ€™s Lymphoma Who Underwent Autologous Hematopoietic Stem Cell Transplantation: A Multicenter Study of the Turkish Pediatric Bone Marrow Transplantation Study. Turkish Journal of Haematology, 2016, 33, 265-272. | 0.2 | 1 |
| 133 | Scurvy: A rare cause of arthritis in a child with neurologic disorder. European Journal of Rheumatology, 2018, 5, 283-284. | 1.3 | 1 |
| 134 | Central Nervous System Fungal Infections in Children with Leukemia, Risk Factors and Outcome: A Multicentric Study. Blood, 2018, 132, 5169-5169. | 0.6 | 1 |
| 135 | Assesment of Patients with Von Willebrand Disease with ISTH/BAT and PBQ Scores. Turkish Journal of Haematology, 2020, 37, 57-58. | 0.2 | 1 |
| 136 | The spectrum of underlying diseases in children with autoimmune hemolytic anemia. Journal of Health Sciences and Medicine, 2021, 4, 772-778. | 0.0 | 1 |
| 137 | Social exclusion and behavior problems in adolescents with cancer and healthy counterparts. Journal of Pediatric Nursing, 2022, 64, e95-e101. | 0.7 | 1 |
| 138 | The effect of methylenetetrahydrofolate reductase polymorphisms on the methotrexate toxicity in children with acute lymphoblastic leukemia. , 2022, , 9-13. | | 1 |
| 139 | The effect of methylenetetrahydrofolate reductase polymorphisms on the methotrexate toxicity in children with acute lymphoblastic leukemia. , 2022, , 9-13. | | 1 |
| 140 | The Real Incidence of Thyroid Carcinoma in Childhood. Laryngoscope, 2006, 116, 2095. | 1.1 | 0 |
| 141 | Importance of chemoprophylaxis and vaccines in the prevention of recurrent meningitis. Pediatrics International, 2008, 50, 416-416. | 0.2 | 0 |
| 142 | Laboratory, Clinical and Genetic Characteristics of Cases with Chronic Granulomatous Diseases: the Erciyes University Experience. Erciyes Tip Dergisi, 2012, 34, 121-126. | 0.1 | 0 |
| 143 | Ataksi telanjiektazi ve ikincil hastalÄ±klar. Turk Pediatri Arsivi, 2012, 47, 38-42. | 0.9 | 0 |
| 144 | Long-Term Patient-Customized Therapy for a Pathogenic EPO Mutation. Med, 2021, 2, 33-37.e1. | 2.2 | 0 |

| # | Article | IF | Citations |
|---|---|---|---|
| 145 | Inherited Biallelic Loss-Of-Function Mutations In CSF3R Define a Novel Type Of Severe Congenital Neutropenia With Full Myeloid Cell Maturation and Refractoriness To RhG-CSF. Blood, 2013, 122, 1025-1025. | 0.6 | 0 |
| 146 | Pansitopeni ile baÅŸvuran metilmalonik asidemi: Olgu sunumu. Medical Journal of Bakirkoy, 2018, , 138-41. | 0.0 | 0 |
| 147 | Next Generation Proteomics of Human Neutrophil Granulocytes in Monogenic Disease. Blood, 2018, 132, 15-15. | 0.6 | 0 |
| 148 | Burkitt Leukemia in a 5-Year-Old Girl with Williamsâ€“Beuren Syndrome: Review of the Literature. Indian Journal of Medical and Paediatric Oncology, 2019, 40, S114-S116. | 0.1 | 0 |
| 149 | Novel Immune Checkpoint Deficiency and Susceptibility to EBV-Associated Lymphoma. Blood, 2019, 134, 2326-2326. | 0.6 | 0 |
| 150 | Machine Learning Unveils Proteotypic Mimicry in Genetically Defined SCN Variants. Blood, 2019, 134, 3580-3580. | 0.6 | 0 |
| 151 | Impact of Intrauterine Growth Restriction Diseases on The Umbilical Cord Blood CD34+ Cell Counts. , 2020, , 56-61. | | 0 |
| 152 | Inflammatory Myofibroblastic Tumor of the Bronchus Mimicking Asthma. , 2020, , 103-104. | | 0 |
| 153 | Congenital afibrinogenemia in a 4-year-old girl complicated with acute lymphoblastic leukemia. Turkish Journal of Pediatrics, 2020, 62, 289. | 0.3 | 0 |
| 154 | THE EFFECT OF NIVOLUMAB IN PEDIATRIC MALIGNANT TUMORS: A SINGLE CENTER EXPERIENCE WITH EIGHT PATIENTS. Hematology, Transfusion and Cell Therapy, 2021, 43, S65. | 0.1 | 0 |
| 155 | Single-center experience of childhood Hodgkin lymphoma treated without radiotherapy. Journal of Health Sciences and Medicine, 2021, 4, 853-857. | 0.0 | 0 |
| 156 | Hemophagocytic lymphohistiocytosis: pouring gasoline on the cytokine storm. , 2022, , 30-34. | | 0 |