

Ekrem Unal

List of Publications by Year in descending order

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Version: 2024-02-01

156
papers

2,570
citations

236925

25
h-index

243625

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. <i>Journal of Allergy and Clinical Immunology</i> , 2015, 136, 402-412.	2.9	163
2	Hypomorphic homozygous mutations in phosphoglucomutase 3 (PGM3) impair immunity and increase serum IgE levels. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 1410-1419.e13.	2.9	160
3	Hematologically important mutations: Leukocyte adhesion deficiency (first update). <i>Blood Cells, Molecules, and Diseases</i> , 2012, 48, 53-61.	1.4	147
4	Genetic, immunological, and clinical features of patients with bacterial and fungal infections due to inherited IL-17RA deficiency. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E8277-E8285.	7.1	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. <i>Blood</i> , 2013, 121, 1345-1356.	1.4	122
6	Functional Selectivity in Cytokine Signaling Revealed Through a Pathogenic EPO Mutation. <i>Cell</i> , 2017, 168, 1053-1064.e15.	28.9	98
7	Inherited biallelic CSF3R mutations in severe congenital neutropenia. <i>Blood</i> , 2014, 123, 3811-3817.	1.4	79
8	Defective glycosylation and multisystem abnormalities characterize the primary immunodeficiency XMEN disease. <i>Journal of Clinical Investigation</i> , 2019, 130, 507-522.	8.2	74
9	Outcomes and Treatment Strategies for Autoimmunity and Hyperinflammation in Patients with RAG Deficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 1970-1985.e4.	3.8	64
10	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. <i>Blood</i> , 2020, 136, 2638-2655.	1.4	64
11	CD137 deficiency causes immune dysregulation with predisposition to lymphomagenesis. <i>Blood</i> , 2019, 134, 1510-1516.	1.4	52
12	Proteome Analysis of Human Neutrophil Granulocytes From Patients With Monogenic Disease Using Data-independent Acquisition. <i>Molecular and Cellular Proteomics</i> , 2019, 18, 760-772.	3.8	52
13	A case of XMEN syndrome presented with severe auto-immune disorders mimicking autoimmune lymphoproliferative disease. <i>Clinical Immunology</i> , 2015, 159, 58-62.	3.2	41
14	Childhood Stroke: Results of 130 Children From a Reference Center in Central Anatolia, Turkey. <i>Pediatric Neurology</i> , 2014, 50, 595-600.	2.1	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. <i>Child's Nervous System</i> , 2014, 30, 227-240.	1.1	37
16	Targeted high-throughput sequencing for genetic diagnostics of hemophagocytic lymphohistiocytosis. <i>Genome Medicine</i> , 2015, 7, 130.	8.2	37
17	Novel Mutations in RASGRP1 are Associated with Immunodeficiency, Immune Dysregulation, and EBV-Induced Lymphoma. <i>Journal of Clinical Immunology</i> , 2018, 38, 699-710.	3.8	37
18	Blood repellent superhydrophobic surfaces constructed from nanoparticle-free and biocompatible materials. <i>Colloids and Surfaces B: Biointerfaces</i> , 2021, 205, 111864.	5.0	35

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19	Genetic Deficiency and Biochemical Inhibition of ITK Affect Human Th17, Treg, and Innate Lymphoid Cells. <i>Journal of Clinical Immunology</i> , 2019, 39, 391-400.	3.8	34
20	Apparent diffusion coefficient in differentiation of pediatric posterior fossa tumors. <i>Japanese Journal of Radiology</i> , 2017, 35, 448-453.	2.4	32
21	Stem cell mobilization and collection from pediatric patients and healthy children. <i>Transfusion and Apheresis Science</i> , 2015, 53, 17-22.	1.0	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. <i>Journal of Clinical Immunology</i> , 2014, 34, 792-795.	3.8	30
23	Torticollis in children: an alert symptom not to be turned away. <i>Child's Nervous System</i> , 2015, 31, 1461-1470.	1.1	30
24	Late-type vitamin K deficiency bleeding: experience from 120 patients. <i>Child's Nervous System</i> , 2012, 28, 247-251.	1.1	28
25	Intracranial hemorrhage in children with congenital factor deficiencies. <i>Child's Nervous System</i> , 2011, 27, 1963-1966.	1.1	27
26	Atypical presentation of chronic granulomatous disease in an adolescent boy with frontal lobe located <i>Aspergillus</i> abscess mimicking intracranial tumor. <i>Child's Nervous System</i> , 2010, 26, 149-154.	1.1	26
27	CARMIL2 Deficiency Presenting as Very Early Onset Inflammatory Bowel Disease. <i>Inflammatory Bowel Diseases</i> , 2019, 25, 1788-1795.	1.9	26
28	Astroblastoma in a child. <i>Child's Nervous System</i> , 2008, 24, 165-168.	1.1	25
29	Hemophagocytic syndrome in a 4-month-old infant with biotinidase deficiency. <i>Pediatric Blood and Cancer</i> , 2012, 59, 191-193.	1.5	23
30	Human FCHO1 deficiency reveals role for clathrin-mediated endocytosis in development and function of T cells. <i>Nature Communications</i> , 2020, 11, 1031.	12.8	23
31	Autoimmune diseases detected in children with primary immunodeficiency diseases: results from a reference centre at middle anatolia. <i>Acta Microbiologica Et Immunologica Hungarica</i> , 2012, 59, 343-353.	0.8	22
32	Whole exome sequencing (WES) approach for diagnosing primary immunodeficiencies (PIDs) in a highly consanguineous community. <i>Clinical Immunology</i> , 2020, 214, 108376.	3.2	22
33	Web-based survey of resources for treatment and long-term follow-up for children with brain tumors in developing countries. <i>Child's Nervous System</i> , 2011, 27, 1957-1961.	1.1	21
34	Mesenchymal Hamartoma of the Liver Mimicking Hepatoblastoma. <i>Journal of Pediatric Hematology/Oncology</i> , 2008, 30, 458-460.	0.6	20
35	The efficacy of <i>Pelargonium sidoides</i> in the treatment of upper respiratory tract infections in children with transient hypogammaglobulinemia of infancy. <i>Phytomedicine</i> , 2012, 19, 958-961.	5.3	20
36	Expanded circulating hematopoietic stem/progenitor cells as novel cell source for the treatment of TCIRG1 osteopetrosis. <i>Haematologica</i> , 2020, 106, 74-86.	3.5	20

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37	Evolution and long-term outcomes of combined immunodeficiency due to CARMIL2 deficiency. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2022, 77, 1004-1019.	5.7	19
38	Diffusion weighted imaging in differentiating malignant and benign neuroblastic tumors. <i>Japanese Journal of Radiology</i> , 2016, 34, 620-624.	2.4	17
39	ILC3 deficiency and generalized ILC abnormalities in DOCK8-deficient patients. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2020, 75, 921-932.	5.7	17
40	Merkel Cell Carcinoma in a Child. <i>Journal of Pediatric Hematology/Oncology</i> , 2009, 31, 359-361.	0.6	16
41	Diagnostic and prognostic significance of glypican-5 and glypican-6 gene expression levels in gastric adenocarcinoma. <i>Molecular and Clinical Oncology</i> , 2015, 3, 584-590.	1.0	16
42	Malignant glioblastomatous transformation of a low-grade glioma in a child. <i>Child's Nervous System</i> , 2008, 24, 1385-1389.	1.1	15
43	CARDIAC FUNCTIONS EVALUATED WITH TISSUE DOPPLER IMAGING IN CHILDHOOD CANCERS TREATED WITH ANTHRACYCLINES. <i>Pediatric Hematology and Oncology</i> , 2010, 27, 13-23.	0.8	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. <i>Child's Nervous System</i> , 2014, 30, 1375-1382.	1.1	15
45	Brain Abscesses in Children. <i>Journal of Child Neurology</i> , 2015, 30, 458-467.	1.4	15
46	A case of congenital afibrinogenemia complicated with thromboembolic events that required repeated amputations. <i>Blood Coagulation and Fibrinolysis</i> , 2015, 26, 354-356.	1.0	14
47	The relationship between hematological parameters and prognosis of children with acute ischemic stroke. <i>Child's Nervous System</i> , 2018, 34, 655-661.	1.1	14
48	Cerebral venous sinus thrombosis in an adolescent with Ewing sarcoma. <i>Child's Nervous System</i> , 2008, 24, 983-986.	1.1	13
49	The Efficacy of Vitamin K2 and Calcitriol Combination on Thalassemic Osteopathy. <i>Journal of Pediatric Hematology/Oncology</i> , 2013, 35, 623-627.	0.6	13
50	Multiple Fungal Brain Abscesses in a Child with Acute Lymphoblastic Leukemia. <i>Mycopathologia</i> , 2012, 174, 505-509.	3.1	12
51	Gastric Signet Ring Carcinoma in a Patient With Ataxia-Telangiectasia. <i>Journal of Pediatric Hematology/Oncology</i> , 2013, 35, e341-e343.	0.6	12
52	Is there relation between COL4A1/A2 mutations and antenatally detected fetal intraventricular hemorrhage?. <i>Child's Nervous System</i> , 2014, 30, 419-424.	1.1	12
53	Acute Colchicine Intoxication Complicated With Extramedullary Hematopoiesis Due to Filgrastim in a Child. <i>Journal of Pediatric Hematology/Oncology</i> , 2014, 36, e460-e462.	0.6	12
54	Genetic Analysis of a Cohort of 275 Patients with Hyper-IgE Syndromes and/or Chronic Mucocutaneous Candidiasis. <i>Journal of Clinical Immunology</i> , 2021, 41, 1804-1838.	3.8	12

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55	Propranolol treatment for chylothorax due to diffuse lymphangiomatosis. <i>Pediatric Blood and Cancer</i> , 2019, 66, e27592.	1.5	11
56	Juvenile Myelomonocytic Leukemia in Turkey: A Retrospective Analysis of Sixty-five Patients. <i>Türkmen Tıp Dergisi</i> . <i>Turkish Journal of Haematology</i> , 2018, 35, 27-34.	0.5	11
57	Neuroblastoma Arising From an Unresected Sacrococcygeal Teratoma in a Child. <i>Journal of Pediatric Hematology/Oncology</i> , 2010, 32, 233-235.	0.6	10
58	Chronic granulomatous disease with markedly elevated IgE levels mimicking hyperimmunoglobulin E syndrome. <i>Acta Microbiologica Et Immunologica Hungarica</i> , 2013, 60, 155-162.	0.8	10
59	X-linked severe combined immunodeficiency due to a novel mutation complicated with hemophagocytic lymphohistiocytosis and presented with invagination: A case report. <i>European Journal of Microbiology and Immunology</i> , 2014, 4, 174-176.	2.8	10
60	Cerebral Sinus Venous Thrombosis and Prothrombotic Risk Factors in Children: A Single-Center Experience From Turkey. <i>Journal of Pediatric Hematology/Oncology</i> , 2018, 40, e369-e372.	0.6	10
61	Quartz Crystal Microbalance Measurements of CD19 Antibody Immobilization on Gold Surface and Capturing B Lymphoblast Cells: Effect of Surface Functionalization. <i>Electroanalysis</i> , 2018, 30, 834-841.	2.9	10
62	A Rare Case of Activated Phosphoinositide 3-Kinase Delta Syndrome (APDS) Presenting With Hemophagocytosis Complicated With Hodgkin Lymphoma. <i>Journal of Pediatric Hematology/Oncology</i> , 2020, 42, 156-159.	0.6	10
63	Effectiveness of Ankaferd BloodStopper in Prophylaxis and Treatment of Oral Mucositis in Childhood Cancers Evaluated with Plasma Citrulline Levels. <i>Turkish Journal of Haematology</i> , 2018, 35, 85-86.	0.5	10
64	Fatal valproate overdose in a newborn baby. <i>Human and Experimental Toxicology</i> , 2007, 26, 453-456.	2.2	9
65	Thyroid Medullary Carcinoma in a Teenager With Cowden Syndrome. <i>Laryngoscope</i> , 2007, 117, 1180-1182.	2.0	9
66	Pituitary-adrenal axis suppression due to topical steroid administration in an infant. <i>Pediatrics International</i> , 2007, 49, 242-244.	0.5	9
67	An uncommon extrapulmonary sequestration located in the upper posterior mediastinum associated with the azygos lobe in a child. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2007, 133, 1110-1111.	0.8	9
68	Release of N-terminal pro-brain natriuretic peptide in children with acute rheumatic carditis. <i>Cardiology in the Young</i> , 2010, 20, 297-301.	0.8	9
69	Cerebellar hemangioblastoma associated with diffuse neonatal hemangiomatosis in an infant. <i>Child's Nervous System</i> , 2012, 28, 1801-1805.	1.1	9
70	Inflammatory Myofibroblastic Tumor of the Kidney and Bilateral Lung Nodules in a Child Mimicking Wilms Tumor With Lung Metastases. <i>Journal of Pediatric Hematology/Oncology</i> , 2015, 37, e390-e393.	0.6	9
71	Neuroblastoma in a Child With Wolf-Hirschhorn Syndrome. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, e224-e226.	0.6	9
72	Microfluidic Chip based direct triple antibody immunoassay for monitoring patient comparative response to leukemia treatment. <i>Biomedical Microdevices</i> , 2020, 22, 48.	2.8	9

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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.	0.5	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.	4.8	9
75	Mycophenolate mofetil-induced pseudotumor cerebri in a boy with autoimmune lymphoproliferative disease. Child's Nervous System, 2011, 27, 853-855.	1.1	8
76	The importance of MTHFR polymorphisms in pediatric cerebral stroke. Child's Nervous System, 2012, 28, 13-13.	1.1	8
77	Fetal intracranial hemorrhage related to maternal autoimmune thrombocytopenic purpura. Child's Nervous System, 2014, 30, 2147-2150.	1.1	8
78	Hepatosplenic Fungal Infections in Children With Leukemia—Risk Factors and Outcome: A Multicentric Study. Journal of Pediatric Hematology/Oncology, 2019, 41, 256-260.	0.6	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.	0.6	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.	0.6	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.	0.6	6
82	Discovery of the First Pathogenic Human EPO Mutation Provides Mechanistic Insight into Cytokine Signaling. Blood, 2016, 128, 331-331.	1.4	6
83	Hashimoto thyroiditis associated with ataxia telangiectasia. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, .	0.9	5
84	Orbital myositis associated with focal active colitis in a teenage girl. Child's Nervous System, 2012, 28, 641-643.	1.1	5
85	Infantile tremor syndrome associated with cobalamin therapy: A case report. Clinical Neurology and Neurosurgery, 2013, 115, 1903-1905.	1.4	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.	1.0	5
87	Pediatric central nervous system tumors in the first 3 years 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.	1.1	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.	2.1	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.	0.6	5
90	Hepatitis-associated aplastic anemia in pediatric patients: single center experience. Transfusion and Apheresis Science, 2020, 59, 102900.	1.0	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. <i>Separation Science and Technology</i> , 2021, 56, 2659-2666.	2.5	5
92	Effect of vitamin K2 and vitamin D3 on bone mineral density in children with acute lymphoblastic leukemia: a prospective cohort study. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2021, 34, 441-447.	0.9	5
93	Vena Cava Superior Syndrome in Children With Mediastinal Tumors: Single Center Experience. <i>Ästanbul Kuzey Klinikleri</i> , 2019, 7, 255-259.	0.3	5
94	Pulmonary arterial pressure in infants with laryngomalacia. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2006, 70, 2067-2071.	1.0	4
95	Hypothermia in a Child With Hodgkin Disease. <i>Journal of Pediatric Hematology/Oncology</i> , 2009, 31, 136-138.	0.6	4
96	A Case of Familial Hemophagocytic Lymphohistiocytosis Type 4 With Involvement of the Central Nervous System Complicated With Infarct. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, e321-e324.	0.6	4
97	Capturing B type acute lymphoblastic leukemia cells using two types of antibodies. <i>Biotechnology Progress</i> , 2019, 35, e2737.	2.6	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. <i>Leukemia and Lymphoma</i> , 2020, 61, 1465-1474.	1.3	4
99	A novel missense mutation outside the <sc>DNAJ</sc> domain of <sc><i>DNAJC21</i></sc> is associated with <sc>Shwachmanâ€“Diamond</sc> syndrome. <i>British Journal of Haematology</i> , 2022, 197, .	2.5	4
100	Letter to the Editor. <i>Inhalation Toxicology</i> , 2007, 19, 587-587.	1.6	3
101	KLUVER-BUCY SYNDROME IN A BOY WITH NON-HODGKIN LYMPHOMA. <i>Pediatric Hematology and Oncology</i> , 2007, 24, 149-152.	0.8	3
102	Pilocytic astrocytoma developing at the site of a previously treated medulloblastoma in a child. <i>Child's Nervous System</i> , 2008, 24, 289-292.	1.1	3
103	Accidental intramuscular overdose administration of vincristine. <i>Drug and Chemical Toxicology</i> , 2012, 35, 232-234.	2.3	3
104	Atrial fibrillation as an uncommon presentation in a large pleomorphic xanthoastrocytoma. <i>Child's Nervous System</i> , 2012, 28, 475-479.	1.1	3
105	Myeloperoxidase Deficiency: The Secret Under the Flag of Unstained Cell. <i>Turkish Journal of Haematology</i> , 2013, 30, 232-233.	0.2	3
106	Neurological complication of non Hodgkin lymphoma in childhood: experience from a single center in Turkey. <i>Child's Nervous System</i> , 2014, 30, 639-645.	1.1	3
107	Cranial metastatic alveolar rhabdomyosarcoma mimicking hematological malignancy in an adolescent boy. <i>Child's Nervous System</i> , 2014, 30, 1737-1741.	1.1	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. <i>Acta Microbiologica Et Immunologica Hungarica</i> , 2015, 62, 267-274.	0.8	3

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127	Phenylketonuria With Acute Myeloblastic Leukemia in a 9-year-old Boy. <i>Journal of Pediatric Hematology/Oncology</i> , 2011, 33, e256-e257.	0.6	1
128	A mummy emerges from the grave: Scurvy confounding the clinical presentation of a child with Fanconi anemia. <i>American Journal of Hematology</i> , 2019, 94, 506-507.	4.1	1
129	Biliary Rhabdomyosarcoma in an Infant Male With Neurofibromatosis Type 1. <i>Journal of Pediatric Hematology/Oncology</i> , 2019, 41, e24-e26.	0.6	1
130	A Novel Intronic Mutation Reduces HAX1 Level and is Associated With Severe Congenital Neutropenia. <i>Journal of Pediatric Hematology/Oncology</i> , 2021, Publish Ahead of Print, .	0.6	1
131	A rare cause of vomiting in an adolescent: gastric burkitt's lymphoma. <i>Turkish Journal of Pediatrics</i> , 2019, 61, 431.	0.6	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. <i>Turkish Journal of Haematology</i> , 2016, 33, 265-272.	0.5	1
133	Scurvy: A rare cause of arthritis in a child with neurologic disorder. <i>European Journal of Rheumatology</i> , 2018, 5, 283-284.	0.6	1
134	Central Nervous System Fungal Infections in Children with Leukemia, Risk Factors and Outcome: A Multicentric Study. <i>Blood</i> , 2018, 132, 5169-5169.	1.4	1
135	Assesment of Patients with Von Willebrand Disease with ISTH/BAT and PBQ Scores. <i>Turkish Journal of Haematology</i> , 2020, 37, 57-58.	0.5	1
136	The spectrum of underlying diseases in children with autoimmune hemolytic anemia. <i>Journal of Health Sciences and Medicine</i> , 2021, 4, 772-778.	0.1	1
137	Social exclusion and behavior problems in adolescents with cancer and healthy counterparts. <i>Journal of Pediatric Nursing</i> , 2022, 64, e95-e101.	1.5	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. <i>Laryngoscope</i> , 2006, 116, 2095.	2.0	0
141	Importance of chemoprophylaxis and vaccines in the prevention of recurrent meningitis. <i>Pediatrics International</i> , 2008, 50, 416-416.	0.5	0
142	Laboratory, Clinical and Genetic Characteristics of Cases with Chronic Granulomatous Diseases: the Erciyes University Experience. <i>Erciyes Tip Dergisi</i> , 2012, 34, 121-126.	0.1	0
143	Ataksi telanjiektazi ve ikincil hastalıklar. <i>Turk Peditri Arsivi</i> , 2012, 47, 38-42.	0.9	0
144	Long-Term Patient-Customized Therapy for a Pathogenic EPO Mutation. <i>Med</i> , 2021, 2, 33-37.e1.	4.4	0

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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.	1.4	0
146	Pansitopeni ile baÅyvuran metilmalonik asidemi: Olgu sunumu. Medical Journal of Bakirkoy, 2018, , 138-41.	0.1	0
147	Next Generation Proteomics of Human Neutrophil Granulocytes in Monogenic Disease. Blood, 2018, 132, 15-15.	1.4	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.	0.2	0
149	Novel Immune Checkpoint Deficiency and Susceptibility to EBV-Associated Lymphoma. Blood, 2019, 134, 2326-2326.	1.4	0
150	Machine Learning Unveils Proteotypic Mimicry in Genetically Defined SCN Variants. Blood, 2019, 134, 3580-3580.	1.4	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.	0.6	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.	0.2	0
155	Single-center experience of childhood Hodgkin lymphoma treated without radiotherapy. Journal of Health Sciences and Medicine, 2021, 4, 853-857.	0.1	0
156	Hemophagocytic lymphohistiocytosis: pouring gasoline on the cytokine storm. , 2022, , 30-34.		0