

Ekrem Unal

List of Publications by Year in descending order

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Version: 2024-02-01

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	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.	2.7	19
2	Social exclusion and behavior problems in adolescents with cancer and healthy counterparts. <i>Journal of Pediatric Nursing</i> , 2022, 64, e95-e101.	0.7	1
3	A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia. <i>Clinica Chimica Acta</i> , 2022, 529, 61-66.	0.5	3
4	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, .	1.2	4
5	Preclinical Studies on Convalescent Human Immune Plasma-Derived Exosome: Omics and Antiviral Properties to SARS-CoV-2. <i>Frontiers in Immunology</i> , 2022, 13, 824378.	2.2	9
6	The Mutation of CD27 Deficiency Presented With Familial Hodgkin Lymphoma and a Review of the Literature. <i>Journal of Pediatric Hematology/Oncology</i> , 2022, 44, e833-e843.	0.3	3
7	Characterization of cord blood CD3⁺TCR\hat{V}7.2⁺CD161^{high} T and innate lymphoid cells in the pregnancies with gestational diabetes, morbidly adherent placenta, and pregnancy hypertension diseases. <i>American Journal of Reproductive Immunology</i> , 2022, 88, .	1.2	3
8	Hemophagocytic lymphohistiocytosis: pouring gasoline on the cytokine storm. , 2022, , 30-34.		0
9	The effect of methylenetetrahydrofolate reductase polymorphisms on the methotrexate toxicity in children with acute lymphoblastic leukemia. , 2022, , 9-13.		1
10	The effect of methylenetetrahydrofolate reductase polymorphisms on the methotrexate toxicity in children with acute lymphoblastic leukemia. , 2022, , 9-13.		1
11	Long-Term Patient-Customized Therapy for a Pathogenic EPO Mutation. <i>Med</i> , 2021, 2, 33-37.e1.	2.2	0
12	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.	1.3	5
13	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, .	0.3	1
14	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.	0.4	5
15	The number and activity of CD3+TCR \hat{V}7.2+CD161+ cells are increased in children with acute rheumatic fever. <i>International Journal of Cardiology</i> , 2021, 333, 174-183.	0.8	3
16	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.	2.0	12
17	Blood repellent superhydrophobic surfaces constructed from nanoparticle-free and biocompatible materials. <i>Colloids and Surfaces B: Biointerfaces</i> , 2021, 205, 111864.	2.5	35
18	Refractory and Fatal Presentation of Severe Autoimmune Hemolytic Anemia in a Child With the DNASE1L3 Mutation Complicated With an Additional DOCK8 Variant. <i>Journal of Pediatric Hematology/Oncology</i> , 2021, 43, e452-e456.	0.3	7

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19	Common Variable Immunodeficiency, Autoimmune Hemolytic Anemia, and Pancytopenia Associated With a Defect in IKAROS. <i>Journal of Pediatric Hematology/Oncology</i> , 2021, 43, e351-e357.	0.3	6
20	Type 1 Plasminogen Deficiency With Pulmonary Involvement: Novel Treatment and Novel Mutation. <i>Journal of Pediatric Hematology/Oncology</i> , 2021, 43, e558-e560.	0.3	2
21	THE EFFECT OF NIVOLUMAB IN PEDIATRIC MALIGNANT TUMORS: A SINGLE CENTER EXPERIENCE WITH EIGHT PATIENTS. <i>Hematology, Transfusion and Cell Therapy</i> , 2021, 43, S65.	0.1	0
22	The spectrum of underlying diseases in children with autoimmune hemolytic anemia. <i>Journal of Health Sciences and Medicine</i> , 2021, 4, 772-778.	0.0	1
23	Single-center experience of childhood Hodgkin lymphoma treated without radiotherapy. <i>Journal of Health Sciences and Medicine</i> , 2021, 4, 853-857.	0.0	0
24	ILC3 deficiency and generalized ILC abnormalities in DOCK8-deficient patients. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2020, 75, 921-932.	2.7	17
25	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.	0.3	10
26	Different Clinical Presentation of 3 Children With Familial Hemophagocytic Lymphohistiocytosis With 2 Novel Mutations. <i>Journal of Pediatric Hematology/Oncology</i> , 2020, 42, e627-e629.	0.3	5
27	Microfluidic Chip based direct triple antibody immunoassay for monitoring patient comparative response to leukemia treatment. <i>Biomedical Microdevices</i> , 2020, 22, 48.	1.4	9
28	Hepatitis-associated aplastic anemia in pediatric patients: single center experience. <i>Transfusion and Apheresis Science</i> , 2020, 59, 102900.	0.5	5
29	Expanded circulating hematopoietic stem/progenitor cells as novel cell source for the treatment of TCIRG1 osteopetrosis. <i>Haematologica</i> , 2020, 106, 74-86.	1.7	20
30	Whole exome sequencing (WES) approach for diagnosing primary immunodeficiencies (PIDs) in a highly consanguineous community. <i>Clinical Immunology</i> , 2020, 214, 108376.	1.4	22
31	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. <i>Blood</i> , 2020, 136, 2638-2655.	0.6	64
32	Human FCHO1 deficiency reveals role for clathrin-mediated endocytosis in development and function of T cells. <i>Nature Communications</i> , 2020, 11, 1031.	5.8	23
33	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.	0.6	4
34	Assesment of Patients with Von Willebrand Disease with ISTH/BAT and PBQ Scores. <i>Turkish Journal of Haematology</i> , 2020, 37, 57-58.	0.2	1
35	Impact of Intrauterine Growth Restriction Diseases on The Umbilical Cord Blood CD34+ Cell Counts. , 2020, , 56-61.		0
36	Inflammatory Myofibroblastic Tumor of the Bronchus Mimicking Asthma. , 2020, , 103-104.		0

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37	Congenital afibrinogenemia in a 4-year-old girl complicated with acute lymphoblastic leukemia. Turkish Journal of Pediatrics, 2020, 62, 289.	0.3	0
38	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
39	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
40	CARMIL2 Deficiency Presenting as Very Early Onset Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2019, 25, 1788-1795.	0.9	26
41	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
42	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
43	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
44	CD137 deficiency causes immune dysregulation with predisposition to lymphomagenesis. Blood, 2019, 134, 1510-1516.	0.6	52
45	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
46	Propranolol treatment for chylothorax due to diffuse lymphangiomatosis. Pediatric Blood and Cancer, 2019, 66, e27592.	0.8	11
47	Capturing B type acute lymphoblastic leukemia cells using two types of antibodies. Biotechnology Progress, 2019, 35, e2737.	1.3	4
48	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
49	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
50	Biliary Rhabdomyosarcoma in an Infant Male With Neurofibromatosis Type 1. Journal of Pediatric Hematology/Oncology, 2019, 41, e24-e26.	0.3	1
51	Defective glycosylation and multisystem abnormalities characterize the primary immunodeficiency XMEN disease. Journal of Clinical Investigation, 2019, 130, 507-522.	3.9	74
52	Vena Cava Superior Syndrome in Children With Mediastinal Tumors: Single Center Experience. İstanbul Kuzey Klinikleri, 2019, 7, 255-259.	0.1	5
53	A rare cause of vomiting in an adolescent: gastric burkitt's lymphoma. Turkish Journal of Pediatrics, 2019, 61, 431.	0.3	1
54	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

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55	Novel Immune Checkpoint Deficiency and Susceptibility to EBV-Associated Lymphoma. <i>Blood</i> , 2019, 134, 2326-2326.	0.6	0
56	Machine Learning Unveils Proteotypic Mimicry in Genetically Defined SCN Variants. <i>Blood</i> , 2019, 134, 3580-3580.	0.6	0
57	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.	0.3	10
58	Quartzâ€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.	1.5	10
59	The relationship between hematological parameters and prognosis of children with acute ischemic stroke. <i>Child's Nervous System</i> , 2018, 34, 655-661.	0.6	14
60	Novel Mutations in RASGRP1 are Associated with Immunodeficiency, Immune Dysregulation, and EBV-Induced Lymphoma. <i>Journal of Clinical Immunology</i> , 2018, 38, 699-710.	2.0	37
61	Genetic Polymorphism of VKORC1-1639 in Children With Intracranial Hemorrhage Due to Vitamin K Deficiency. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2018, 24, 89S-93S.	0.7	3
62	Juvenile Myelomonocytic Leukemia in Turkey: A Retrospective Analysis of Sixty-five Patientsâ€zlem TÃ¼fekÅŸi. <i>Turkish Journal of Haematology</i> , 2018, 35, 27-34.	0.2	11
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.	0.2	10
64	Scurvy: A rare cause of arthritis in a child with neurologic disorder. <i>European Journal of Rheumatology</i> , 2018, 5, 283-284.	1.3	1
65	Pansitopeni ile baÅŸvuran metilmalonik asidemi: Olgu sunumu. <i>Medical Journal of Bakirkoy</i> , 2018, , 138-41.	0.0	0
66	Next Generation Proteomics of Human Neutrophil Granulocytes in Monogenic Disease. <i>Blood</i> , 2018, 132, 15-15.	0.6	0
67	Central Nervous System Fungal Infections in Children with Leukemia, Risk Factors and Outcome: A Multicentric Study. <i>Blood</i> , 2018, 132, 5169-5169.	0.6	1
68	Functional Selectivity in Cytokine Signaling Revealed Through a Pathogenic EPO Mutation. <i>Cell</i> , 2017, 168, 1053-1064.e15.	13.5	98
69	A Newborn With Familial Hemophagocytic Lymphohistiocytosis Complicated With Transfusion Associated Graft Versus Host Disease. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, e309-e311.	0.3	6
70	Apparent diffusion coefficient in differentiation of pediatric posterior fossa tumors. <i>Japanese Journal of Radiology</i> , 2017, 35, 448-453.	1.0	32
71	Pediatric central nervous system tumors in the first 3Ã¥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. <i>Child's Nervous System</i> , 2017, 33, 233-238.	0.6	5
72	Neuroblastoma in a Child With Wolf-Hirschhorn Syndrome. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, e224-e226.	0.3	9

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73	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.	0.3	4
74	Evaluation of childhood pancreas solid pseudopapillary tumors. <i>Ä°stanbul Kuzey Klinikleri</i> , 2017, 5, 207-210.	0.1	3
75	Hematopoietic Stem Cell Transplant for Primary Immunodeficiency Diseases: A Single-Center Experience. <i>Experimental and Clinical Transplantation</i> , 2017, 15, 337-343.	0.2	3
76	Utility of the Aspergillus galactomannan antigen testing for neutropenic paediatric patients. <i>Infezioni in Medicina</i> , 2017, 25, 38-44.	0.7	2
77	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.	3.3	137
78	Diffusion weighted imaging in differentiating malignant and benign neuroblastic tumors. <i>Japanese Journal of Radiology</i> , 2016, 34, 620-624.	1.0	17
79	Fibromuscular Dysplasia Complicated With Cerebral Stroke in a Child With Congenital Dyserythropoietic Anemia Type II. <i>Journal of Pediatric Hematology/Oncology</i> , 2016, 38, e333-e335.	0.3	3
80	Familial moyamoya disease in two Turkish siblings with same polymorphism in RNF213 gene but different clinical features. <i>Child's Nervous System</i> , 2016, 32, 569-573.	0.6	3
81	Prognostic Factors and a New Prognostic Index Model for Children and Adolescents with Hodgkin'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.	0.2	1
82	Discovery of the First Pathogenic Human EPO Mutation Provides Mechanistic Insight into Cytokine Signaling. <i>Blood</i> , 2016, 128, 331-331.	0.6	6
83	Targeted high-throughput sequencing for genetic diagnostics of hemophagocytic lymphohistiocytosis. <i>Genome Medicine</i> , 2015, 7, 130.	3.6	37
84	Autosomal recessive hyper IgM syndrome associated with activation-induced cytidine deaminase gene in three Turkish siblings presented with tuberculosis lymphadenitis – Case report. <i>Acta Microbiologica Et Immunologica Hungarica</i> , 2015, 62, 267-274.	0.4	3
85	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.	0.4	16
86	A case of congenital afibrinogenemia complicated with thromboembolic events that required repeated amputations. <i>Blood Coagulation and Fibrinolysis</i> , 2015, 26, 354-356.	0.5	14
87	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.	0.3	9
88	Outcome of autologous hematopoietic stem cell transplantation in children and adolescents with relapsed or refractory Hodgkin's lymphoma. <i>Pediatric Transplantation</i> , 2015, 19, 745-752.	0.5	5
89	Malignancies in Primary Immunodeficiencies: A Single Center Experience. <i>Pediatric, Allergy, Immunology, and Pulmonology</i> , 2015, 28, 47-54.	0.3	3
90	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.	1.5	163

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91	A case of XMEN syndrome presented with severe auto-immune disorders mimicking autoimmune lymphoproliferative disease. <i>Clinical Immunology</i> , 2015, 159, 58-62.	1.4	41
92	Torticollis in children: an alert symptom not to be turned away. <i>Child's Nervous System</i> , 2015, 31, 1461-1470.	0.6	30
93	Stem cell mobilization and collection from pediatric patients and healthy children. <i>Transfusion and Apheresis Science</i> , 2015, 53, 17-22.	0.5	31
94	Brain Abscesses in Children. <i>Journal of Child Neurology</i> , 2015, 30, 458-467.	0.7	15
95	The Importance of Nucleated Red Blood Cells in Patients with Beta Thalassemia Major and Comparison of Two Automated Systems with. <i>Clinical Laboratory</i> , 2015, 61, 1289-95.	0.2	2
96	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.	2.0	30
97	Inherited biallelic CSF3R mutations in severe congenital neutropenia. <i>Blood</i> , 2014, 123, 3811-3817.	0.6	79
98	Childhood Stroke: Results of 130 Children From a Reference Center in Central Anatolia, Turkey. <i>Pediatric Neurology</i> , 2014, 50, 595-600.	1.0	38
99	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.	0.6	37
100	Is there relation between COL4A1/A2 mutations and antenatally detected fetal intraventricular hemorrhage?. <i>Child's Nervous System</i> , 2014, 30, 419-424.	0.6	12
101	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.	0.6	3
102	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.	0.6	15
103	Cranial metastatic alveolar rhabdomyosarcoma mimicking hematological malignancy in an adolescent boy. <i>Child's Nervous System</i> , 2014, 30, 1737-1741.	0.6	3
104	Fetal intracranial hemorrhage related to maternal autoimmune thrombocytopenic purpura. <i>Child's Nervous System</i> , 2014, 30, 2147-2150.	0.6	8
105	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.	1.5	160
106	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.	1.5	10
107	Acute Colchicine Intoxication Complicated With Extramedullary Hematopoiesis Due to Filgrastim in a Child. <i>Journal of Pediatric Hematology/Oncology</i> , 2014, 36, e460-e462.	0.3	12
108	Infantile tremor syndrome associated with cobalamin therapy: A case report. <i>Clinical Neurology and Neurosurgery</i> , 2013, 115, 1903-1905.	0.6	5

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109	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.	0.6	122
110	Myeloperoxidase Deficiency: The Secret Under the Flag of Unstained Cell. <i>Turkish Journal of Haematology</i> , 2013, 30, 232-233.	0.2	3
111	The Efficacy of Vitamin K2 and Calcitriol Combination on Thalassemic Osteopathy. <i>Journal of Pediatric Hematology/Oncology</i> , 2013, 35, 623-627.	0.3	13
112	Gastric Signet Ring Carcinoma in a Patient With Ataxia-Telangiectasia. <i>Journal of Pediatric Hematology/Oncology</i> , 2013, 35, e341-e343.	0.3	12
113	Chronic granulomatous disease with markedly elevated IgE levels mimicking hyperimmunoglobulin E syndrome. <i>Acta Microbiologica Et Immunologica Hungarica</i> , 2013, 60, 155-162.	0.4	10
114	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. <i>Blood</i> , 2013, 122, 1025-1025.	0.6	0
115	Laboratory, Clinical and Genetic Characteristics of Cases with Chronic Granulomatous Diseases: the Erciyes University Experience. <i>Erciyes Tıp Dergisi</i> , 2012, 34, 121-126.	0.1	0
116	Ataksi telanjiektazi ve ikincil hastalıklar. <i>Türk Pediatri Arsivi</i> , 2012, 47, 38-42.	0.9	0
117	Coenzyme Q ₁₀ Levels in β^2 -Thalassemia and its Association with Ferritin Levels and Chelation Therapy. <i>Hemoglobin</i> , 2012, 36, 219-229.	0.4	2
118	Hashimoto thyroiditis associated with ataxia telangiectasia. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2012, 25, .	0.4	5
119	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.	0.4	22
120	Cerebellar hemangioblastoma associated with diffuse neonatal hemangiomatosis in an infant. <i>Child's Nervous System</i> , 2012, 28, 1801-1805.	0.6	9
121	Multiple Fungal Brain Abscesses in a Child with Acute Lymphoblastic Leukemia. <i>Mycopathologia</i> , 2012, 174, 505-509.	1.3	12
122	The efficacy of Pelargonium sidoides in the treatment of upper respiratory tract infections in children with transient hypogammaglobulinemia of infancy. <i>Phytomedicine</i> , 2012, 19, 958-961.	2.3	20
123	Accidental intramuscular overdose administration of vincristine. <i>Drug and Chemical Toxicology</i> , 2012, 35, 232-234.	1.2	3
124	Hematologically important mutations: Leukocyte adhesion deficiency (first update). <i>Blood Cells, Molecules, and Diseases</i> , 2012, 48, 53-61.	0.6	147
125	Hemophagocytic syndrome in a 4-month-old infant with biotinidase deficiency. <i>Pediatric Blood and Cancer</i> , 2012, 59, 191-193.	0.8	23
126	Orbital myositis associated with focal active colitis in a teenage girl. <i>Child's Nervous System</i> , 2012, 28, 641-643.	0.6	5

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127	Late-type vitamin K deficiency bleeding: experience from 120 patients. <i>Child's Nervous System</i> , 2012, 28, 247-251.	0.6	28
128	The importance of MTHFR polymorphisms in pediatric cerebral stroke. <i>Child's Nervous System</i> , 2012, 28, 13-13.	0.6	8
129	Atrial fibrillation as an uncommon presentation in a large pleomorphic xanthoastrocytoma. <i>Child's Nervous System</i> , 2012, 28, 475-479.	0.6	3
130	Hashimoto thyroiditis associated with ataxia telangiectasia. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2012, 25, 349-52.	0.4	2
131	Phenylketonuria With Acute Myeloblastic Leukemia in a 9-year-old Boy. <i>Journal of Pediatric Hematology/Oncology</i> , 2011, 33, e256-e257.	0.3	1
132	Mycophenolate mofetil-induced pseudotumor cerebri in a boy with autoimmune lymphoproliferative disease. <i>Child's Nervous System</i> , 2011, 27, 853-855.	0.6	8
133	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.	0.6	21
134	Intracranial hemorrhage in children with congenital factor deficiencies. <i>Child's Nervous System</i> , 2011, 27, 1963-1966.	0.6	27
135	Neuroblastoma Arising From an Unresected Sacrococcygeal Teratoma in a Child. <i>Journal of Pediatric Hematology/Oncology</i> , 2010, 32, 233-235.	0.3	10
136	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.	0.6	26
137	Reply from the authors of the article entitled "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, 735-735.	0.6	1
138	Release of N-terminal pro-brain natriuretic peptide in children with acute rheumatic carditis. <i>Cardiology in the Young</i> , 2010, 20, 297-301.	0.4	9
139	CARDIAC FUNCTIONS EVALUATED WITH TISSUE DOPPLER IMAGING IN CHILDHOOD CANCERS TREATED WITH ANTHRACYCLINES. <i>Pediatric Hematology and Oncology</i> , 2010, 27, 13-23.	0.3	15
140	Hypothermia in a Child With Hodgkin Disease. <i>Journal of Pediatric Hematology/Oncology</i> , 2009, 31, 136-138.	0.3	4
141	Merkel Cell Carcinoma in a Child. <i>Journal of Pediatric Hematology/Oncology</i> , 2009, 31, 359-361.	0.3	16
142	Astroblastoma in a child. <i>Child's Nervous System</i> , 2008, 24, 165-168.	0.6	25
143	Pilocytic astrocytoma developing at the site of a previously treated medulloblastoma in a child. <i>Child's Nervous System</i> , 2008, 24, 289-292.	0.6	3
144	Cerebral venous sinus thrombosis in an adolescent with Ewing sarcoma. <i>Child's Nervous System</i> , 2008, 24, 983-986.	0.6	13

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145	Malignant glioblastomatous transformation of a low-grade glioma in a child. <i>Child's Nervous System</i> , 2008, 24, 1385-1389.	0.6	15
146	Importance of chemoprophylaxis and vaccines in the prevention of recurrent meningitis. <i>Pediatrics International</i> , 2008, 50, 416-416.	0.2	0
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