Ekrem Unal

List of Publications by Year
in descending order

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	Evolution and longâ€term outcomes of combined immunodeficiency due to CARMIL2 deficiency. Allergy: European Journal of Allergy and Clinical Immunology, 2022, 77, 1004-1019.	5.7	19
2	Social exclusion and behavior problems in adolescents with cancer and healthy counterparts. Journal of Pediatric Nursing, 2022, 64, e95-e101.	1.5	1
3	A teenager boy with a novel variant of Sitosterolemia presented with pancytopenia. Clinica Chimica Acta, 2022, 529, 61-66.	1.1	3
4	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, .	2.5	4
5	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
6	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.	0.6	3
7	Characterization of cord blood CD3⁺TCRVÎ±7.2⁺CD161^{high} 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
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. Med, 2021, 2, 33-37.e1.	4.4	0
12	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.	2.5	5
13	A Novel Intronic Mutation Reduces HAX1 Level and is Associated With Severe Congenital Neutropenia. Journal of Pediatric Hematology/Oncology, 2021, Publish Ahead of Print, .	0.6	1
14	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.	0.9	5
15	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.	1.7	3
16	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.	3.8	12
17	Blood repellent superhydrophobic surfaces constructed from nanoparticle-free and biocompatible materials. Colloids and Surfaces B: Biointerfaces, 2021, 205, 111864.	5.0	35
18	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

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19	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
20	Type 1 Plasminogen Deficiency With Pulmonary Involvement: Novel Treatment and Novel Mutation. Journal of Pediatric Hematology/Oncology, 2021, 43, e558-e560.	0.6	2
21	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
22	The spectrum of underlying diseases in children with autoimmune hemolytic anemia. Journal of Health Sciences and Medicine, 2021, 4, 772-778.	0.1	1
23	Single-center experience of childhood Hodgkin lymphoma treated without radiotherapy. Journal of Health Sciences and Medicine, 2021, 4, 853-857.	0.1	0
24	ILC3 deficiency and generalized ILC abnormalities in DOCK8â€deficient patients. Allergy: European Journal of Allergy and Clinical Immunology, 2020, 75, 921-932.	5.7	17
25	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.	0.6	10
26	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
27	Microfluidic Chip based direct triple antibody immunoassay for monitoring patient comparative response to leukemia treatment. Biomedical Microdevices, 2020, 22, 48.	2.8	9
28	Hepatitis-associated aplastic anemia in pediatric patients: single center experience. Transfusion and Apheresis Science, 2020, 59, 102900.	1.0	5
29	Expanded circulating hematopoietic stem/progenitor cells as novel cell source for the treatment of TCIRG1 osteopetrosis. Haematologica, 2020, 106, 74-86.	3.5	20
30	Whole exome sequencing (WES) approach for diagnosing primary immunodeficiencies (PIDs) in a highly consanguineous community. Clinical Immunology, 2020, 214, 108376.	3.2	22
31	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. Blood, 2020, 136, 2638-2655.	1.4	64
32	Human FCHO1 deficiency reveals role for clathrin-mediated endocytosis in development and function of T cells. Nature Communications, 2020, 11, 1031.	12.8	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. Leukemia and Lymphoma, 2020, 61, 1465-1474.	1.3	4
34	Assesment of Patients with Von Willebrand Disease with ISTH/BAT and PBQ Scores. Turkish Journal of Haematology, 2020, 37, 57-58.	0.5	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.	0.6	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.	0.5	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.	2.1	5
40	CARMIL2 Deficiency Presenting as Very Early Onset Inflammatory Bowel Disease. Inflammatory Bowel Diseases, 2019, 25, 1788-1795.	1.9	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.	3.8	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.	3.8	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.	0.6	8
44	CD137 deficiency causes immune dysregulation with predisposition to lymphomagenesis. Blood, 2019, 134, 1510-1516.	1.4	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.	0.9	3
46	Propranolol treatment for chylothorax due to diffuse lymphangiomatosis. Pediatric Blood and Cancer, 2019, 66, e27592.	1.5	11
47	Capturing B type acute lymphoblastic leukemia cells using two types of antibodies. Biotechnology Progress, 2019, 35, e2737.	2.6	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.	3.8	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.	4.1	1
50	Biliary Rhabdomyosarcoma in an Infant Male With Neurofibromatosis Type 1. Journal of Pediatric Hematology/Oncology, 2019, 41, e24-e26.	0.6	1
51	Defective glycosylation and multisystem abnormalities characterize the primary immunodeficiency XMEN disease. Journal of Clinical Investigation, 2019, 130, 507-522.	8.2	74
52	Vena Cava Superior Syndrome in Children With Mediastinal Tumors: Single Center Experience. Ä°stanbul Kuzey Klinikleri, 2019, 7, 255-259.	0.3	5
53	A rare cause of vomiting in an adolescent: gastric burkitt`s lymphoma. Turkish Journal of Pediatrics, 2019, 61, 431.	0.6	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.	0.2	0

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55	Novel Immune Checkpoint Deficiency and Susceptibility to EBV-Associated Lymphoma. Blood, 2019, 134, 2326-2326.	1.4	0
56	Machine Learning Unveils Proteotypic Mimicry in Genetically Defined SCN Variants. Blood, 2019, 134, 3580-3580.	1.4	0
57	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.	0.6	10
58	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.	2.9	10
59	The relationship between hematological parameters and prognosis of children with acute ischemic stroke. Child's Nervous System, 2018, 34, 655-661.	1.1	14
60	Novel Mutations in RASGRP1 are Associated with Immunodeficiency, Immune Dysregulation, and EBV-Induced Lymphoma. Journal of Clinical Immunology, 2018, 38, 699-710.	3.8	37
61	Genetic Polymorphism of VKORC1-1639 in Children With Intracranial Hemorrhage Due to Vitamin K Deficiency. Clinical and Applied Thrombosis/Hemostasis, 2018, 24, 89S-93S.	1.7	3
62	Juvenile Myelomonocytic Leukemia in Turkey: A Retrospective Analysis of Sixty-five PatientsÃ–zlem TÃ¼fekÃ§i1. Turkish Journal of Haematology, 2018, 35, 27-34.	0.5	11
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.	0.5	10
64	Scurvy: A rare cause of arthritis in a child with neurologic disorder. European Journal of Rheumatology, 2018, 5, 283-284.	0.6	1
65	Pansitopeni ile baÅŸvuran metilmalonik asidemi: Olgu sunumu. Medical Journal of Bakirkoy, 2018, , 138-41.	0.1	0
66	Next Generation Proteomics of Human Neutrophil Granulocytes in Monogenic Disease. Blood, 2018, 132, 15-15.	1.4	0
67	Central Nervous System Fungal Infections in Children with Leukemia, Risk Factors and Outcome: A Multicentric Study. Blood, 2018, 132, 5169-5169.	1.4	1
68	Functional Selectivity in Cytokine Signaling Revealed Through a Pathogenic EPO Mutation. Cell, 2017, 168, 1053-1064.e15.	28.9	98
69	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
70	Apparent diffusion coefficient in differentiation of pediatric posterior fossa tumors. Japanese Journal of Radiology, 2017, 35, 448-453.	2.4	32
71	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.	1.1	5
72	Neuroblastoma in a Child With Wolf-Hirschhorn Syndrome. Journal of Pediatric Hematology/Oncology, 2017, 39, e224-e226.	0.6	9

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73	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.	0.6	4
74	Evaluation of childhood pancreas solid pseudopapillary tumors. Ä°stanbul Kuzey Klinikleri, 2017, 5, 207-210.	0.3	3
75	Hematopoietic Stem Cell Transplant for Primary Immunodeficiency Diseases: A Single-Center Experience. Experimental and Clinical Transplantation, 2017, 15, 337-343.	0.5	3
76	Utility of the Aspergillus galactomannan antigen testing for neutropenic paediatric patients. Infezioni in Medicina, 2017, 25, 38-44.	1.1	2
77	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.	7.1	137
78	Diffusion weighted imaging in differentiating malignant and benign neuroblastic tumors. Japanese Journal of Radiology, 2016, 34, 620-624.	2.4	17
79	Fibromuscular Dysplasia Complicated With Cerebral Stroke in a Child With Congenital Dyserythropoietic Anemia Type II. Journal of Pediatric Hematology/Oncology, 2016, 38, e333-e335.	0.6	3
80	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.	1.1	3
81	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.	0.5	1
82	Discovery of the First Pathogenic Human EPO Mutation Provides Mechanistic Insight into Cytokine Signaling. Blood, 2016, 128, 331-331.	1.4	6
83	Targeted high-throughput sequencing for genetic diagnostics of hemophagocytic lymphohistiocytosis. Genome Medicine, 2015, 7, 130.	8.2	37
84	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.	0.8	3
85	Diagnostic and prognostic significance of glypicanÂ5 and glypicanÂ6 gene expression levels in gastric adenocarcinoma. Molecular and Clinical Oncology, 2015, 3, 584-590.	1.0	16
86	A case of congenital afibrinogenemia complicated with thromboembolic events that required repeated amputations. Blood Coagulation and Fibrinolysis, 2015, 26, 354-356.	1.0	14
87	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.	0.6	9
88	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
89	Malignancies in Primary Immunodeficiencies: A Single Center Experience. Pediatric, Allergy, Immunology, and Pulmonology, 2015, 28, 47-54.	0.8	3
90	The extended clinical phenotype of 64 patients with dedicator of cytokinesis 8 deficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 402-412.	2.9	163

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91	A case of XMEN syndrome presented with severe auto-immune disorders mimicking autoimmune lymphoproliferative disease. Clinical Immunology, 2015, 159, 58-62.	3.2	41
92	Torticollis in children: an alert symptom not to be turned away. Child's Nervous System, 2015, 31, 1461-1470.	1.1	30
93	Stem cell mobilization and collection from pediatric patients and healthy children. Transfusion and Apheresis Science, 2015, 53, 17-22.	1.0	31
94	Brain Abscesses in Children. Journal of Child Neurology, 2015, 30, 458-467.	1.4	15
95	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.	0.5	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. Journal of Clinical Immunology, 2014, 34, 792-795.	3.8	30
97	Inherited biallelic CSF3R mutations in severe congenital neutropenia. Blood, 2014, 123, 3811-3817.	1.4	79
98	Childhood Stroke: Results of 130 Children From a Reference Center in Central Anatolia, Turkey. Pediatric Neurology, 2014, 50, 595-600.	2.1	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. Child's Nervous System, 2014, 30, 227-240.	1.1	37
100	Is there relation between COL4A1/A2 mutations and antenatally detected fetal intraventricular hemorrhage?. Child's Nervous System, 2014, 30, 419-424.	1.1	12
101	Neurological complication of non Hodgkin lymphoma in childhood: experience from a single center in Turkey. Child's Nervous System, 2014, 30, 639-645.	1.1	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. Child's Nervous System, 2014, 30, 1375-1382.	1.1	15
103	Cranial metastatic alveolar rhabdomyosarcoma mimicking hematological malignancy in an adolescent boy. Child's Nervous System, 2014, 30, 1737-1741.	1.1	3
104	Fetal intracranial hemorrhage related to maternal autoimmune thrombocytopenic purpura. Child's Nervous System, 2014, 30, 2147-2150.	1.1	8
105	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.	2.9	160
106	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.	2.8	10
107	Acute Colchicine Intoxication Complicated With Extramedullary Hematopoiesis Due to Filgrastim in a Child. Journal of Pediatric Hematology/Oncology, 2014, 36, e460-e462.	0.6	12
108	Infantile tremor syndrome associated with cobalamin therapy: A case report. Clinical Neurology and Neurosurgery, 2013, 115, 1903-1905.	1.4	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. Blood, 2013, 121, 1345-1356.	1.4	122
110	Myeloperoxidase Deficiency: The Secret Under the Flag of Unstained Cell. Turkish Journal of Haematology, 2013, 30, 232-233.	0.2	3
111	The Efficacy of Vitamin K2 and Calcitriol Combination on Thalassemic Osteopathy. Journal of Pediatric Hematology/Oncology, 2013, 35, 623-627.	0.6	13
112	Gastric Signet Ring Carcinoma in a Patient With Ataxia-Telangiectasia. Journal of Pediatric Hematology/Oncology, 2013, 35, e341-e343.	0.6	12
113	Chronic granulomatous disease with markedly elevated IgE levels mimicking hyperimmunoglobulin E syndrome. Acta Microbiologica Et Immunologica Hungarica, 2013, 60, 155-162.	0.8	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. Blood, 2013, 122, 1025-1025.	1.4	0
115	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
116	Ataksi telanjiektazi ve ikincil hastalÄ±klar. Turk Pediatri Arsivi, 2012, 47, 38-42.	0.9	0
117	Coenzyme Q₁₀Levels in Î²-Thalassemia and its Association with Ferritin Levels and Chelation Therapy. Hemoglobin, 2012, 36, 219-229.	0.8	2
118	Hashimoto thyroiditis associated with ataxia telangiectasia. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, .	0.9	5
119	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.	0.8	22
120	Cerebellar hemangioblastoma associated with diffuse neonatal hemangiomatosis in an infant. Child's Nervous System, 2012, 28, 1801-1805.	1.1	9
121	Multiple Fungal Brain Abscesses in a Child with Acute Lymphoblastic Leukemia. Mycopathologia, 2012, 174, 505-509.	3.1	12
122	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.	5.3	20
123	Accidental intramuscular overdose administration of vincristine. Drug and Chemical Toxicology, 2012, 35, 232-234.	2.3	3
124	Hematologically important mutations: Leukocyte adhesion deficiency (first update). Blood Cells, Molecules, and Diseases, 2012, 48, 53-61.	1.4	147
125	Hemophagocytic syndrome in a 4â€monthâ€old infant with biotinidase deficiency. Pediatric Blood and Cancer, 2012, 59, 191-193.	1.5	23
126	Orbital myositis associated with focal active colitis in a teenage girl. Child's Nervous System, 2012, 28, 641-643.	1.1	5

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127	Late-type vitamin K deficiency bleeding: experience from 120 patients. Child's Nervous System, 2012, 28, 247-251.	1.1	28
128	The importance of MTHFR polymorphisms in pediatric cerebral stroke. Child's Nervous System, 2012, 28, 13-13.	1.1	8
129	Atrial fibrillation as an uncommon presentation in a large pleomorphic xanthoastrocytoma. Child's Nervous System, 2012, 28, 475-479.	1.1	3
130	Hashimoto thyroiditis associated with ataxia telangiectasia. Journal of Pediatric Endocrinology and Metabolism, 2012, 25, 349-52.	0.9	2
131	Phenylketonuria With Acute Myeloblastic Leukemia in a 9-year-old Boy. Journal of Pediatric Hematology/Oncology, 2011, 33, e256-e257.	0.6	1
132	Mycophenolate mofetil-induced pseudotumor cerebri in a boy with autoimmune lymphoproliferative disease. Child's Nervous System, 2011, 27, 853-855.	1.1	8
133	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.	1.1	21
134	Intracranial hemorrhage in children with congenital factor deficiencies. Child's Nervous System, 2011, 27, 1963-1966.	1.1	27
135	Neuroblastoma Arising From an Unresected Sacrococcygeal Teratoma in a Child. Journal of Pediatric Hematology/Oncology, 2010, 32, 233-235.	0.6	10
136	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.	1.1	26
137	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.	1.1	1
138	Release of N-terminal pro-brain natriuretic peptide in children with acute rheumatic carditis. Cardiology in the Young, 2010, 20, 297-301.	0.8	9
139	CARDIAC FUNCTIONS EVALUATED WITH TISSUE DOPPLER IMAGING IN CHILDHOOD CANCERS TREATED WITH ANTHRACYCLINES. Pediatric Hematology and Oncology, 2010, 27, 13-23.	0.8	15
140	Hypothermia in a Child With Hodgkin Disease. Journal of Pediatric Hematology/Oncology, 2009, 31, 136-138.	0.6	4
141	Merkel Cell Carcinoma in a Child. Journal of Pediatric Hematology/Oncology, 2009, 31, 359-361.	0.6	16
142	Astroblastoma in a child. Child's Nervous System, 2008, 24, 165-168.	1.1	25
143	Pilocytic astrocytoma developing at the site of a previously treated medulloblastoma in a child. Child's Nervous System, 2008, 24, 289-292.	1.1	3
144	Cerebral venous sinus thrombosis in an adolescent with Ewing sarcoma. Child's Nervous System, 2008, 24, 983-986.	1.1	13

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145	Malignant glioblastomatous transformation of a low-grade glioma in a child. Child's Nervous System, 2008, 24, 1385-1389.	1.1	15
146	Importance of chemoprophylaxis and vaccines in the prevention of recurrent meningitis. Pediatrics International, 2008, 50, 416-416.	0.5	0
147	Mesenchymal Hamartoma of the Liver Mimicking Hepatoblastoma. Journal of Pediatric Hematology/Oncology, 2008, 30, 458-460.	0.6	20
148	Fatal valproate overdose in a newborn baby. Human and Experimental Toxicology, 2007, 26, 453-456.	2.2	9
149	Letter to the Editor. Inhalation Toxicology, 2007, 19, 587-587.	1.6	3
150	KLUVER-BUCY SYNDROME IN A BOY WITH NON-HODGKIN LYMPHOMA. Pediatric Hematology and Oncology, 2007, 24, 149-152.	0.8	3
151	Thyroid Medullary Carcinoma in a Teenager With Cowden Syndrome. Laryngoscope, 2007, 117, 1180-1182.	2.0	9
152	Pituitary?adrenal axis suppression due to topical steroid administration in an infant. Pediatrics International, 2007, 49, 242-244.	0.5	9
153	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.	0.8	9
154	Pulmonary arterial pressure in infants with laryngomalacia. International Journal of Pediatric Otorhinolaryngology, 2006, 70, 2067-2071.	1.0	4
155	The Real Incidence of Thyroid Carcinoma in Childhood. Laryngoscope, 2006, 116, 2095.	2.0	0
156	Refractory cutaneous leishmaniasis in an adolescent: Initial manifestation of type 1 diabetes. Journal of Infection, 2006, 53, 290-291.	3.3	1