Tiraje Celkan

List of Publications by Year
in descending order

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127 papers	1,391 citations	430874 18 h-index	414414 32 g-index
130 all docs	130 docs citations	130 times ranked	2041 citing authors

#	Article	IF	Citations
1	Thrombosis in the Vena Cava Inferior and Right Atrium in a Patient with Wilms Tumor. , 2023, 56, 535-536.		0
2	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
3	Impact of having a sibling with cancer or type I diabetes mellitus on psychopathology and self-conscious emotions in adolescents: a comparative study including controls. Supportive Care in Cancer, 2022, 30, 635-645.	2.2	1
4	Adverse COVIDâ€19 outcomes in immune deficiencies: Inequality exists between subclasses. Allergy: European Journal of Allergy and Clinical Immunology, 2022, 77, 282-295.	5.7	29
5	Increased risk for kidney sequelae surrogates in survivors of Wilms tumor. Pediatric Nephrology, 2022, 37, 2415-2426.	1.7	2
6	Telemedicine Applications in a Tertiary Pediatric Hospital in Turkey During COVID-19 Pandemic. Telemedicine Journal and E-Health, 2021, 27, 1180-1187.	2.8	16
7	Comparison of outcomes of children with acute lymphoblastic leukemia treated with BMF protocol across 2 decades. Pediatric Hematology and Oncology, 2021, 38, 134-146.	0.8	1
8	Investigation of (epi)genotype causes and followâ€up manifestations in the patients with classical and atypical phenotype of Beckwithâ€Wiedemann spectrum. American Journal of Medical Genetics, Part A, 2021, 185, 1721-1731.	1.2	5
9	Clinical course of pediatric large vascular anomalies located in the extremities. Turkish Archives of Pediatrics, 2021, 56, 213-218.	0.4	4
10	Neurofibromatosis type 1: Expanded variant spectrum with multiplex ligationâ€dependent probe amplification and genotypeâ€“phenotype correlation in 138 Turkish patients. Annals of Human Genetics, 2021, 85, 155-165.	0.8	4
11	Prognostic evidence of LEF1 isoforms in childhood acute lymphoblastic leukemia. International Journal of Laboratory Hematology, 2021, 43, 1093-1103.	1.3	4
12	Neurofibromatosis Type 1 in Children: A Single-Center Experience. , 2021, 56, 339-343.		0
13	A Peculiar Disease of a Young Woman who Wants to Get Pregnant. Turkish Journal of Haematology, 2021, 38, 333-334.	0.5	0
14	Hypoxic Ischemic Encephalopathy in Forensic Medicine. Adli TÄ±p BÃ¼lteni, 2021, 26, 205-209.	0.1	0
15	Adrenal masses in children: Imaging, surgical treatment and outcome. Asian Journal of Surgery, 2020, 43, 207-212.	0.4	12
16	A novel combined treatment for plasminogen deficiency with lung involvement. Pediatric Pulmonology, 2020, 55, E1-E3.	2.0	3
17	Hemogram bize neler sÃ¶yler?. Turk Pediatri Arsivi, 2020, 55, 103-116.	0.9	10
18	Ligneous gingivitis: Hard to diagnose and treat. Haemophilia, 2020, 26, e49-e50.	2.1	3

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19	Accidental High Dose Intrathecal Treatment - Late Result of A Patient. Turkish Journal of Haematology, 2020, 37, 64-65.	0.5	0
20	PTEN and AKT1 Variations in Childhood T-cell Acute Lymphoblastic Leukemia. Turkish Journal of Haematology, 2020, 37, 98-103.	0.5	1
21	Cross-sectional study: long term follow-up care for pediatric cancer survivors in a developing country, Turkey: current status, challenges, and future perspectives. Turkish Journal of Medical Sciences, 2020, 50, 1916-1921.	0.9	6
22	OUTCOMES OF ELTROMBOPAG TREATMENT AND DEVELOPMENT OF IRON DEFICIENCY IN CHILDREN WITH IMMUNE THROMBOCYTOPENIA IN TURKEY. Turkish Journal of Haematology, 2020, 37, 139-144.	0.5	6
23	An Unusual Presentation of Carney Complex. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 117-121.	0.9	3
24	Homozygous c.130â€“131 ins A (pW44X) mutation in the HAX1 gene as the most common cause of congenital neutropenia in Turkey: Report from the Turkish Severe Congenital Neutropenia Registry. Pediatric Blood and Cancer, 2019, 66, e27923.	1.5	16
25	LANGERANS CELL HISTIOCYTOSIS: SINGLE CENTER EXPERIENCE OF 25 YEARS. Mediterranean Journal of Hematology and Infectious Diseases, 2019, 11, e2019035.	1.3	12
26	Posterior reversible encephalopathy syndrome in children with acute lymphoblastic leukemia: Experience of a single center using BFM protocols. Pediatric Blood and Cancer, 2019, 66, e27711.	1.5	8
27	Frequency of ALK and GD2 Expression in Neuroblastoma. Fetal and Pediatric Pathology, 2019, 38, 326-334.	0.7	8
28	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
29	Pediatric Stroke: A Single-Center Experience. Journal of Pediatric Hematology/Oncology, 2019, 41, 519-524.	0.6	8
30	Ã‡ocukluk Ã§aÄŸÄ± medulloblastom olgularÄ±nda P53, ERBB2, c-Kit ve BCL2 Ekspresyonunun prognostik ve klinik Ã¶nemi. Cukurova Medical Journal, 2019, 44, 1-7.	0.2	0
31	Homozygous protein C deficiency presenting as neonatal purpura fulminans: management with fresh frozen plasma, low molecular weight heparin and protein C concentrate. Journal of Thrombosis and Thrombolysis, 2018, 45, 315-318.	2.1	8
32	Sirolimus Experience in Blue Rubber Bleb Nevus Syndrome. Journal of Pediatric Hematology/Oncology, 2018, 40, 168-169.	0.6	10
33	Treatment of plasminogen deficiency patients with fresh frozen plasma. Pediatric Blood and Cancer, 2018, 65, e26779.	1.5	17
34	Coagulation Disturbances in Patients with Argininemia. Acta Haematologica, 2018, 140, 221-225.	1.4	5
35	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
36	Use of a High-Purity Factor X Concentrate in Turkish Subjects with Hereditary Factor X Deficiency: Post Hoc Cohort Subanalysis of a Phase 3 Study. Turkish Journal of Haematology, 2018, 35, 129-133.	0.5	2

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37	Thrombosis in children: Which test to whom, when and how much necessary?. Turk Pediatri Arsivi, 2018, 53, 1-9.	0.9	26
38	Association of the TLR4 and NOD2 Polymorphisms with Childhood Acute Lymphoblastic Leukemia. BezmiÃ¢lem Science, 2018, 6, 119-125.	0.2	1
39	Human Plasminogen Replacement Therapy Vs. Best Supportive Care in Patients with Congenital Plasminogen Deficiency: A Comparison of Health Care Resource Use over a 48-Week Period. Blood, 2018, 132, 2232-2232.	1.4	0
40	Central Nervous System Fungal Infections in Children with Leukemia, Risk Factors and Outcome: A Multicentric Study. Blood, 2018, 132, 5169-5169.	1.4	1
41	Influence of Paroxysmal Nocturnal Hemoglobinuria Clone Positivity on Outcome of Childhood Acquried Aplastic Anemia: A Multicenter Center Study. Blood, 2018, 132, 5101-5101.	1.4	0
42	Evaluation of high-risk features of primary enucleation of patients with retinoblastoma in a tertiary center of a developing country in the era of intra-arterial chemotherapy. International Ophthalmology, 2017, 38, 151-156.	1.4	2
43	Sinusoidal Obstruction Syndrome During Chemotherapy of Pediatric Cancers and its Successful Management With Defibrotide. Journal of Pediatric Hematology/Oncology, 2017, 39, e373-e376.	0.6	7
44	Plasminogen deficiency. Journal of Thrombosis and Thrombolysis, 2017, 43, 132-138.	2.1	29
45	A child presenting with hypercalcemia. Turk Pediatri Arsivi, 2017, 49, 81-83.	0.9	8
46	Wernickeâ€™s Encephalopathy in a Child with Acute Lymphoblastic Leukemia. Turkish Journal of Haematology, 2017, 34, 99-100.	0.5	0
47	Hemoglobin H Disease in Turkey: Experience from Eight Centers. Turkish Journal of Haematology, 2016, 33, 56-59.	0.5	4
48	Prenatal unilateral cerebellar hypoplasia diagnosed as PHACE syndrome. Child's Nervous System, 2016, 32, 587-588.	1.1	3
49	Novel plasminogen gene mutations in Turkish patients with type I plasminogen deficiency. Blood Coagulation and Fibrinolysis, 2016, 27, 637-644.	1.0	5
50	Turkish National Severe Congenital Neutropenia Registry. Blood, 2016, 128, 4916-4916.	1.4	2
51	Febrile neutropenia in children with acute lymphoblastic leukemia: single center experience. Turk Pediatri Arsivi, 2016, 51, 79-86.	0.9	15
52	Five-year-old girl with tongue bleeding. Turk Pediatri Arsivi, 2016, 51, 117-119.	0.9	0
53	DNA Repair Gene Polymorphisms and Their Relation With DNA Damage, DNA Repair, and Total Antioxidant Capacity in Childhood Acute Lymphoblastic Leukemia Survivors. Journal of Pediatric Hematology/Oncology, 2015, 37, 344-350.	0.6	12
54	Approach to the patient with neutropenia in childhood. Turk Pediatri Arsivi, 2015, 50, 136-144.	0.9	30

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55	Childhood mastocytosis: results of a single center. Turk Pediatri Arsivi, 2015, 50, 108-113.	0.9	6
56	Role of glutathione S-transferase M1, T1 and P1 gene polymorphisms in childhood acute lymphoblastic leukemia susceptibility in a Turkish population. Meta Gene, 2015, 5, 115-119.	0.6	7
57	Initial Management of Childhood Acute Immune Thrombocytopenia: Single-Center Experience of 32 Years. Pediatric Hematology and Oncology, 2015, 32, 406-414.	0.8	4
58	A possible role for WNT5A hypermethylation in Pediatric Acute Lymphoblastic Leukemia. Turkish Journal of Haematology, 2015, 32, 127-135.	0.5	10
59	Evaluation of Alpha-Thalassemia Mutations in Cases with Hypochromic Microcytic Anemia: The Ä°stanbul Perspective. Turkish Journal of Haematology, 2015, 32, 344-350.	0.5	11
60	The Role of Epstein-Barr Virus LMP-1 Immunohistochemical Staining in Childhood Hodgkin Lymphoma. Iranian Journal of Pediatrics, 2015, 25, e2359.	0.3	5
61	Successful management of hepatic mucormycosis in an acute lymphoblastic leukaemia patient: a case report and review of the literature. Mycoses, 2014, 57, 513-518.	4.0	6
62	Chediak-Higashi Syndrome: A Case Report of a Girl Without Silvery Hair and Oculocutaneous Albinism Presenting with Hemophagocytic Lymphohistiocytosis. Turkish Journal of Haematology, 2014, 31, 426-427.	0.5	0
63	Blue Toe Syndromeâ€"Reply. JAMA Ophthalmology, 2014, 132, 654.	2.5	0
64	A 17-year experience with ALL-BFM protocol in acute lymphoblastic leukemia: Prognostic predictors and interruptions during protocol. Leukemia Research, 2014, 38, 699-705.	0.8	10
65	Spontaneous splenic rupture in a patient with congenital afibrinogenemia. Turk Pediatri Arsivi, 2014, 49, 247-249.	0.9	14
66	A rare cause of thrombocyte dysfunction: Hermansky-Pudlak syndrome. Turk Pediatri Arsivi, 2014, 49, 163-166.	0.9	5
67	An infant with chronic hemolytic anemia. Turk Pediatri Arsivi, 2014, 49, 264-268.	0.9	0
68	Febrile Neutropenia in Children with Acute Lymphoblastic Leukemia Treated with BFM Protocols. Blood, 2014, 124, 5245-5245.	1.4	0
69	Hodgkinâ€™s Disease (1975-2012): Long-Term Results of a Single Center. Blood, 2014, 124, 5362-5362.	1.4	0
70	Novel use of propranolol for management of pain in children with vertebral hemangioma: report of two cases. Child's Nervous System, 2013, 29, 855-860.	1.1	19
71	Effect of Prothrombotic Mutations on Factor Consumption in Children With Hemophilia. Clinical and Applied Thrombosis/Hemostasis, 2013, 19, 445-448.	1.7	4
72	Tumour lysis syndrome; new approaches at diagnose, follow up and treatment. Turk Pediatri Arsivi, 2013, 48, 188-194.	0.9	0

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73	A newborn with subcutaneous nodules. Turk Pediatri Arsivi, 2013, 48, 259-262.	0.9	0
74	KaraciÄŸer hastalÄ±klarÄ±nda pÄ±htÄ±laÅŸma-tromboz mekanizmasÄ±na neler oluyor?. Turk Pediatri Arsivi, 2013, 48, 94-101.	0.9	0
75	On dÃ¶rt yaÅŸÄ±nda akut lenfoblastik lÃ¶semili hastada dÃ¼zelmeyen taÅŸikardi. Turk Pediatri Arsivi, 2013, 48, 176-178.	0.9	0
76	Macrocephaly-Capillary Malformation Syndrome in a Newborn With Tetralogy of Fallot and Sagittal Sinus Thrombosis. Journal of Child Neurology, 2013, 28, 115-119.	1.4	6
77	Voriconazole Induced Bradycardia. Pediatric Hematology and Oncology, 2013, 30, 674-676.	0.8	5
78	Genetic and Epigenetic Profile Of Early Relapsed Childhood ALL. Blood, 2013, 122, 1380-1380.	1.4	0
79	Analysis of Chromosomal Aberrations and FLT3 gene Mutations in Childhood Acute Myelogenous Leukemia Patients. Turkish Journal of Haematology, 2012, 29, 225-232.	0.5	6
80	RARE COAGULATION DISORDERS. Retrospective analyses of 156 patients in TURKEY. Turkish Journal of Haematology, 2012, 29, 48-54.	0.5	11
81	Upregulation of T-Cell-Specific Transcription Factor Expression in Pediatric T-Cell Acute Lymphoblastic Leukemia (T-ALL). Turkish Journal of Haematology, 2012, 29, 325-333.	0.2	5
82	Erythrocyte membrane protein defects in hereditary spherocytosis patients in Turkish population. Hematology, 2012, 17, 232-236.	1.5	3
83	Evaluation of PAX5 gene in the early stages of leukemic B cells in the childhood B cell acute lymphoblastic leukemia. Leukemia Research, 2012, 36, 87-92.	0.8	12
84	DNA repair gene XPD and XRCC1 polymorphisms and the risk of febrile neutropenia and mucositis in children with leukemia and lymphoma. Leukemia Research, 2012, 36, 565-569.	0.8	18
85	2011 yÄ±lÄ±nda Ã§ocukluk Ã§aÄŸÄ± immÃ¼n trombositopenik purpura hastalÄ±ÄŸÄ± izlem ve tedavisinde deÄŸiÅŸiklikler. Turk Pediatri Arsivi, 2012, 47, 8-16.	0.9	1
86	Novel Influenza a (H1N1) Infection in a Pediatric Hematology Oncology Clinic During the 2009â€“2010 Pandemia. Pediatric Hematology and Oncology, 2011, 28, 288-293.	0.8	16
87	Daily and every other day use of iron prophylaxis in the first year of life. Turk Pediatri Arsivi, 2011, 46, 184.	0.9	0
88	Cytopenia associated with iron deficiency anemia and iron therapy: A report of two cases. Turkish Journal of Haematology, 2011, 28, 243-244.	0.5	5
89	Kanama bozukluÄŸu olan Ã§ocuklarda sÃ¼nnet deneyimi. Turk Pediatri Arsivi, 2011, 46, 313-317.	0.9	0
90	Trombositozu olan bir yaÅŸÄ±nda erkek Ã§ocuk. Turk Pediatri Arsivi, 2011, 46, 351-352.	0.9	0

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91	Ã‡ocukluk Ã§aÄŸÄ± akut lenfoblastik lÃ¶semisinde tedavinin ortalama eritrosit hacimleri Ã¼zerine etkisi. Turk Pediatri Arsivi, 2011, 46, 133-136.	0.9	1
92	Niemann-Pick disease. Turkish Journal of Haematology, 2011, 28, 84-85.	0.5	0
93	Non-Hodgkinâ€™s lymphoma in a hemophilic patient with a traumatic hematoma. Turkish Journal of Haematology, 2011, 28, 237-238.	0.5	1
94	Treatment of Pediatric Burkitt Lymphoma in Turkey. Journal of Pediatric Hematology/Oncology, 2010, 32, e279-e284.	0.6	6
95	Spontaneous intracranial bleeding in a neonate with congenital afibrinogenemia. Blood Coagulation and Fibrinolysis, 2010, 21, 592-594.	1.0	6
96	Acute Promyelocytic Leukemia Treated With Idarubicin Complicated by Focal Segmental Glomerulosclerosis. Journal of Pediatric Hematology/Oncology, 2010, 32, e82-e84.	0.6	9
97	Prognostic Significance ofNOTCH1andFBXW7Mutations in Pediatric T-ALL. Disease Markers, 2010, 28, 353-360.	1.3	25
98	Clinical and laboratory data of primary hemophagocytic lymphohistiocytosis: A retrospective review of the Turkish Histiocyte Study Group. Turkish Journal of Haematology, 2010, 27, 257-262.	0.5	4
99	PSEUDOPROGRESSION AFTER RADIOTHERAPY WITH CONCURRENT TEMOZOLOMIDE IN A CHILD WITH ANAPLASTIC ASTROCYTOMA. Pediatric Hematology and Oncology, 2010, 27, 317-319.	0.8	2
100	Prognostic significance of NOTCH1 and FBXW7 mutations in pediatric T-ALL. Disease Markers, 2010, 28, 353-60.	1.3	18
101	Aplastic anemia presenting as hemophagocytic lymphohistiocytosis. Turkish Journal of Haematology, 2010, 27, 38-42.	0.5	1
102	ASSOCIATION BETWEEN GENETIC POLYMORPHISM IN DNA REPAIR GENES AND RISK OF B-CELL LYMPHOMA. Pediatric Hematology and Oncology, 2009, 26, 467-472.	0.8	14
103	A rare cause of ischemic stroke: fibromuscular dysplasia. Neurological Sciences, 2009, 30, 77-79.	1.9	5
104	DNA repair gene XPD and XRCC1 polymorphisms and the risk of childhood acute lymphoblastic leukemia. Leukemia Research, 2009, 33, 759-763.	0.8	62
105	FatalTrichodermaharzianuminfection in a leukemic pediatric patient. Medical Mycology, 2009, 47, 207-215.	0.7	26
106	Is Familial Mediterranean Fever a thrombotic disease or not?. European Journal of Pediatrics, 2008, 167, 279-285.	2.7	34
107	Coinheritance of sickle cell anemia and hereditary spherocytosis. Pediatric Blood and Cancer, 2008, 51, 560-563.	1.5	0
108	Prognostic factors in children with Hodgkin disease. Pediatric Blood and Cancer, 2008, 51, 712-712.	1.5	0

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109	Syntaxinâ€11 is expressed in primary human monocytes/macrophages and acts as a negative regulator of macrophage engulfment of apoptotic cells and IgGâ€opsonized target cells. British Journal of Haematology, 2008, 142, 469-479.	2.5	29
110	The difference between pre-B cell acute lymphoblastic leukemia and Burkitt lymphoma in relation to DNA damage repair gene polymorphisms in childhood. Leukemia and Lymphoma, 2008, 49, 1638-1640.	1.3	16
111	CONTROL OF BLEEDING ASSOCIATED WITH HEMOPHAGOCYTIC SYNDROME IN CHILDREN: An Audit of the Clinical use of Recombinant Activated Factor VII. Pediatric Hematology and Oncology, 2007, 24, 117-121.	0.8	10
112	Defective cytotoxic lymphocyte degranulation in syntaxin-11â€“deficient familial hemophagocytic lymphohistiocytosis 4 (FHL4) patients. Blood, 2007, 110, 1906-1915.	1.4	272
113	Antiviral prophylaxis with continuous low dose acyclovir in childhood cancer. Leukemia and Lymphoma, 2006, 47, 1418-1420.	1.3	3
114	Retinoblastoma in Turkey: survival and clinical characteristics 1981Ã¢Â€Â“2004. Pediatrics International, 2006, 48, 369-373.	0.5	38
115	Venous and Intrapericardial Thrombosis. Pediatric Cardiology, 2006, 27, 497-499.	1.3	6
116	Varicella-induced hemolytic anemia with hepatitis. Annals of Hematology, 2006, 85, 64-65.	1.8	7
117	Antibody Titers and Immune Response to Diphtheria-Tetanus-Pertussis and Measles-Mumps-Rubella Vaccination in Children Treated for Acute Lymphoblastic Leukemia. Journal of Pediatric Hematology/Oncology, 2005, 27, 273-277.	0.6	54
118	Prevalence of Anemia and the Risk Factors Among Schoolchildren in Istanbul. Journal of Tropical Pediatrics, 2005, 51, 346-350.	1.5	16
119	THE ANEMIA OF FAMILIAL MEDITERRANEAN FEVER DISEASE. Pediatric Hematology and Oncology, 2005, 22, 657-665.	0.8	22
120	Outcome after Relapse in Childhood Acute Lymphoblastic Leukaemia - Results of a Single Center for a Period of 15 Years.. Blood, 2005, 106, 4582-4582.	1.4	0
121	NAD(P)H:quinone oxidoreductase 1 null genotype is not associated with pediatric de novo acute leukemia. Pediatric Blood and Cancer, 2004, 43, 568-570.	1.5	31
122	Anemic or not?. Turkish Journal of Pediatrics, 2003, 45, 329-34.	0.6	0
123	Bacteremia in Childhood Cancer. Journal of Tropical Pediatrics, 2002, 48, 373-376.	1.5	25
124	Increased factor VIII activity and dural sinus thrombosis. Medical and Pediatric Oncology, 2002, 39, 71-72.	1.0	3
125	Radiological features in paediatric primary gastric MALT lymphoma and association with Helicobacter pylori. Pediatric Radiology, 2002, 32, 82-87.	2.0	30
126	Toxic Epidermal Necrolysis After the Use of Highâ€Dose Cytosine Arabinoside. Pediatric Dermatology, 2001, 18, 38-40.	0.9	26

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127	The vanishing bile duct syndrome in a child with Hodgkin disease. Medical and Pediatric Oncology, 2001, 36, 398-399.	1.0	11