Tiraje Celkan

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1094628/publications.pdf

Version: 2024-02-01

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	Defective cytotoxic lymphocyte degranulation in syntaxin-11â€“deficient familial hemophagocytic lymphohistiocytosis 4 (FHL4) patients. Blood, 2007, 110, 1906-1915.	1.4	272
2	DNA repair gene XPD and XRCC1 polymorphisms and the risk of childhood acute lymphoblastic leukemia. Leukemia Research, 2009, 33, 759-763.	0.8	62
3	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
4	Retinoblastoma in Turkey: survival and clinical characteristics 1981Ã¢Â€Â“2004. Pediatrics International, 2006, 48, 369-373.	0.5	38
5	Is Familial Mediterranean Fever a thrombotic disease or not?. European Journal of Pediatrics, 2008, 167, 279-285.	2.7	34
6	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
7	Radiological features in paediatric primary gastric MALT lymphoma and association with Helicobacter pylori. Pediatric Radiology, 2002, 32, 82-87.	2.0	30
8	Approach to the patient with neutropenia in childhood. Turk Pediatri Arsivi, 2015, 50, 136-144.	0.9	30
9	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
10	Plasminogen deficiency. Journal of Thrombosis and Thrombolysis, 2017, 43, 132-138.	2.1	29
11	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
12	Toxic Epidermal Necrolysis After the Use of Highâ€Dose Cytosine Arabinoside. Pediatric Dermatology, 2001, 18, 38-40.	0.9	26
13	FatalTrichodermaharzianuminfection in a leukemic pediatric patient. Medical Mycology, 2009, 47, 207-215.	0.7	26
14	Thrombosis in children: Which test to whom, when and how much necessary?. Turk Pediatri Arsivi, 2018, 53, 1-9.	0.9	26
15	Bacteremia in Childhood Cancer. Journal of Tropical Pediatrics, 2002, 48, 373-376.	1.5	25
16	Prognostic Significance ofNOTCH1andFBXW7Mutations in Pediatric T-ALL. Disease Markers, 2010, 28, 353-360.	1.3	25
17	THE ANEMIA OF FAMILIAL MEDITERRANEAN FEVER DISEASE. Pediatric Hematology and Oncology, 2005, 22, 657-665.	0.8	22
18	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

#	Article	IF	Citations
19	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
20	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
21	Prognostic significance of NOTCH1 and FBXW7 mutations in pediatric T-ALL. Disease Markers, 2010, 28, 353-60.	1.3	18
22	Treatment of plasminogen deficiency patients with fresh frozen plasma. Pediatric Blood and Cancer, 2018, 65, e26779.	1.5	17
23	Prevalence of Anemia and the Risk Factors Among Schoolchildren in Istanbul. Journal of Tropical Pediatrics, 2005, 51, 346-350.	1.5	16
24	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
25	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
26	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
27	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
28	Febrile neutropenia in children with acute lymphoblastic leukemia: single center experience. Turk Pediatri Arsivi, 2016, 51, 79-86.	0.9	15
29	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
30	Spontaneous splenic rupture in a patient with congenital afibrinogenemia. Turk Pediatri Arsivi, 2014, 49, 247-249.	0.9	14
31	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
32	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
33	LANGERANS CELL HISTIOCYTOSIS: SINGLE CENTER EXPERIENCE OF 25 YEARS. Mediterranean Journal of Hematology and Infectious Diseases, 2019, 11, e2019035.	1.3	12
34	Adrenal masses in children: Imaging, surgical treatment and outcome. Asian Journal of Surgery, 2020, 43, 207-212.	0.4	12
35	The vanishing bile duct syndrome in a child with Hodgkin disease. Medical and Pediatric Oncology, 2001, 36, 398-399.	1.0	11
36	RARE COAGULATION DISORDERS. Retrospective analyses of 156 patients in TURKEY. Turkish Journal of Haematology, 2012, 29, 48-54.	0.5	11

#	Article	IF	Citations
37	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
38	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
39	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
40	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
41	Sirolimus Experience in Blue Rubber Bleb Nevus Syndrome. Journal of Pediatric Hematology/Oncology, 2018, 40, 168-169.	0.6	10
42	Hemogram bize neler sÃ¶yler?. Turk Pediatri Arsivi, 2020, 55, 103-116.	0.9	10
43	A possible role for WNT5A hypermethylation in Pediatric Acute Lymphoblastic Leukemia. Turkish Journal of Haematology, 2015, 32, 127-135.	0.5	10
44	Acute Promyelocytic Leukemia Treated With Idarubicin Complicated by Focal Segmental Glomerulosclerosis. Journal of Pediatric Hematology/Oncology, 2010, 32, e82-e84.	0.6	9
45	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
46	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
47	Frequency of ALK and GD2 Expression in Neuroblastoma. Fetal and Pediatric Pathology, 2019, 38, 326-334.	0.7	8
48	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
49	Pediatric Stroke: A Single-Center Experience. Journal of Pediatric Hematology/Oncology, 2019, 41, 519-524.	0.6	8
50	A child presenting with hypercalcemia. Turk Pediatri Arsivi, 2017, 49, 81-83.	0.9	8
51	Varicella-induced hemolytic anemia with hepatitis. Annals of Hematology, 2006, 85, 64-65.	1.8	7
52	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
53	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
54	Venous and Intrapericardial Thrombosis. Pediatric Cardiology, 2006, 27, 497-499.	1.3	6

#	Article	IF	Citations
55	Treatment of Pediatric Burkitt Lymphoma in Turkey. Journal of Pediatric Hematology/Oncology, 2010, 32, e279-e284.	0.6	6
56	Spontaneous intracranial bleeding in a neonate with congenital afibrinogenemia. Blood Coagulation and Fibrinolysis, 2010, 21, 592-594.	1.0	6
57	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
58	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
59	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
60	Childhood mastocytosis: results of a single center. Turk Pediatri Arsivi, 2015, 50, 108-113.	0.9	6
61	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
62	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
63	A rare cause of ischemic stroke: fibromuscular dysplasia. Neurological Sciences, 2009, 30, 77-79.	1.9	5
64	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
65	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
66	Voriconazole Induced Bradycardia. Pediatric Hematology and Oncology, 2013, 30, 674-676.	0.8	5
67	Novel plasminogen gene mutations in Turkish patients with type I plasminogen deficiency. Blood Coagulation and Fibrinolysis, 2016, 27, 637-644.	1.0	5
68	Coagulation Disturbances in Patients with Argininemia. Acta Haematologica, 2018, 140, 221-225.	1.4	5
69	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
70	A rare cause of thrombocyte dysfunction: Hermansky-Pudlak syndrome. Turk Pediatri Arsivi, 2014, 49, 163-166.	0.9	5
71	The Role of Epstein-Barr Virus LMP-1 Immunohistochemical Staining in Childhood Hodgkin Lymphoma. Iranian Journal of Pediatrics, 2015, 25, e2359.	0.3	5
72	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

#	Article	IF	Citations
73	Effect of Prothrombotic Mutations on Factor Consumption in Children With Hemophilia. Clinical and Applied Thrombosis/Hemostasis, 2013, 19, 445-448.	1.7	4
74	Initial Management of Childhood Acute Immune Thrombocytopenia: Single-Center Experience of 32 Years. Pediatric Hematology and Oncology, 2015, 32, 406-414.	0.8	4
75	Hemoglobin H Disease in Turkey: Experience from Eight Centers. Turkish Journal of Haematology, 2016, 33, 56-59.	0.5	4
76	Clinical course of pediatric large vascular anomalies located in the extremities. Turkish Archives of Pediatrics, 2021, 56, 213-218.	0.4	4
77	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
78	Prognostic evidence of LEF1 isoforms in childhood acute lymphoblastic leukemia. International Journal of Laboratory Hematology, 2021, 43, 1093-1103.	1.3	4
79	Increased factor VIII activity and dural sinus thrombosis. Medical and Pediatric Oncology, 2002, 39, 71-72.	1.0	3
80	Antiviral prophylaxis with continuous low dose acyclovir in childhood cancer. Leukemia and Lymphoma, 2006, 47, 1418-1420.	1.3	3
81	Erythrocyte membrane protein defects in hereditary spherocytosis patients in Turkish population. Hematology, 2012, 17, 232-236.	1.5	3
82	Prenatal unilateral cerebellar hypoplasia diagnosed as PHACE syndrome. Child's Nervous System, 2016, 32, 587-588.	1.1	3
83	A novel combined treatment for plasminogen deficiency with lung involvement. Pediatric Pulmonology, 2020, 55, E1-E3.	2.0	3
84	Ligneous gingivitis: Hard to diagnose and treat. Haemophilia, 2020, 26, e49-e50.	2.1	3
85	An Unusual Presentation of Carney Complex. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2020, 12, 117-121.	0.9	3
86	PSEUDOPROGRESSION AFTER RADIOTHERAPY WITH CONCURRENT TEMOZOLOMIDE IN A CHILD WITH ANAPLASTIC ASTROCYTOMA. Pediatric Hematology and Oncology, 2010, 27, 317-319.	0.8	2
87	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
88	Turkish National Severe Congenital Neutropenia Registry. Blood, 2016, 128, 4916-4916.	1.4	2
89	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
90	Increased risk for kidney sequelae surrogates in survivors of Wilms tumor. Pediatric Nephrology, 2022, 37, 2415-2426.	1.7	2

#	Article	IF	Citations
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	Non-Hodgkinâ€™s lymphoma in a hemophilic patient with a traumatic hematoma. Turkish Journal of Haematology, 2011, 28, 237-238.	0.5	1
93	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
94	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
95	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
96	Association of the TLR4 and NOD2 Polymorphisms with Childhood Acute Lymphoblastic Leukemia. BezmiÃ¢lem Science, 2018, 6, 119-125.	0.2	1
97	Central Nervous System Fungal Infections in Children with Leukemia, Risk Factors and Outcome: A Multicentric Study. Blood, 2018, 132, 5169-5169.	1.4	1
98	PTEN and AKT1 Variations in Childhood T-cell Acute Lymphoblastic Leukemia. Turkish Journal of Haematology, 2020, 37, 98-103.	0.5	1
99	Aplastic anemia presenting as hemophagocytic lymphohistiocytosis. Turkish Journal of Haematology, 2010, 27, 38-42.	0.5	1
100	Coinheritance of sickle cell anemia and hereditary spherocytosis. Pediatric Blood and Cancer, 2008, 51, 560-563.	1.5	0
101	Prognostic factors in children with Hodgkin disease. Pediatric Blood and Cancer, 2008, 51, 712-712.	1.5	0
102	Daily and every other day use of iron prophylaxis in the first year of life. Turk Pediatri Arsivi, 2011, 46, 184.	0.9	0
103	Kanama bozukluÄŸu olan Ã§ocuklarda sÃ¼nnet deneyimi. Turk Pediatri Arsivi, 2011, 46, 313-317.	0.9	0
104	Trombositozu olan bir yaÅŸÄ±nda erkek Ã§ocuk. Turk Pediatri Arsivi, 2011, 46, 351-352.	0.9	0
105	Niemann-Pick disease. Turkish Journal of Haematology, 2011, 28, 84-85.	0.5	0
106	Tumour lysis syndrome; new approaches at diagnose, follow up and treatment. Turk Pediatri Arsivi, 2013, 48, 188-194.	0.9	0
107	A newborn with subcutaneous nodules. Turk Pediatri Arsivi, 2013, 48, 259-262.	0.9	0
108	KaraciÄŸer hastalÄ±klarÄ±nda pÄ±htÄ±laÅŸma-tromboz mekanizmasÄ±na neler oluyor?. Turk Pediatri Arsivi, 2013, 48, 94-101.	0.9	0

#	Article	IF	Citations
109	On dÃ¶rt yaÅŸÄ±nda akut lenfoblastik lÃ¶semili hastada dÃ¼zelmeyen taÅŸikardi. Turk Pediatri Arsivi, 2013, 48, 176-178.	0.9	0
110	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
111	Blue Toe Syndromeâ€"Reply. JAMA Ophthalmology, 2014, 132, 654.	2.5	0
112	Neurofibromatosis Type 1 in Children: A Single-Center Experience. , 2021, 56, 339-343.		0
113	A Peculiar Disease of a Young Woman who Wants to Get Pregnant. Turkish Journal of Haematology, 2021, 38, 333-334.	0.5	0
114	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
115	Genetic and Epigenetic Profile Of Early Relapsed Childhood ALL. Blood, 2013, 122, 1380-1380.	1.4	0
116	An infant with chronic hemolytic anemia. Turk Pediatri Arsivi, 2014, 49, 264-268.	0.9	0
117	Febrile Neutropenia in Children with Acute Lymphoblastic Leukemia Treated with BFM Protocols. Blood, 2014, 124, 5245-5245.	1.4	0
118	Hodgkinâ€™s Disease (1975-2012): Long-Term Results of a Single Center. Blood, 2014, 124, 5362-5362.	1.4	0
119	Five-year-old girl with tongue bleeding. Turk Pediatri Arsivi, 2016, 51, 117-119.	0.9	0
120	Wernickeâ€™s Encephalopathy in a Child with Acute Lymphoblastic Leukemia. Turkish Journal of Haematology, 2017, 34, 99-100.	0.5	0
121	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
122	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
123	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
124	Accidental High Dose Intrathecal Treatment - Late Result of A Patient. Turkish Journal of Haematology, 2020, 37, 64-65.	0.5	0
125	Hypoxic Ischemic Encephalopathy in Forensic Medicine. Adli TÄ±p BÃ¼lteni, 2021, 26, 205-209.	0.1	0
126	Anemic or not?. Turkish Journal of Pediatrics, 2003, 45, 329-34.	0.6	0

#	Article	IF	Citations
127	Thrombosis in the Vena Cava Inferior and Right Atrium in a Patient with Wilms Tumor. , 2023, 56, 535-536.		0