Aydan Ä°kincioÄullarÄ±

List of Publications by Year
in descending order

91 papers	3,827 citations	147801 31 h-index	128289 60 g-index
92 all docs	92 docs citations	92 times ranked	5556 citing authors

#	Article	IF	Citations
1	Scales of Magt1 Gene: Novel Mutations, Different Presentations. Iranian Journal of Allergy, Asthma and Immunology, 2022, 21, 92-97.	0.4	0
2	Single-Center Study of 72 Patients with Severe Combined Immunodeficiency: Clinical and Laboratory Features and Outcomes. Journal of Clinical Immunology, 2021, 41, 1563-1573.	3.8	3
3	Impaired respiratory burst contributes to infections in PKCÎ´-deficient patients. Journal of Experimental Medicine, 2021, 218, .	8.5	23
4	Expanding the Nude SCID/CID Phenotype Associated with FOXN1 Homozygous, Compound Heterozygous, or Heterozygous Mutations. Journal of Clinical Immunology, 2021, 41, 756-768.	3.8	13
5	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. Blood, 2020, 136, 2638-2655.	1.4	64
6	Does the Hyper IgM Phenotype Affect Prognosis in Ataxia Telangiectasia?. Asim, Allerji, Immunoloji, 2020, 18, 38-46.	0.0	1
7	An Unexpected Infection in Loss-of-function Mutations in STAT3: Malignant Alveolar Echinococcosis in Liver. Iranian Journal of Allergy, Asthma and Immunology, 2020, 19, 667-675.	0.4	2
8	Allogeneic hematopoietic stem cell and liver transplantation in a young girl with dedicator of cytokinesis 8 protein deficiency. Pediatric Transplantation, 2019, 23, e13545.	1.0	2
9	Hematopoietic stem cell transplantation for CD40 ligand deficiency: Results from an EBMT/ESID-IEWP-SCETIDE-PIDTC study. Journal of Allergy and Clinical Immunology, 2019, 143, 2238-2253.	2.9	60
10	Clinical Features and HSCT Outcome for SCID in Turkey. Journal of Clinical Immunology, 2019, 39, 316-323.	3.8	17
11	X-linked agammaglobulinemia (XLA): Phenotype, diagnosis, and therapeutic challenges around the world. World Allergy Organization Journal, 2019, 12, 100018.	3.5	83
12	Hematopoietic Stem Cell Transplantation as Treatment for Patients with DOCK8 Deficiency. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 848-855.	3.8	67
13	Recurrent Infections and Chronic Diarrhea. , 2019, , 453-458.		0
14	Treatment of severe forms of LPS-responsive beige-like anchor protein deficiency with allogeneic hematopoietic stem cell transplantation. Journal of Allergy and Clinical Immunology, 2018, 141, 770-775.e1.	2.9	52
15	Monogenic Diabetes Not Caused By Mutations in Mody Genes: A Very Heterogenous Group of Diabetes. Experimental and Clinical Endocrinology and Diabetes, 2018, 126, 612-618.	1.2	12
16	Human IFN-Î³ immunity to mycobacteria is governed by both IL-12 and IL-23. Science Immunology, 2018, 3, .	11.9	152
17	Outcome of treosulfanâ€based reducedâ€toxicity conditioning regimens for <scp>HSCT</scp> in highâ€risk patients with primary immune deficiencies. Pediatric Transplantation, 2018, 22, e13266.	1.0	5
18	Disruption of an antimycobacterial circuit between dendritic and helper T cells in human SPPL2a deficiency. Nature Immunology, 2018, 19, 973-985.	14.5	96

#	Article	IF	Citations
19	Both granulocytic and non-granulocytic blood cells are affected in patients with severe congenital neutropenia and their non-neutropenic family members: An evaluation of morphology, function and cell death. Turkish Journal of Haematology, 2018, 35, 229-259.	0.5	2
20	T Regulatory Cells in Children with Atopic Dermatitis. Journal of Ankara University Faculty of Medicine, 2018, 71, 118-122.	0.1	0
21	Combined immunodeficiency and Epstein-Barr virusâ€“induced B cell malignancy in humans with inherited CD70 deficiency. Journal of Experimental Medicine, 2017, 214, 91-106.	8.5	134
22	Multiple Presentations of LRBA Deficiency: a Single-Center Experience. Journal of Clinical Immunology, 2017, 37, 790-800.	3.8	64
23	Analysis of the recovery of CD247 expression in a PID patient: insights into the spontaneous repair of defective genes. Blood, 2017, 130, 1205-1208.	1.4	12
24	Primary T-cell immunodeficiency with functional revertant somatic mosaicism in CD247. Journal of Allergy and Clinical Immunology, 2017, 139, 347-349.e8.	2.9	17
25	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. Frontiers in Immunology, 2017, 8, 685.	4.8	50
26	Extended clinical and genetic spectrum associated with biallelic RTEL1 mutations. Blood Advances, 2016, 1, 36-46.	5.2	19
27	Neutrophilic Alveolitis and High Serum Pro-Brain Natriuretic Peptide Level may be Indicators of Pulmonary Functional Impairment in Connective Tissue Disorders. Archives of Rheumatology, 2016, 31, 14-23.	0.9	0
28	Unique and shared signaling pathways cooperate to regulate the differentiation of human CD4+ T cells into distinct effector subsets. Journal of Experimental Medicine, 2016, 213, 1589-1608.	8.5	77
29	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
30	HSCT for DOCK8 Deficiency - an International Study on 74 Patients. Biology of Blood and Marrow Transplantation, 2016, 22, S103-S104.	2.0	2
31	A Clinical Approach to a Child with Hypoalbuminemia and Lymphopenia. Journal of Clinical Immunology, 2016, 36, 370-373.	3.8	2
32	Lymphoma Secondary to Congenital and Acquired Immunodeficiency Syndromes at a Turkish Pediatric Oncology Center. Journal of Clinical Immunology, 2016, 36, 667-676.	3.8	21
33	Natural killer cell hyporesponsiveness and impaired development in a CD247-deficient patient. Journal of Allergy and Clinical Immunology, 2016, 137, 942-945.e4.	2.9	12
34	Immunologic alterations and efficacy of subcutaneous immunotherapy with Dermatophagoides pteronyssinus in monosensitized and polysensitized patients. Annals of Allergy, Asthma and Immunology, 2016, 116, 244-251.e2.	1.0	16
35	A Successful HSCT in a Girl with Novel LRBA Mutation with Refractory Celiac Disease. Journal of Clinical Immunology, 2016, 36, 8-11.	3.8	38
36	Regulatory T Cells and Vitamin D Status in Children with Chronic Autoimmune Thyroiditis. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2016, 8, 276-281.	0.9	20

#	Article	IF	Citations
37	Serum IL-13 levels at diagnosis and remission in children with malignant lymphoma. Turkish Journal of Pediatrics, 2016, 58, 246-253.	0.6	1
38	Inherited DOCK2 Deficiency in Patients with Early-Onset Invasive Infections. New England Journal of Medicine, 2015, 372, 2409-2422.	27.0	169
39	Î²2-Microglobulin deficiency causes a complex immunodeficiency of the innate and adaptive immune system. Journal of Allergy and Clinical Immunology, 2015, 136, 392-401.	2.9	66
40	Monogenic mutations differentially affect the quantity and quality of T follicular helper cells in patients with human primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2015, 136, 993-1006.e1.	2.9	181
41	Inherited IL-17RC deficiency in patients with chronic mucocutaneous candidiasis. Journal of Experimental Medicine, 2015, 212, 619-631.	8.5	162
42	The impact of donor age and sex on the nucleated cell count and CD34 count in healthy bone marrow donors. Pediatric Transplantation, 2015, 19, 385-390.	1.0	4
43	Toll-like receptors; integrate innate and adaptive immunity, promise innovative therapies. Journal of Pediatric Infectious Diseases, 2015, 01, 195-203.	0.2	0
44	Short-Term Preseasonal Immunotherapy: Is Early Clinical Efficacy Related to the Basophil Response?. International Archives of Allergy and Immunology, 2014, 164, 237-245.	2.1	27
45	Oral Lactoferrin to Prevent Nosocomial Sepsis and Necrotizing Enterocolitis of Premature Neonates and Effect on T-Regulatory Cells. American Journal of Perinatology, 2014, 31, 1111-1120.	1.4	86
46	The seroprevalence of Kaposiâ€²s sarcoma associated herpes virus and human herpes virus-6 in pediatric patients with cancer and healthy children in a Turkish pediatric oncology center. Indian Journal of Medical and Paediatric Oncology, 2014, 35, 221-225.	0.2	2
47	c.761C>T Mutation Linked Hyper IgM Syndrome Presenting with Hypertransaminasemia and Arthritis. Turkish Journal of Haematology, 2014, 31, 420-421.	0.5	0
48	Patients with Primary Immunodeficiencies in Pediatric Intensive Care Unit: Outcomes and Mortality-Related Risk Factors. Journal of Clinical Immunology, 2014, 34, 309-315.	3.8	19
49	BCG vaccination in patients with severe combined immunodeficiency: Complications, risks, and vaccination policies. Journal of Allergy and Clinical Immunology, 2014, 133, 1134-1141.	2.9	212
50	Clinical Features of Candidiasis in Patients With Inherited Interleukin 12 Receptor Î²1 Deficiency. Clinical Infectious Diseases, 2014, 58, 204-213.	5.8	98
51	Diagnosis of immunodeficiency caused by a purine nucleoside phosphorylase defect by using tandem mass spectrometry on dried blood spots. Journal of Allergy and Clinical Immunology, 2014, 134, 155-159.e3.	2.9	56
52	Early-onset inflammatory bowel disease and common variable immunodeficiencyâ€“like disease caused by IL-21 deficiency. Journal of Allergy and Clinical Immunology, 2014, 133, 1651-1659.e12.	2.9	124
53	Biallelic loss-of-function mutation in NIK causes a primary immunodeficiency with multifaceted aberrant lymphoid immunity. Nature Communications, 2014, 5, 5360.	12.8	116
54	Lymphoma Patients Secondary to Congenital and Acquired Immunodeficiency Syndroms in a Turkish Pediatric Oncology Center. Blood, 2014, 124, 5445-5445.	1.4	0

#	Article	IF	Citations
55	A Novel G6PC3 Gene Mutation in a Patient With Severe Congenital Neutropenia. Journal of Pediatric Hematology/Oncology, 2013, 35, e81-e83.	0.6	12
56	Selective IgA Deficiency: Clinical and Laboratory Features of 118 Children in Turkey. Journal of Clinical Immunology, 2012, 32, 961-966.	3.8	57
57	Selenium in the Prevention of Anthracycline-Induced Cardiac Toxicity in Children with Cancer. Journal of Oncology, 2012, 2012, 1-6.	1.3	4
58	Diagnosis of Immediate Hypersensitivity to Î²-Lactam Antibiotics Can Be Made Safely with Current Approaches. International Archives of Allergy and Immunology, 2012, 157, 311-317.	2.1	19
59	HLAâ€haploidentical transplantations for primary immunodeficiencies: A singleâ€center experience. Pediatric Transplantation, 2012, 16, 451-457.	1.0	7
60	Reliability of Basophil Activation Test Using CD203c Expression in Diagnosis of Pollen Allergy. American Journal of Rhinology and Allergy, 2011, 25, e225-e231.	2.0	25
61	The Effect of Mode of Delivery on T Regulatory (Treg) Cells of Cord Blood. Indian Journal of Pediatrics, 2011, 78, 1234-1238.	0.8	13
62	Two Patients with Partial DiGeorge Syndrome Presenting with Attention Disorder and Learning Difficulties - Case Report. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2011, 3, 95-97.	0.9	9
63	Revisiting Human IL-12RÎ²1 Deficiency. Medicine (United States), 2010, 89, 381-402.	1.0	367
64	Purine nucleoside phosphorylase deficiency with fatal course in two sisters. European Journal of Pediatrics, 2010, 169, 311-314.	2.7	22
65	Reply to correspondence letter by Luis Ignacio Gonzalez-Granado. European Journal of Pediatrics, 2010, 169, 519-519.	2.7	0
66	Eponym. European Journal of Pediatrics, 2010, 169, 657-660.	2.7	9
67	Kostmann disease with developmental delay in three patients. European Journal of Pediatrics, 2010, 169, 759-762.	2.7	10
68	Late onset hemorrhagic cystitis in a hematopoietic stem cell recipient: Treatment with intravesical hyaluronic acid. Pediatric Transplantation, 2010, 14, E79-E82.	1.0	12
69	DOES SERUM SOLUBLE VASCULAR ENDOTHELIAL GROWTH FACTOR LEVELS HAVE DIFFERENT IMPORTANCE IN PEDIATRIC ACUTE LEUKEMIA AND MALIGNANT LYMPHOMA PATIENTS?. Pediatric Hematology and Oncology, 2010, 27, 503-516.	0.8	9
70	Asymptomatic catheter related Rhizobium radiobacter infection in a haploidentical hemapoetic stem cell recipient. Journal of Infection in Developing Countries, 2010, 4, 530-532.	1.2	6
71	Selective Immunoglobulin M Deficiency Presenting with Recurrent Impetigo: A Case Report and Review of the Literature. International Archives of Allergy and Immunology, 2009, 149, 283-288.	2.1	18
72	Upregulation of CD63 or CD203c Alone or in Combination Is Not Sensitive in the Diagnosis of Nonsteroidal Anti-Inflammatory Drug Intolerance. International Archives of Allergy and Immunology, 2009, 150, 261-270.	2.1	39

#	ARTICLE	IF	CITATIONS
73	Defects along the TH17 differentiation pathway underlie genetically distinct forms of the hyper IgE syndrome. Journal of Allergy and Clinical Immunology, 2009, 124, 342-348.e5.	2.9	94
74	G-CSF-mobilized haploidentical peripheral blood stem cell transplantation in children with poor prognostic nonmalignant disorders. American Journal of Hematology, 2008, 83, 133-136.	4.1	9
75	An unconditioned bone marrow transplantation in a child with purine nucleoside phosphorylase deficiency and its unique complication. Pediatric Transplantation, 2008, 12, 479-482.	1.0	26
76	Long-term outcome following hematopoietic stem-cell transplantation in Wiskott-Aldrich syndrome: collaborative study of the European Society for Immunodeficiencies and European Group for Blood and Marrow Transplantation. Blood, 2008, 111, 439-445.	1.4	216
77	A novel mutation for TAP deficiency and its possible association with Toxoplasmosis. Parasitology International, 2006, 55, 219-222.	1.3	19
78	Tonsillectomy and the immune system. International Journal of Pediatric Otorhinolaryngology, 2006, 70, 175-176.	1.0	0
79	Stable mixed chimerism after hematopoietic stem cell transplantation in Wiskott-Aldrich syndrome. Pediatric Transplantation, 2006, 10, 395-399.	1.0	5
80	Complete deficiency of the IL-12 receptor Î²1Âchain: three unrelated Turkish children with unusual clinical features. European Journal of Pediatrics, 2006, 165, 415-417.	2.7	19
81	Meningococccal meningitis and complement component 6 deficiency associated with oculocutaneous albinism. European Journal of Pediatrics, 2005, 164, 177-179.	2.7	7
82	Granulocyte Transfusions in Children With Chronic Granulomatous Disease and Invasive Aspergillosis. Therapeutic Apheresis and Dialysis, 2005, 9, 137-141.	0.9	41
83	Unusual cause of respiratory distress: Chilaiditi syndrome. Pediatrics International, 2004, 46, 188-190.	0.5	13
84	Peripheral blood lymphocyte subsets in healthy Turkish children. Turkish Journal of Pediatrics, 2004, 46, 125-30.	0.6	55
85	Evaluation of Nutritional Status and Factors Related to Malnutrition in Children on CAPD. Peritoneal Dialysis International, 2003, 23, 557-562.	2.3	13
86	RECURRENT ARTERIAL THROMBOSIS IN A CHILD: Primary Antiphospholipid Antibody Syndrome. Pediatric Hematology and Oncology, 2002, 19, 59-66.	0.8	12
87	Is immune system influenced by adenotonsillectomy in children?. International Journal of Pediatric Otorhinolaryngology, 2002, 66, 251-257.	1.0	41
88	Serum levels and differential expression of CD44 in childhood leukemia and malignant lymphoma: Correlation with prognostic criteria and survival. Pediatrics International, 2001, 43, 354-360.	0.5	12
89	Zinc status and cytokine profile in pediatric Hodgkin's disease. Journal of Trace Elements in Experimental Medicine, 2001, 14, 25-30.	0.8	2
90	Polyglandular autoimmune syndrome accompanied by Munchausen syndrome. Pediatrics International, 2000, 42, 386-388.	0.5	5

#	Article	IF	Citations
91	NATURAL KILLER CELL NUMBERS AND CYTOTOXIC ACTIVITY IN PEDIATRIC HODGKIN DISEASE. Pediatric Hematology and Oncology, 2000, 17, 133-139.	0.8	2