Deniz Ãagdas Ayvaz

List of Publications by Year
in descending order

124 papers	2,638 citations	279701 23 h-index	214721 47 g-index
137 all docs	137 docs citations	137 times ranked	4100 citing authors

#	Article	IF	Citations
1	Clinical, Laboratory Features and Clinical Courses of Patients with Wiskott Aldrich Syndrome and Xâ€“linked Thrombocytopeniaâ€“A single center study. Immunological Investigations, 2022, 51, 1272-1283.	1.0	0
2	TMC8 mutation in a Turkish family with epidermodysplasia verruciformis including laryngeal papilloma and recurrent skin carcinoma. Journal of Cosmetic Dermatology, 2022, 21, 2263-2267.	0.8	1
3	Nutritional status of children with primary immunodeficiency: A single center experience. Pediatrics International, 2022, 64, .	0.2	0
4	Long Term Follow-Up of the Patients with Severe Combined Immunodeficiency After Hematopoietic Stem Cell Transplantation: A Single-Center Study. Immunological Investigations, 2022, 51, 739-747.	1.0	4
5	STK4 deficiency and EBV-associated lymphoproliferative disorders, emphasis on histomorphology, and review of literature. Virchows Archiv Fur Pathologische Anatomie Und Physiologie Und Fur Klinische Medizin, 2022, 480, 393-401.	1.4	5
6	Multicentric Castleman disease in a DOCK8â€deficient patient with Orf virus infection. Pediatric Allergy and Immunology, 2022, 33, .	1.1	3
7	Evaluation of the peripheral blood T and B cell subsets and ‹scp›IRF‹/scp›â€7 variants in adult patients with severe influenza virus infection. Health Science Reports, 2022, 5, e492.	0.6	1
8	Combined immunodeficiency due to purine nucleoside phosphorylase deficiency: Outcome of three patients. European Journal of Medical Genetics, 2022, 65, 104428.	0.7	10
9	Evaluation of Health Status and Quality of Life in Patients Using Intravenous and Subcutaneous Forms of Immunoglobulin Replacement. Journal of Ankara University Faculty of Medicine, 2022, 75, 77-83.	0.0	0
10	First allogeneic hematopoietic stem cell transplantation in RASGRP1 deficiency: long-term follow-up. Bone Marrow Transplantation, 2022, , .	1.3	0
11	GIMAP6 regulates autophagy, immune competence, and inflammation in mice and humans. Journal of Experimental Medicine, 2022, 219, .	4.2	4
12	Antimycobacterial prophylaxis regarding Bacillus Calmette-GuÃ©rin -associated complications in children with primary immunodeficiency. Respiratory Medicine, 2022, , 106919.	1.3	1
13	Frequency of HLA Class I and Class II Alleles in Patients with CVID from Turkey. Immunological Investigations, 2021, 50, 363-371.	1.0	0
14	Evaluation of periodontal status and cytokine/chemokine profile of GCF in patients with severe congenital neutropenia. Odontology / the Society of the Nippon Dental University, 2021, 109, 474-482.	0.9	7
15	Flow Cytometric Analysis of T Cell VÎ² Repertoire in Common Variable Immunodeficiency Patients with TACI Mutations. Turkish Journal of Immunology, 2021, , .	0.1	0
16	A Patient With AIRE Mutation Who Presented With Severe Diarrhea and Lung Abscess. Pediatric Infectious Disease Journal, 2021, 40, 66-69.	1.1	3
17	Characteristics of patients with C1 esterase inhibitor deficiency: a single center study. European Annals of Allergy and Clinical Immunology, 2021, 53, 75.	0.4	3
18	Living the SARS-CoV-2 pandemic in Turkey. Nature Immunology, 2021, 22, 260-260.	7.0	2

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19	Clinical and Immunological Characteristics of 63 Patients with Chronic Granulomatous Disease: Hacettepe Experience. Journal of Clinical Immunology, 2021, 41, 992-1003.	2.0	8
20	Selective IgM deficiency: Followâ€up and outcome. Pediatric Allergy and Immunology, 2021, 32, 1327-1334.	1.1	8
21	Convalescent plasma and hyperimmune globulin therapy in COVID-19. Expert Review of Clinical Immunology, 2021, 17, 309-315.	1.3	10
22	Genomic Spectrum and Phenotypic Heterogeneity of Human IL-21 Receptor Deficiency. Journal of Clinical Immunology, 2021, 41, 1272-1290.	2.0	25
23	Expression of HLA class I and class II genes in patients with multiple skin warts. Experimental Dermatology, 2021, 30, 1642-1649.	1.4	1
24	Diversity in Serine/Threonine Protein Kinase-4 Deficiency and Review of the Literature. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 3752-3766.e4.	2.0	13
25	Inherited PD-1 deficiency underlies tuberculosis and autoimmunity in a child. Nature Medicine, 2021, 27, 1646-1654.	15.2	65
26	Differential diagnosis of primary immunodeficiency in patients with BCGitis and BCGosis: A singleâ€centre study. Scandinavian Journal of Immunology, 2021, 94, e13084.	1.3	8
27	COVID-19 in Patients with Primary Immunodeficiency. Journal of Clinical Immunology, 2021, 41, 1515-1522.	2.0	38
28	Clinical and laboratory findings in patients with leukocyte adhesion deficiency type I: A multicenter study in Turkey. Clinical and Experimental Immunology, 2021, 206, 47-55.	1.1	10
29	Respiratory system findings in pediatric patients with primary immunodeficiency. Pediatric Pulmonology, 2021, 56, 4011-4019.	1.0	2
30	Hematologically important mutations: X-linked chronic granulomatous disease (fourth update). Blood Cells, Molecules, and Diseases, 2021, 90, 102587.	0.6	22
31	Hematologically important mutations: The autosomal forms of chronic granulomatous disease (third) Tj ETQq1 1 0.784314 rgBT /Ove	0.6	22
32	A RARE INVOLVEMENT OF CENTRAL NERVOUS SYSTEM INVOLVEMENT DUE TO CTLA-4 GENE DEFECT. Noropsikiyatri Arsivi, 2021, , .	0.2	0
33	In case of recurrent wheezing and bronchiolitis: Think again, it may be a primary immunodeficiency. Asian Pacific Journal of Allergy and Immunology, 2021, , .	0.2	1
34	Cytokine profile in serum and gingival crevicular fluid of children with inflammatory bowel disease: A caseâ€control study. Journal of Periodontology, 2021, , .	1.7	2
35	Maternofetal Transfusion, Maternal Chimerism, and Maternal Engraftment: A Mystery in Health and Disease. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 4154-4155.	2.0	0
36	Hematopoietic stem cell transplantation complicated with ebv associated hemophagocytic lymphohistiocytosis in a patient with dock2 deficiency. Turkish Journal of Pediatrics, 2021, 63, 1072.	0.3	5

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37	A Monogenic Disease with a Variety of Phenotypes: Deficiency of Adenosine Deaminase 2. Journal of Rheumatology, 2020, 47, 117-125.	1.0	65
38	A clinical score to guide in decision making for monogenic type I IFNopathies. Pediatric Research, 2020, 87, 745-752.	1.1	16
39	Recurrent Demyelinating Episodes as Sole Manifestation of Inherited CD59 Deficiency. Neuropediatrics, 2020, 51, 206-210.	0.3	6
40	Adenosine Deaminase Type II Deficiency: Severe Chronic Neutropenia, Lymphoid Infiltration in Bone Marrow, and Inflammatory Features. Immunological Investigations, 2020, , 1-9.	1.0	3
41	Hemophagocytic Lymphohistiocytosis in Patients With Primary Immunodeficiency. Journal of Pediatric Hematology/Oncology, 2020, 42, e434-e439.	0.3	7
42	Recurrent skin abscesses in a female Xâ€linked chronic granulomatous disease carrier. Journal of Cosmetic Dermatology, 2020, 19, 1810-1812.	0.8	1
43	Impact of mannoseâ€binding lectin 2 gene polymorphisms on disease severity in noncystic fibrosis bronchiectasis in children. Pediatric Pulmonology, 2020, 55, 1190-1198.	1.0	5
44	Autoinflammation in addition to combined immunodeficiency: SLC29A3 gene defect. Molecular Immunology, 2020, 121, 28-37.	1.0	18
45	A rare form of congenital neutropenia: VPS45 deficiency. Scandinavian Journal of Immunology, 2020, 91, e12871.	1.3	5
46	Lymphocyte Subgroups and KREC Numbers in Common Variable Immunodeficiency: A Single Center Study. Journal of Clinical Immunology, 2020, 40, 494-502.	2.0	2
47	Mutational landscape of severe combined immunodeficiency patients from Turkey. International Journal of Immunogenetics, 2020, 47, 529-538.	0.8	14
48	Retroperitoneal Abscess in Severe Combined Immunodeficiency Probably Due to BCG Vaccine. Asim, Allerji, Immunoloji, 2020, 18, 98-101.	0.2	1
49	Primary Immunodeficiency Disorders in Children with Non Cystic Fibrosis Bronchiectasis. European Annals of Allergy and Clinical Immunology, 2020, 52, 271.	0.4	6
50	Three patients with glucose-6 phosphatase catalytic subunit 3 deficiency. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 957-961.	0.4	0
51	A Spectrum of Clinical Findings from ALPS to CVID: Several Novel LRBA Defects. Journal of Clinical Immunology, 2019, 39, 726-738.	2.0	45
52	ADA2 deficiency in a patient with Noonan syndromeâ€like disorder with loose anagen hair: The coâ€occurrence of two rare syndromes. American Journal of Medical Genetics, Part A, 2019, 179, 2474-2480.	0.7	5
53	Bone marrow transplantation with Favorable outcome in three patients with LPS-responsive beige-like anchor (LRBA) deficiency. Clinical Immunology, 2019, 203, 162-165.	1.4	7
54	Clinical Features and HSCT Outcome for SCID in Turkey. Journal of Clinical Immunology, 2019, 39, 316-323.	2.0	17

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55	AB0574â€…A MONOGENIC DISEASE WITH WIDE RANGE OF SYMPTOMS: DEFICIENCY OF ADENOSINE DEAMINASE 2. , 2019, , .		1
56	Selective loss of function variants in IL6ST cause Hyper-IgE syndrome with distinct impairments of T-cell phenotype and function. Haematologica, 2019, 104, 609-621.	1.7	74
57	Tumor necrosis factor Î±â€308 G/A and interleukin 1 Î²â€511 C/T gene polymorphisms in patients with scarring acne. Journal of Cosmetic Dermatology, 2019, 18, 395-400.	0.8	10
58	Primer Ä°mmÃ¼n Yetersizliklerde Western Blot YÃ¶nteminin Ã–nemi: Ä°ki Aile Olgusundan Ã–rnekler. Turkish Journal of Immunology, 2019, 7, .	0.1	0
59	Axillary Lymphadenopathy: Only Presentation in an Infant Diagnosed with Chronic Granulomatous Disease. Kafkas Journal of Medical Sciences, 2019, 9, 217-220.	0.1	0
60	The Association Between Vitamin D Levels and Infections in Patients with Primary Immunodeficiency. Turkish Journal of Immunology, 2019, 7, .	0.1	0
61	Clinical and genetic features of the patients with Xâ€linked agammaglobulinemia from Turkey: Singleâ€centre experience. Scandinavian Journal of Immunology, 2018, 87, e12647.	1.3	7
62	Neurologic Involvement in Primary Immunodeficiency Disorders. Journal of Child Neurology, 2018, 33, 320-328.	0.7	12
63	Type I IFNâ€“related NETosis in ataxia telangiectasia and Artemis deficiency. Journal of Allergy and Clinical Immunology, 2018, 142, 246-257.	1.5	47
64	A young girl with severe cerebral fungal infection due to card 9 deficiency. Clinical Immunology, 2018, 191, 21-26.	1.4	27
65	Successful hematopoietic stem cell transplantation after myeloablative conditioning in three patients with dedicator of cytokinesis 8 deficiency (DOCK8) related Hyper IgE syndrome. Bone Marrow Transplantation, 2018, 53, 339-343.	1.3	8
66	A Rare Cause of Secondary Immunodeficiency: Generalized Lymphatic Anomaly. Journal of Pediatric Hematology/Oncology, 2018, 40, 248-251.	0.3	0
67	Hypomorphic RAG1 defect in a child presented with pulmonary hemorrhage and digital necrosis. Clinical Immunology, 2018, 187, 92-94.	1.4	6
68	B lymphocyte subsets and outcomes in patients with an initial diagnosis of transient hypogammaglobulinemia of infancy. Scandinavian Journal of Immunology, 2018, 88, e12709.	1.3	10
69	A Novel Homozygous Mutation With Different Clinical Presentations in 2 IRAK-4â€“Deficient Siblings: First Case With Recurrent Salmonellosis and Non-Hodgkin Lymphoma. Journal of Investigational Allergology and Clinical Immunology, 2018, 28, 271-273.	0.6	6
70	ADA Deficiency: Evaluation of the Clinical and Laboratory Features and the Outcome. Journal of Clinical Immunology, 2018, 38, 484-493.	2.0	26
71	Two siblings with PRKDC defect who presented with cutaneous granulomas and review of the literature. Clinical Immunology, 2018, 197, 1-5.	1.4	18
72	Advantage of the subcutaneous immunoglobulin replacement therapy in primary immunodeficient patients with or without secondary protein loss. Turkish Journal of Pediatrics, 2018, 60, 270-276.	0.3	2

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73	Investigation of Genetic Defects in Severe Combined Immunodeficiency Patients from Turkey by Targeted Sequencing. Scandinavian Journal of Immunology, 2017, 85, 227-234.	1.3	30
74	A novel mutation in TAP1 gene leading to MHC class I deficiency: Report of two cases and review of the literature. Clinical Immunology, 2017, 178, 74-78.	1.4	31
75	Acute myeloid leukemia in a child with dedicator of cytokinesis 8 (DOCK8) deficiency. Pediatric Blood and Cancer, 2017, 64, e26695.	0.8	6
76	Diagnosis of Interstitial Lung Disease Caused by Possible Hypersensitivity Pneumonitis in a Child: Think CGD. Journal of Clinical Immunology, 2017, 37, 269-272.	2.0	9
77	Low T Cell Numbers Resembling Tâ˜B+ SCID in a Patient with Wiskottâ€“Aldrich Syndrome and the Outcome of Two Hematopoietic Stem Cell Transplantations. Journal of Clinical Immunology, 2017, 37, 18-21.	2.0	0
78	Hematopoietic stem cell transplantation in children with Griscelli syndrome: A singleâ€¢center experience. Pediatric Transplantation, 2017, 21, e13040.	0.5	12
79	Progressive multifocal leukoencephalopathy in a patient with lymphoma and presumptive hyper IgE syndrome. Journal of NeuroVirology, 2017, 23, 632-636.	1.0	2
80	Course of IL-2-inducible T-cell kinase deficiency in a family: lymphomatoid granulomatosis, lymphoma and allogeneic bone marrow transplantation in one sibling; and death in the other. Bone Marrow Transplantation, 2017, 52, 126-129.	1.3	21
81	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. Frontiers in Immunology, 2017, 8, 685.	2.2	50
82	Defective pneumococcal antibody response in patients with recurrent respiratory tract infections. Turkish Journal of Pediatrics, 2017, 59, 555-560.	0.3	1
83	Detection of parasites in children with chronic diarrhea. Pediatrics International, 2016, 58, 531-533.	0.2	12
84	Heterozygous STAT1 gain-of-function mutations underlie an unexpectedly broad clinical phenotype. Blood, 2016, 127, 3154-3164.	0.6	465
85	RASGRP1 deficiency causes immunodeficiency with impaired cytoskeletal dynamics. Nature Immunology, 2016, 17, 1352-1360.	7.0	115
86	An infant with ZAP-70 deficiency with disseminated mycobacterial disease. Journal of Clinical Immunology, 2016, 36, 103-106.	2.0	11
87	Successful outcome with second hematopoietic stem cell transplantation in a patient with IL-10R deficiency. Bone Marrow Transplantation, 2016, 51, 615-616.	1.3	5
88	Recurrent viral infections associated with a homozygous CORO1A mutation that disrupts oligomerization and cytoskeletal association. Journal of Allergy and Clinical Immunology, 2016, 137, 879-888.e2.	1.5	41
89	Clinical and genetic features of IL12Rb1 deficiency: Single center experience of 18 patients. Turkish Journal of Pediatrics, 2016, 58, 356-361.	0.3	15
90	CVID Associated with Systemic Amyloidosis. Case Reports in Immunology, 2015, 2015, 1-4.	0.2	8

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91	Combined immunodeficiency with CD4 lymphopenia and sclerosing cholangitis caused by a novel loss-of-function mutation affecting IL21R. Haematologica, 2015, 100, e216-e219.	1.7	46
92	DOCK8 Deficiency: Clinical and Immunological Phenotype and Treatment Options - a Review of 136 Patients. Journal of Clinical Immunology, 2015, 35, 189-198.	2.0	284
93	STK4 (MST1) deficiency in two siblings with autoimmune cytopenias: A novel mutation. Clinical Immunology, 2015, 161, 316-323.	1.4	73
94	Diagnosis: Melanoderma after Hematopoietic Stem Cell Transplantation. Turkish Journal of Haematology, 2015, 32, 378-379.	0.2	1
95	65 Year Old Patient with Common Variable Immunodeficiency: 3 Years Followup and Development of Lung Adeno Carcinoma, is it a Coincidence?. Journal of Pulmonary & Respiratory Medicine, 2015, 05, .	0.1	0
96	Polymorphisms in FAS and CASP8 genes may contribute to the development of ALPS phenotype: a study in 25 patients with probable ALPS. Turkish Journal of Pediatrics, 2015, 57, 141-5.	0.3	4
97	A Novel Mutation in Leukocyte Adhesion Deficiency Type II/CDGIIc. Journal of Clinical Immunology, 2014, 34, 1009-1014.	2.0	21
98	Progressive Neurodegenerative Syndrome in a Patient with X-Linked Agammaglobulinemia Receiving Intravenous Immunoglobulin Therapy. Cognitive and Behavioral Neurology, 2014, 27, 155-159.	0.5	9
99	Metamizole-Induced Bicytopenia Reversed by G-CSF and IVIG Treatment in a Child. Pediatric Hematology and Oncology, 2014, 31, 117-119.	0.3	5
100	Coronin-1A Oligomerization Is Critical For Host Defense Against Viral Pathogens. Journal of Allergy and Clinical Immunology, 2014, 133, AB94.	1.5	2
101	Successful treatment of severe myasthenia gravis developed after allogeneic hematopoietic stem cell transplantation with plasma exchange and rituximab. Pediatric Blood and Cancer, 2014, 61, 928-930.	0.8	14
102	Identification of ITK deficiency as a novel genetic cause of idiopathic CD4+ T-cell lymphopenia. Blood, 2014, 124, 655-657.	0.6	51
103	Griscelli syndrome type 3â€like phenotype with <scp>MYO</scp>â€5A exonâ€F deletion. Pediatric Allergy and Immunology, 2014, 25, 817-819.	1.1	7
104	IL-12RÎ²1 Deficiency: Mutation Update and Description of theIL12RB1Variation Database. Human Mutation, 2013, 34, 1329-1339.	1.1	81
105	Atypical combined immunodeficiency due to Artemis defect: A case presenting as hyperimmunoglobulin M syndrome and with LGLL. Molecular Immunology, 2013, 56, 354-357.	1.0	21
106	A Case ofDOCK8Deficient Hyper-IgE Syndrome Presenting Primarily With Eczema, Food Allergy, and Asthma. Pediatric, Allergy, Immunology, and Pulmonology, 2013, 26, 48-51.	0.3	0
107	The Role of IL-12RÃŸ1 Expression Analysis Using Flow Cytometry in the Diagnosis of IL-12RÃŸ1 Deficiency. Turkish Journal of Immunology, 2013, 1, 1-4.	0.1	0
108	Apoptosis in Autoimmune Lymphoproliferative Syndrome Suspected Patients with Clinical and Laboratory Findings. Turkish Journal of Immunology, 2013, 1, 5-12.	0.1	0

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109	Griscelli syndrome types 1 and 3: analysis of four new cases and long-term evaluation of previously diagnosed patients. European Journal of Pediatrics, 2012, 171, 1527-1531.	1.3	31
110	Additional Diverse Findings Expand the Clinical Presentation of DOCK8 Deficiency. Journal of Clinical Immunology, 2012, 32, 698-708.	2.0	84
111	Two SCID cases with Cernunnosâ€XLF deficiency successfully treated by hematopoietic stem cell transplantation. Pediatric Transplantation, 2012, 16, E167-71.	0.5	22
112	SaÄŸlÄ±k meslek lisesi Ã¶ÄŸrencilerinin beslenme alÄ±ÅŸkanlÄ±klarÄ±nÄ±n, beslenme bilgi dÃ¼zeylerinin ve vÃ¼cut bileÅŸimlerinin deÄŸerlendirilmesi. Turk Pediatri Arsivi, 2012, 47, 181-188.	0.9	3
113	Congenital Left Ventricular Apical Diverticulum. Echocardiography, 2010, 27, 476-477.	0.3	1
114	Swyer-James syndrome or destroyed lung?/Coronary artery bypass in a patient with Swyer-James syndrome due to pulmonary tuberculosis. Anatolian Journal of Cardiology, 2010, 10, 383-384.	0.4	0
115	Hepatopulmonary syndrome associated with Budd-Chiari syndrome. Anatolian Journal of Cardiology, 2010, 10, 286-288.	0.4	1
116	Axenfeldâ€“Rieger syndrome and pseudotruncus arteriosus. International Journal of Cardiology, 2008, 126, e4-e7.	0.8	4
117	Glucocorticoid-induced Diabetic Ketoacidosis in Acute Rheumatic Fever. Journal of Cardiovascular Pharmacology and Therapeutics, 2008, 13, 298-300.	1.0	26
118	Treatment of massive cardiac thrombi in a patient with protein C and protein S deficiency. Blood Coagulation and Fibrinolysis, 2007, 18, 699-702.	0.5	13
119	Left main coronary artery and aortic root compression associated with atrial septal defect and pulmonary hypertension. International Journal of Cardiology, 2007, 118, e41-e43.	0.8	10
120	Familial Mediterranean fever and mesangial proliferative glomerulonephritis: report of a case and review of the literature. Pediatric Nephrology, 2005, 20, 1352-1354.	0.9	11
121	Right-sided Duane retraction syndrome associated with multiple malformations. Pediatrics International, 2003, 45, 577-579.	0.2	0
122	Does Retirement Hurt Weil-Being? Factors Influencing Self-Esteem and Depression Among Retires and Workers. Gerontologist, The, 1996, 36, 649-656.	2.3	131
123	Premature ventricular contractions in normal children. Journal of Pediatrics, 1978, 92, 36-38.	0.9	85
124	Flow Cytometry is a Reliable Tool in the Diagnosis of STK4 Deficiency. Asim, Allerji, Immunoloji, 0, , .	0.2	2