

# Figen Dogu

## List of Publications by Year in descending order

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107  
papers

4,131  
citations

147801

31  
h-index

123424

61  
g-index

109  
all docs

109  
docs citations

109  
times ranked

5675  
citing authors

#	ARTICLE	IF	CITATIONS
1	Hematopoietic cell transplantation in severe combined immunodeficiency: The SCETIDE 2006-2014 European cohort. <i>Journal of Allergy and Clinical Immunology</i> , 2022, 149, 1744-1754.e8.	2.9	51
2	Scales of Magt1 Gene: Novel Mutations, Different Presentations. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2022, 21, 92-97.	0.4	0
3	Single-Center Study of 72 Patients with Severe Combined Immunodeficiency: Clinical and Laboratory Features and Outcomes. <i>Journal of Clinical Immunology</i> , 2021, 41, 1563-1573.	3.8	3
4	Inherited PD-1 deficiency underlies tuberculosis and autoimmunity in a child. <i>Nature Medicine</i> , 2021, 27, 1646-1654.	30.7	65
5	Impaired respiratory burst contributes to infections in PKC $\delta$ -deficient patients. <i>Journal of Experimental Medicine</i> , 2021, 218, .	8.5	23
6	Expanding the Nude SCID/CID Phenotype Associated with FOXP1 Homozygous, Compound Heterozygous, or Heterozygous Mutations. <i>Journal of Clinical Immunology</i> , 2021, 41, 756-768.	3.8	13
7	Long-term outcome of LRBA deficiency in 76 patients after various treatment modalities as evaluated by the immune deficiency and dysregulation activity (IDDA) score. <i>Journal of Allergy and Clinical Immunology</i> , 2020, 145, 1452-1463.	2.9	112
8	Dominant-negative mutations in human <i>IL6ST</i> underlie hyper-IgE syndrome. <i>Journal of Experimental Medicine</i> , 2020, 217, .	8.5	64
9	Clinical, immunological features and follow up of 20 patients with dedicator of cytokinesis 8 (DOCK8) deficiency. <i>Pediatric Allergy and Immunology</i> , 2020, 31, 515-527.	2.6	17
10	Extended clinical and immunological phenotype and transplant outcome in CD27 and CD70 deficiency. <i>Blood</i> , 2020, 136, 2638-2655.	1.4	64
11	Clinical Features and Outcomes of 23 Patients with Wiskott- Aldrich Syndrome: A Single-Center Experience. <i>Turkish Journal of Haematology</i> , 2020, 37, 271-281.	0.5	7
12	An Unexpected Infection in Loss-of-function Mutations in STAT3: Malignant Alveolar Echinococcosis in Liver. <i>Iranian Journal of Allergy, Asthma and Immunology</i> , 2020, 19, 667-675.	0.4	2
13	Underlying Diseases and Causative Microorganisms of Recurrent Pneumonia in Children: A 13-Year Study in a University Hospital. <i>Journal of Tropical Pediatrics</i> , 2019, 65, 224-230.	1.5	2
14	PROMIS1: AAT-cell receptor 1 signature associated with immunodeficiencies caused by V(D)J recombination defects. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 325-334.e2.	2.9	43
15	Allogeneic hematopoietic stem cell and liver transplantation in a young girl with dedicator of cytokinesis 8 protein deficiency. <i>Pediatric Transplantation</i> , 2019, 23, e13545.	1.0	2
16	Hematopoietic stem cell transplantation for CD40 ligand deficiency: Results from an EBMT/ESID-IEWP-SCETIDE-PIDTC study. <i>Journal of Allergy and Clinical Immunology</i> , 2019, 143, 2238-2253.	2.9	60
17	Clinical Features and HSCT Outcome for SCID in Turkey. <i>Journal of Clinical Immunology</i> , 2019, 39, 316-323.	3.8	17
18	X-linked agammaglobulinemia (XLA): Phenotype, diagnosis, and therapeutic challenges around the world. <i>World Allergy Organization Journal</i> , 2019, 12, 100018.	3.5	83

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19	Hematopoietic Stem Cell Transplantation as Treatment for Patients with DOCK8 Deficiency. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2019, 7, 848-855.	3.8	67
20	Treatment of severe forms of LPS-responsive beige-like anchor protein deficiency with allogeneic hematopoietic stem cell transplantation. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 141, 770-775.e1.	2.9	52
21	Monogenic Diabetes Not Caused By Mutations in Mody Genes: A Very Heterogenous Group of Diabetes. <i>Experimental and Clinical Endocrinology and Diabetes</i> , 2018, 126, 612-618.	1.2	12
22	Human IFN- $\beta$ immunity to mycobacteria is governed by both IL-12 and IL-23. <i>Science Immunology</i> , 2018, 3, .	11.9	152
23	Disruption of an antimycobacterial circuit between dendritic and helper T cells in human SPPL2a deficiency. <i>Nature Immunology</i> , 2018, 19, 973-985.	14.5	96
24	T Regulatory Cells in Children with Atopic Dermatitis. <i>Journal of Ankara University Faculty of Medicine</i> , 2018, 71, 118-122.	0.1	0
25	Expanding the mutation spectrum in <scp>ICF</scp> syndrome: Evidence for a gender bias in <scp>ICF2</scp>. <i>Clinical Genetics</i> , 2017, 92, 380-387.	2.0	28
26	Combined immunodeficiency and Epstein-Barr virus-induced B cell malignancy in humans with inherited CD70 deficiency. <i>Journal of Experimental Medicine</i> , 2017, 214, 91-106.	8.5	134
27	Multiple Presentations of LRBA Deficiency: a Single-Center Experience. <i>Journal of Clinical Immunology</i> , 2017, 37, 790-800.	3.8	64
28	P271â€œ...Underlying disease and causative microorganisms of recurrent pneumonia in children: 13-year study in a university hospital. , 2017, , .		0
29	Primary T-cell immunodeficiency with functional revertant somatic mosaicism in CD247. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 139, 347-349.e8.	2.9	17
30	P274â€œ...An unexpected disease in an infant with pancytopenia and pulmonary abscess: glycogen storage disease type 1b. , 2017, , .		0
31	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. <i>Frontiers in Immunology</i> , 2017, 8, 685.	4.8	50
32	Extended clinical and genetic spectrum associated with biallelic RTEL1 mutations. <i>Blood Advances</i> , 2016, 1, 36-46.	5.2	19
33	Genetic, immunological, and clinical features of patients with bacterial and fungal infections due to inherited IL-17RA deficiency. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, E8277-E8285.	7.1	137
34	HSCT for DOCK8 Deficiency - an International Study on 74 Patients. <i>Biology of Blood and Marrow Transplantation</i> , 2016, 22, S103-S104.	2.0	2
35	A Clinical Approach to a Child with Hypoalbuminemia and Lymphopenia. <i>Journal of Clinical Immunology</i> , 2016, 36, 370-373.	3.8	2
36	Lymphoma Secondary to Congenital and Acquired Immunodeficiency Syndromes at a Turkish Pediatric Oncology Center. <i>Journal of Clinical Immunology</i> , 2016, 36, 667-676.	3.8	21

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37	Economic Burden of Primary Immunodeficiency (PIDD) In Turkey. Value in Health, 2016, 19, A585.	0.3	1
38	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
39	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
40	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
41	Serum IL-13 levels at diagnosis and remission in children with malignant lymphoma. Turkish Journal of Pediatrics, 2016, 58, 246-253.	0.6	1
42	ITK Deficiency: How can EBV be Treated Before Lymphoma?. Pediatric Blood and Cancer, 2015, 62, 2247-2248.	1.5	27
43	Inherited and acquired immunodeficiencies underlying tuberculosis in childhood. Immunological Reviews, 2015, 264, 103-120.	6.0	180
44	Inherited DOCK2 Deficiency in Patients with Early-Onset Invasive Infections. New England Journal of Medicine, 2015, 372, 2409-2422.	27.0	169
45	Inherited IL-17RC deficiency in patients with chronic mucocutaneous candidiasis. Journal of Experimental Medicine, 2015, 212, 619-631.	8.5	162
46	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
47	Intractable colitis associated with chronic granulomatous disease in a young girl. Turkish Journal of Pediatrics, 2015, 57, 189-91.	0.6	0
48	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
49	The seroprevalence of Kaposi's 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
50	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
51	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
52	Clinical Features of Candidiasis in Patients With Inherited Interleukin 12 Receptor Î²1 Deficiency. Clinical Infectious Diseases, 2014, 58, 204-213.	5.8	98
53	Early-onset inflammatory bowel disease and common variable immunodeficiency-like disease caused by IL-21 deficiency. Journal of Allergy and Clinical Immunology, 2014, 133, 1651-1659.e12.	2.9	124
54	Biallelic loss-of-function mutation in NIK causes a primary immunodeficiency with multifaceted aberrant lymphoid immunity. Nature Communications, 2014, 5, 5360.	12.8	116

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55	NK Cell Killer Ig-like Receptor Repertoire Acquisition and Maturation Are Strongly Modulated by HLA Class I Molecules. <i>Journal of Immunology</i> , 2014, 192, 2602-2610.	0.8	19
56	A Novel G6PC3 Gene Mutation in a Patient With Severe Congenital Neutropenia. <i>Journal of Pediatric Hematology/Oncology</i> , 2013, 35, e81-e83.	0.6	12
57	B-cell subsets in patients with transient hypogammaglobulinemia of infancy, partial IgA deficiency, and selective IgM deficiency. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2013, 23, 94-100.	1.3	17
58	815 Patients with Primary Immunodeficiency Disorders in Pediatric Intensive Care Unit: Outcomes and Mortality-Associated Risk Factors. <i>Archives of Disease in Childhood</i> , 2012, 97, A234-A235.	1.9	0
59	Pressure-induced angioedema associated with endotracheal tube: successful treatment with epinephrine in two cases. <i>European Journal of Pediatrics</i> , 2012, 171, 1573-1575.	2.7	3
60	Selective IgA Deficiency: Clinical and Laboratory Features of 118 Children in Turkey. <i>Journal of Clinical Immunology</i> , 2012, 32, 961-966.	3.8	57
61	Diagnosis of Immediate Hypersensitivity to $\beta$ -Lactam Antibiotics Can Be Made Safely with Current Approaches. <i>International Archives of Allergy and Immunology</i> , 2012, 157, 311-317.	2.1	19
62	HLA haploidentical transplantations for primary immunodeficiencies: A single-center experience. <i>Pediatric Transplantation</i> , 2012, 16, 451-457.	1.0	7
63	Oral Lactoferrin Prophylaxis to Prevent Sepsis and Necrotising Enterocolitis of Very Low Birth Weight Neonates in Neonatal Intensive Care Unit and Effect on T-Regulatory Cells. <i>Pediatric Research</i> , 2011, 70, 471-471.	2.3	0
64	The Effect of Mode of Delivery on T Regulatory (Treg) Cells of Cord Blood. <i>Indian Journal of Pediatrics</i> , 2011, 78, 1234-1238.	0.8	13
65	Two Patients with Partial DiGeorge Syndrome Presenting with Attention Disorder and Learning Difficulties - Case Report. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2011, 3, 95-97.	0.9	9
66	Hyperimmunoglobulin E Job s Syndrome and Staphylococcal Botryomycosis in a Child. <i>Gazi Medical Journal</i> , 2011, 22, 124-126.	0.0	1
67	Bacille Calmette-Guérin lymphadenitis and recurrent oral candidiasis in an infant with a new mutation leading to interleukin-12 receptor beta-1 deficiency. <i>Journal of Investigational Allergology and Clinical Immunology</i> , 2011, 21, 401-4.	1.3	22
68	CD40 ligand deficiency with grade III liver fibrosis, transplanted by a treosulphan-based conditioning regimen. <i>Experimental and Clinical Transplantation</i> , 2011, 9, 349-52.	0.2	0
69	Revisiting Human IL-12R $\beta$ 1 Deficiency. <i>Medicine (United States)</i> , 2010, 89, 381-402.	1.0	367
70	Purine nucleoside phosphorylase deficiency with fatal course in two sisters. <i>European Journal of Pediatrics</i> , 2010, 169, 311-314.	2.7	22
71	Reply to correspondence letter by Luis Ignacio Gonzalez-Granado. <i>European Journal of Pediatrics</i> , 2010, 169, 519-519.	2.7	0
72	Eponym. <i>European Journal of Pediatrics</i> , 2010, 169, 657-660.	2.7	9

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73	Kostmann disease with developmental delay in three patients. <i>European Journal of Pediatrics</i> , 2010, 169, 759-762.	2.7	10
74	Late onset hemorrhagic cystitis in a hematopoietic stem cell recipient: Treatment with intravesical hyaluronic acid. <i>Pediatric Transplantation</i> , 2010, 14, E79-E82.	1.0	12
75	Soy Isoflavones Ameliorate the Adverse Effects of Chemotherapy in Children. <i>Nutrition and Cancer</i> , 2010, 62, 1001-1005.	2.0	25
76	Invasive <i>Haemophilus influenzae</i> Infections in Two Children who Vaccinated with <i>Haemophilus influenzae</i> type b Vaccine. <i>Cocuk Enfeksiyon Dergisi</i> , 2010, 4, 76-78.	0.1	3
77	DOES SERUM SOLUBLE VASCULAR ENDOTHELIAL GROWTH FACTOR LEVELS HAVE DIFFERENT IMPORTANCE IN PEDIATRIC ACUTE LEUKEMIA AND MALIGNANT LYMPHOMA PATIENTS?. <i>Pediatric Hematology and Oncology</i> , 2010, 27, 503-516.	0.8	9
78	Combined Immunodeficiencies: Clinical, Immunological Features And Outcome Of 56 Cases From A Single Institution. <i>Journal of Allergy and Clinical Immunology</i> , 2010, 125, AB75.	2.9	0
79	CD27 expression on lymphocyte and sCD27 levels in children with asthma. <i>Allergologia Et Immunopathologia</i> , 2010, 38, 327-332.	1.7	1
80	Asymptomatic catheter related <i>Rhizobium radiobacter</i> infection in a haploidentical hemapoetic stem cell recipient. <i>Journal of Infection in Developing Countries</i> , 2010, 4, 530-532.	1.2	6
81	Effects of soy isoflavones (genistein) on chemotherapy and radiotherapy toxicities in childhood cancer patients.. <i>Journal of Clinical Oncology</i> , 2010, 28, e20008-e20008.	1.6	0
82	Primary immune deficiency disease awareness among a group of Turkish physicians. <i>Turkish Journal of Pediatrics</i> , 2010, 52, 372-7.	0.6	7
83	Selective Immunoglobulin M Deficiency Presenting with Recurrent Impetigo: A Case Report and Review of the Literature. <i>International Archives of Allergy and Immunology</i> , 2009, 149, 283-288.	2.1	18
84	Soluble CD27 Levels in Children with Acute and Chronic Renal Failure. <i>Journal of Allergy and Clinical Immunology</i> , 2009, 123, S226-S226.	2.9	0
85	Percutaneous transcatheter retrieval of intracardiac central venous catheter fragments in two infants using Amplatz Goose Neck snare. <i>Turkish Journal of Pediatrics</i> , 2009, 51, 519-23.	0.6	4
86	An unconditioned bone marrow transplantation in a child with purine nucleoside phosphorylase deficiency and its unique complication. <i>Pediatric Transplantation</i> , 2008, 12, 479-482.	1.0	26
87	HLA Class I Deficiency: Clinical, Immunological and Genetic Characteristics of Three Patient. <i>Journal of Allergy and Clinical Immunology</i> , 2008, 121, S86-S86.	2.9	0
88	Mutations in <i>STAT3</i> and <i>IL12RB1</i> impair the development of human IL-17-producing T cells. <i>Journal of Experimental Medicine</i> , 2008, 205, 1543-1550.	8.5	406
89	A novel mutation for TAP deficiency and its possible association with Toxoplasmosis. <i>Parasitology International</i> , 2006, 55, 219-222.	1.3	19
90	Tonsillectomy and the immune system. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2006, 70, 175-176.	1.0	0

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91	Stable mixed chimerism after hematopoietic stem cell transplantation in Wiskott-Aldrich syndrome. <i>Pediatric Transplantation</i> , 2006, 10, 395-399.	1.0	5
92	Complete deficiency of the IL-12 receptor $\beta$ 1 chain: three unrelated Turkish children with unusual clinical features. <i>European Journal of Pediatrics</i> , 2006, 165, 415-417.	2.7	19
93	A case of Wiskott-Aldrich syndrome with de novo mutation at exon 4. <i>Turkish Journal of Pediatrics</i> , 2006, 48, 66-8.	0.6	2
94	Meningococcal meningitis and complement component 6 deficiency associated with oculocutaneous albinism. <i>European Journal of Pediatrics</i> , 2005, 164, 177-179.	2.7	7
95	Granulocyte Transfusions in Children With Chronic Granulomatous Disease and Invasive Aspergillosis. <i>Therapeutic Apheresis and Dialysis</i> , 2005, 9, 137-141.	0.9	41
96	Serum sFas and sFas-Ligand Levels in Childhood Asthma. <i>Pediatric Asthma, Allergy and Immunology</i> , 2004, 17, 126-130.	0.2	2
97	Unusual cause of respiratory distress: Chilaiditi syndrome. <i>Pediatrics International</i> , 2004, 46, 188-190.	0.5	13
98	Transient hypogammaglobulinemia of infancy and early childhood: outcome of 30 cases. <i>Turkish Journal of Pediatrics</i> , 2004, 46, 120-4.	0.6	16
99	Peripheral blood lymphocyte subsets in healthy Turkish children. <i>Turkish Journal of Pediatrics</i> , 2004, 46, 125-30.	0.6	55
100	Direct Genetic Correction as a New Method for Diagnosis and Molecular Characterization of MHC Class II Deficiency. <i>Molecular Therapy</i> , 2002, 6, 824-829.	8.2	10
101	RECURRENT ARTERIAL THROMBOSIS IN A CHILD: Primary Antiphospholipid Antibody Syndrome. <i>Pediatric Hematology and Oncology</i> , 2002, 19, 59-66.	0.8	12
102	Is immune system influenced by adenotonsillectomy in children?. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2002, 66, 251-257.	1.0	41
103	657del5 mutation in the NBS1 gene is associated with Nijmegen breakage syndrome in a Turkish family. <i>Clinical Genetics</i> , 2002, 62, 84-88.	2.0	7
104	An intensive approach to the treatment of disseminated BCG infection in a SCID patient. <i>Bone Marrow Transplantation</i> , 2002, 30, 45-47.	2.4	22
105	Circulating adhesion molecule levels in childhood asthma. <i>Indian Pediatrics</i> , 2002, 39, 1017-21.	0.4	2
106	NATURAL KILLER CELL NUMBERS AND CYTOTOXIC ACTIVITY IN PEDIATRIC HODGKIN DISEASE. <i>Pediatric Hematology and Oncology</i> , 2000, 17, 133-139.	0.8	2
107	Disseminated Bacille Calmette-Guerin (BCG) Disease in A Successfully Bone Marrow Transplanted Scid Patient. <i>Pediatric Research</i> , 1999, 45, 768-768.	2.3	0