Alessandro Plebani

List of Publications by Year
in descending order

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206	17,138	23567 58	15732 125
papers	citations	h-index	g-index

210	210	210	17533
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Immunological Evaluation of Patients Affected with Jacobsen Syndrome Reveals Profound Not Age-Related Lymphocyte Alterations. Journal of Clinical Immunology, 2022, 42, 365-374.	3.8	4
2	Lack of DOCK8 impairs the primary biologic functions of human NK cells and abrogates CCR7 surface expression in a WASP-independent manner. Clinical Immunology, 2022, 237, 108974.	3.2	2
3	The Impact of SARS-CoV-2 Infection in Patients with Inborn Errors of Immunity: the Experience of the Italian Primary Immunodeficiencies Network (IPINet). Journal of Clinical Immunology, 2022, 42, 935-946.	3.8	21
4	Exploration of a panel of urine biomarkers of kidney disease in two paediatric cohorts with Type 1 diabetes mellitus of differing duration. Diabetology and Metabolic Syndrome, 2022, 14, 71.	2.7	0
5	Follicular helper T cell signature of replicative exhaustion, apoptosis, and senescence in common variable immunodeficiency. European Journal of Immunology, 2022, 52, 1171-1189.	2.9	9
6	Coronavirus disease 2019 in patients with inborn errors of immunity: An international study. Journal of Allergy and Clinical Immunology, 2021, 147, 520-531.	2.9	278
7	Fatal SARS-CoV-2 infection in a male patient with Good's syndrome. Clinical Immunology, 2021, 223, 108644.	3.2	15
8	Predominantly Antibody Deficiencies. , 2021, , .		0
9	Modeling, optimization, and comparable efficacy of T cell and hematopoietic stem cell gene editing for treating hyperâ€IgM syndrome. EMBO Molecular Medicine, 2021, 13, e13545.	6.9	36
10	Granulomatous Lymphocytic Interstitial Lung Disease (GLILD) in Common Variable Immunodeficiency (CVID): A Multicenter Retrospective Study of Patients From Italian PID Referral Centers. Frontiers in Immunology, 2021, 12, 627423.	4.8	25
11	Temporal viral loads in respiratory and gastrointestinal tract and serum antibody responses during SARS-CoV-2 infection in an Italian pediatric cohort. Clinical Immunology, 2021, 225, 108695.	3.2	2
12	Clinical outcome, incidence, and SARS-CoV-2 infection-fatality rates in Italian patients with inborn errors of immunity. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 2904-2906.e2.	3.8	56
13	Complete CD95/FAS deficiency due to complex homozygous germline TNFRSF6 mutations in an adult patient with mild autoimmune lymphoproliferative syndrome (ALPS). Clinical Immunology, 2021, 228, 108757.	3.2	3
14	Known and potential molecules associated with altered B cell development leading to predominantly antibody deficiencies. Pediatric Allergy and Immunology, 2021, 32, 1601-1615.	2.6	10
15	Simple Measurement of IgA Predicts Immunity and Mortality in Ataxia-Telangiectasia. Journal of Clinical Immunology, 2021, 41, 1878-1892.	3.8	9
16	Combined immunodeficiency with autoimmunity caused by a homozygous missense mutation in inhibitor of nuclear factor ?B kinase alpha (IKKÎ±). Science Immunology, 2021, 6, eabf6723.	11.9	6
17	Emergence of Letermovir-resistant HCMV UL56 mutant during rescue treatment in a liver transplant recipient with ganciclovir-resistant infection HCMV: a case report. BMC Infectious Diseases, 2021, 21, 994.	2.9	11
18	Factors Associated With Severe Gastrointestinal Diagnoses in Children With SARS-CoV-2 Infection or Multisystem Inflammatory Syndrome. JAMA Network Open, 2021, 4, e2139974.	5.9	24

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19	Health-Related Quality of Life and Emotional Difficulties in Chronic Granulomatous Disease: Data on Adult and Pediatric Patients from Italian Network for Primary Immunodeficiency (IPINet). Journal of Clinical Immunology, 2020, 40, 289-298.	3.8	11
20	NFKB2 regulates human Tfh and Tfr pool formation and germinal center potential. Clinical Immunology, 2020, 210, 108309.	3.2	14
21	Mycoplasma infection may complicate the clinical course of SARS-Co-V-2 associated Kawasaki-like disease in children. Clinical Immunology, 2020, 221, 108613.	3.2	10
22	Activated Phosphoinositide 3-Kinase Delta Syndrome 1: Clinical and Immunological Data from an Italian Cohort of Patients. Journal of Clinical Medicine, 2020, 9, 3335.	2.4	23
23	Successful hematopoietic stem cell transplantation for complete CTLA-4 haploinsufficiency due to a de novo monoallelic 2q33.2-2q33.3 deletion. Clinical Immunology, 2020, 220, 108589.	3.2	2
24	Paediatric MAS/HLH caused by a novel monoallelic activating mutation in p110Î´. Clinical Immunology, 2020, 219, 108543.	3.2	8
25	The Italian Registry for Primary Immunodeficiencies (Italian Primary Immunodeficiency Network;) Tj ETQq1 1 0.784314 rgBT /Overlock	3.8	15
26	Consensus of the Italian Primary Immunodeficiency Network on transition management from pediatric to adult care in patients affected with childhood-onset inborn errors of immunity. Journal of Allergy and Clinical Immunology, 2020, 146, 967-983.	2.9	12
27	RAC2 and primary human immune deficiencies. Journal of Leukocyte Biology, 2020, 108, 687-696.	3.3	31
28	Long-term follow-up of 168 patients with X-linked agammaglobulinemia reveals increased morbidity and mortality. Journal of Allergy and Clinical Immunology, 2020, 146, 429-437.	2.9	59
29	Diagnostic approach of hypogammaglobulinemia in infancy. Pediatric Allergy and Immunology, 2020, 31, 11-12.	2.6	2
30	Characterization of the clinical and immunologic phenotype and management of 157 individuals with 56 distinct heterozygous NFKB1 mutations. Journal of Allergy and Clinical Immunology, 2020, 146, 901-911.	2.9	78
31	A possible role for B cells in COVID-19? Lesson from patients with agammaglobulinemia. Journal of Allergy and Clinical Immunology, 2020, 146, 211-213.e4.	2.9	275
32	Immunophenotype Anomalies Predict the Development of Autoimmune Cytopenia in 22q11.2 Deletion Syndrome. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2369-2376.	3.8	38
33	From Natural Killer Cell Receptor Discovery to Characterization of Natural Killer Cell Defects in Primary Immunodeficiencies. Frontiers in Immunology, 2019, 10, 1757.	4.8	2
34	Long-Term Outcome of WHIM Syndrome in 18 Patients: High Risk of Lung Disease and HPV-Related Malignancies. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1568-1577.	3.8	40
35	Autosomal-dominant hyper-IgE syndrome is associated with appearance of infections early in life and/or neonatal rash: Evidence from the Italian cohort of 61 patients with elevated IgE. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2072-2075.e4.	3.8	10
36	Clinical, Immunological, and Molecular Features of Typical and Atypical Severe Combined Immunodeficiency: Report of the Italian Primary Immunodeficiency Network. Frontiers in Immunology, 2019, 10, 1908.	4.8	41

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37	The RAC2-PI3K axis regulates human NK cell maturation and function. Clinical Immunology, 2019, 208, 108257.	3.2	11
38	Clinical and Laboratory Features of 184 Italian Pediatric Patients Affected with Selective IgA Deficiency (SIgAD): a Longitudinal Single-Center Study. Journal of Clinical Immunology, 2019, 39, 470-475.	3.8	27
39	Double-blind, placebo-controlled, randomized trial on low-dose azithromycin prophylaxis in patients with primary antibody deficiencies. Journal of Allergy and Clinical Immunology, 2019, 144, 584-593.e7.	2.9	54
40	Homozygous frameshift mutations in FAT1 cause a syndrome characterized by colobomatous-microphthalmia, ptosis, nephropathy and syndactyly. Nature Communications, 2019, 10, 1180.	12.8	27
41	X-linked agammaglobulinemia (XLA): Phenotype, diagnosis, and therapeutic challenges around the world. World Allergy Organization Journal, 2019, 12, 100018.	3.5	83
42	A monoallelic activating mutation in RAC2 resulting in a combined immunodeficiency. Journal of Allergy and Clinical Immunology, 2019, 143, 1649-1653.e3.	2.9	37
43	Imaging of Bronchial Pathology in Antibody Deficiency: Data from the European Chest CT Group. Journal of Clinical Immunology, 2019, 39, 45-54.	3.8	32
44	A novel monoallelic gain of function mutation in p110Î´ causing atypical activated phosphoinositide 3-kinase Î´ syndrome (APDS-1). Clinical Immunology, 2019, 200, 31-34.	3.2	11
45	Early B cell developmental impairment with progressive B cell deficiency in NFKB2 mutated CVID disease without autoimmunity. Clinical Immunology, 2019, 205, 153-155.	3.2	4
46	Progressive severe B cell and NK cell deficiency with T cell senescence in adult CD40L deficiency. Clinical Immunology, 2018, 190, 11-14.	3.2	14
47	Recurrent Bleedings in Newborn: A Factor VII Deficiency Case Report. Transfusion Medicine and Hemotherapy, 2018, 45, 104-106.	1.6	3
48	Long term outcome of eight patients with type 1 Leukocyte Adhesion Deficiency (LAD-1): Not only infections, but high risk of autoimmune complications. Clinical Immunology, 2018, 191, 75-80.	3.2	33
49	Impaired platelet activation in patients with hereditary deficiency of p47^{phox}. British Journal of Haematology, 2018, 180, 454-456.	2.5	5
50	In vivo effects of dexamethasone on blood gene expression in ataxia telangiectasia. Molecular and Cellular Biochemistry, 2018, 438, 153-166.	3.1	20
51	Neurovisual Assessment in Children with Ataxia Telangiectasia. Neuropediatrics, 2018, 49, 026-034.	0.6	6
52	Novel biallelic TRNT1 mutations resulting in sideroblastic anemia, combined B and T cell defects, hypogammaglobulinemia, recurrent infections, hypertrophic cardiomyopathy and developmental delay. Clinical Immunology, 2018, 188, 20-22.	3.2	24
53	Chronic Granulomatous Disease in children: a single center experience. Clinical Immunology, 2018, 188, 12-19.	3.2	10
54	A de novo monoallelic CTLA-4 deletion causing pediatric onset CVID with recurrent autoimmune cytopenias and severe enteropathy. Clinical Immunology, 2018, 197, 186-188.	3.2	4

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55	Phenotype, penetrance, and treatment of 133 cytotoxic T-lymphocyte antigen 4â€“insufficient subjects. Journal of Allergy and Clinical Immunology, 2018, 142, 1932-1946.	2.9	344
56	CTLA-4 regulates human Natural Killer cell effector functions. Clinical Immunology, 2018, 194, 43-45.	3.2	30
57	Circulating Follicular Helper and Follicular Regulatory T Cells Are Severely Compromised in Human CD40 Deficiency: A Case Report. Frontiers in Immunology, 2018, 9, 1761.	4.8	27
58	Response to the Letter to the Editor Regarding â€œFunctional evaluation of natural killer cell cytotoxic activity in NFKB-2 mutated patientsâ€: Immunology Letters, 2018, 200, 16-17.	2.5	0
59	Impaired natural killer cell functions in patients with signal transducer and activator of transcription 1 (STAT1) gain-of-function mutations. Journal of Allergy and Clinical Immunology, 2017, 140, 553-564.e4.	2.9	58
60	Beta2 integrins are required for follicular helper T cell differentiation in humans. Clinical Immunology, 2017, 180, 60-62.	3.2	3
61	Early Identification of Lung Fungal Infections in Chronic Granulomatous Disease (CGD) Using Multidetector Computer Tomography. Journal of Clinical Immunology, 2017, 37, 36-41.	3.8	8
62	Shift from intravenous or 16% subcutaneous replacement therapy to 20% subcutaneous immunoglobulin in patients with primary antibody deficiencies. International Journal of Immunopathology and Pharmacology, 2017, 30, 73-82.	2.1	21
63	NFKB1 regulates human NK cell maturation and effector functions. Clinical Immunology, 2017, 175, 99-108.	3.2	38
64	Response to the Letter to the Editor Regarding â€œKinetics of Radiological Response of Thoracic Invasive Fungal Disease in Chronic Granulomatous Diseaseâ€: Journal of Clinical Immunology, 2017, 37, 744-745.	3.8	0
65	Early and late B-cell developmental impairment in nuclear factor kappa B, subunit 1â€“mutated common variable immunodeficiency disease. Journal of Allergy and Clinical Immunology, 2017, 139, 349-352.e1.	2.9	30
66	Natural Killer Cells from Patients with Recombinase-Activating Gene and Non-Homologous End Joining Gene Defects Comprise a Higher Frequency of CD56bright NKG2A+++ Cells, and Yet Display Increased Degranulation and Higher Perforin Content. Frontiers in Immunology, 2017, 8, 798.	4.8	41
67	NOX 5 is expressed in platelets from patients with chronic granulomatous disease. Thrombosis and Haemostasis, 2016, 116, 198-200.	3.4	8
68	Diagnostics of Primary Immunodeficiencies through Next-Generation Sequencing. Frontiers in Immunology, 2016, 7, 466.	4.8	80
69	Progressive severe B cell deficiency in pediatric Rubinstein-Taybi syndrome. Clinical Immunology, 2016, 173, 181-183.	3.2	7
70	Impairment of dendritic cell functions in patients with adaptor protein-3 complex deficiency. Blood, 2016, 127, 3382-3386.	1.4	11
71	Common variants at PVT1, ATG13â€“AMBRA1, AHI1 and CLEC16A are associated with selective IgA deficiency. Nature Genetics, 2016, 48, 1425-1429.	21.4	67
72	Novel compound heterozygous mutations in a child with Ataxia-Telangiectasia showing unrelated cerebellar disorders. Journal of the Neurological Sciences, 2016, 371, 48-53.	0.6	7

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73	Proteus syndrome: evaluation of the immunological profile. Orphanet Journal of Rare Diseases, 2016, 11, 3.	2.7	5
74	Monoallelic BAFFR P21R/H159Y Mutations and Familiar Primary Antibody Deficiencies. Journal of Clinical Immunology, 2016, 36, 1-3.	3.8	9
75	Clinical and immunological data of nine patients with chronic mucocutaneous candidiasis disease. Data in Brief, 2016, 7, 311-315.	1.0	10
76	p85Î± is an intrinsic regulator of human natural killer cell effector functions. Journal of Allergy and Clinical Immunology, 2016, 138, 605-608.e3.	2.9	7
77	Clinical heterogeneity of dominant chronic mucocutaneous candidiasis disease: presenting as treatment-resistant candidiasis and chronic lung disease. Clinical Immunology, 2016, 164, 1-9.	3.2	27
78	Reduced germinal center follicular helper T cells but normal follicular regulatory T cells in the tonsils of a patient with a mutation in the PI3KR1 gene. Clinical Immunology, 2016, 164, 43-44.	3.2	11
79	Cohort of Iranian Patients with Congenital Agammaglobulinemia: Mutation Analysis and Novel Gene Defects. Expert Review of Clinical Immunology, 2016, 12, 479-486.	3.0	22
80	Reduction of CRKL expression in patients with partial DiGeorge syndrome is associated with impairment of T-cell functions. Journal of Allergy and Clinical Immunology, 2016, 138, 229-240.e3.	2.9	16
81	Correlation of bone marrow abnormalities, peripheral lymphocyte subsets and clinical features in uncomplicated common variable immunodeficiency (CVID) patients. Clinical Immunology, 2016, 163, 10-13.	3.2	10
82	Altered germinal center reaction and abnormal B cell peripheral maturation in PI3KR1-mutated patients presenting with HIGM-like phenotype. Clinical Immunology, 2015, 159, 33-36.	3.2	51
83	The Tec Kinaseâ€“Regulated Phosphoproteome Reveals a Mechanism for the Regulation of Inhibitory Signals in Murine Macrophages. Journal of Immunology, 2015, 195, 246-256.	0.8	31
84	Positive effect of erythrocyte-delivered dexamethasone in ataxia-telangiectasia. Neurology: Neuroimmunology and NeuroInflammation, 2015, 2, e98.	6.0	59
85	Defective natural killerâ€“cell cytotoxic activity in NFKB2-mutated CVID-like disease. Journal of Allergy and Clinical Immunology, 2015, 135, 1641-1643.e3.	2.9	68
86	Gastrointestinal Pathologic Abnormalities in Pediatric- and Adult-Onset Common Variable Immunodeficiency. Digestive Diseases and Sciences, 2015, 60, 2384-2389.	2.3	17
87	Severe congenital neutropenia due to G6PC3 deficiency: early and delayed phenotype in two patients with two novel mutations. Italian Journal of Pediatrics, 2014, 40, 80.	2.6	8
88	A CXCR1 haplotype hampers HIV-1 matrix protein p17 biological activity. Aids, 2014, 28, 2355-2364.	2.2	5
89	Different Degrees of NADPH Oxidase 2 Regulation and In Vivo Platelet Activation: Lesson From Chronic Granulomatous Disease. Journal of the American Heart Association, 2014, 3, e000920.	3.7	31
90	Intergenerational and intrafamilial phenotypic variability in 22q11.2 Deletion syndrome subjects. BMC Medical Genetics, 2014, 15, 1.	2.1	48

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91	Bruton tyrosine kinase mediates TLR9-dependent human dendritic cell activation. Journal of Allergy and Clinical Immunology, 2014, 133, 1644-1650.e4.	2.9	62
92	Autosomal Recessive Agammaglobulinemia: A Novel Non-sense Mutation in CD79a. Journal of Clinical Immunology, 2014, 34, 138-141.	3.8	16
93	Autosomal Recessive Agammaglobulinemia: The Third Case of Igî² Deficiency Due to a Novel Non-sense Mutation. Journal of Clinical Immunology, 2014, 34, 425-427.	3.8	11
94	Expansion of CCR4+ activated T cells is associated with memory B cell reduction in DOCK8-deficient patients. Clinical Immunology, 2014, 152, 164-170.	3.2	11
95	A common single nucleotide polymorphism impairs B-cell activating factor receptor's multimerization, contributing to common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2014, 133, 1222-1225.e10.	2.9	60
96	BAFF-R mutations in Good's syndrome. Clinical Immunology, 2014, 153, 91-93.	3.2	20
97	Body experiences, emotional competence, and psychosocial functioning in juvenile idiopathic arthritis. Rheumatology International, 2013, 33, 2045-2052.	3.0	39
98	Does NADPH Oxidase Deficiency Cause Artery Dilatation in Humans?. Antioxidants and Redox Signaling, 2013, 18, 1491-1496.	5.4	27
99	Reduced Atherosclerotic Burden in Subjects With Genetically Determined Low Oxidative Stress. Arteriosclerosis, Thrombosis, and Vascular Biology, 2013, 33, 406-412.	2.4	52
100	Ribosomal Protein SA Haploinsufficiency in Humans with Isolated Congenital Asplenia. Science, 2013, 340, 976-978.	12.6	176
101	The Significance of Duodenal Mucosal Atrophy in Patients With Common Variable Immunodeficiency. American Journal of Clinical Pathology, 2012, 138, 185-189.	0.7	101
102	A novel primary human immunodeficiency due to deficiency in the WASP-interacting protein WIP. Journal of Experimental Medicine, 2012, 209, 29-34.	8.5	158
103	A randomized trial of oral betamethasone to reduce ataxia symptoms in ataxia telangiectasia. Movement Disorders, 2012, 27, 1312-1316.	3.9	73
104	DOCK8 functions as an adaptor that links TLR-MyD88 signaling to B cell activation. Nature Immunology, 2012, 13, 612-620.	14.5	205
105	Exome sequencing reveals a pallidin mutation in a Hermansky-Pudlakâ€“like primary immunodeficiency syndrome. Blood, 2012, 119, 3185-3187.	1.4	76
106	Soluble BAFF Levels Inversely Correlate with Peripheral B Cell Numbers and the Expression of BAFF Receptors. Journal of Immunology, 2012, 188, 497-503.	0.8	155
107	Deleterious Mutations in LRBA Are Associated with a Syndrome of Immune Deficiency and Autoimmunity. American Journal of Human Genetics, 2012, 90, 986-1001.	6.2	452
108	X-linked lymphoproliferative disease due to SAP/SH2D1A deficiency: a multicenter study on the manifestations, management and outcome of the disease. Blood, 2011, 117, 53-62.	1.4	268

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109	Clinical heterogeneity and diagnostic delay of autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy syndrome. Clinical Immunology, 2011, 139, 6-11.	3.2	49
110	Effectiveness of Immunoglobulin Replacement Therapy on Clinical Outcome in Patients with Primary Antibody Deficiencies: Results from a Multicenter Prospective Cohort Study. Journal of Clinical Immunology, 2011, 31, 315-322.	3.8	252
111	Thymic and Bone Marrow Output in Patients with Common Variable Immunodeficiency. Journal of Clinical Immunology, 2011, 31, 540-549.	3.8	35
112	SH2â€domain mutations in STAT3 in hyperâ€IgE syndrome patients result in impairment of ILâ€10 function. European Journal of Immunology, 2011, 41, 3075-3084.	2.9	26
113	Use of linezolid in infants and children: a retrospective multicentre study of the Italian Society for Paediatric Infectious Diseases. Journal of Antimicrobial Chemotherapy, 2011, 66, 2393-2397.	3.0	31
114	Inherited Human gp91 ^{phox} Deficiency Is Associated With Impaired Isoprostane Formation and Platelet Dysfunction. Arteriosclerosis, Thrombosis, and Vascular Biology, 2011, 31, 423-434.	2.4	124
115	Favourable and sustained response to anakinra in tumour necrosis factor receptor-associated periodic syndrome (TRAPS) with or without AA amyloidosis. Annals of the Rheumatic Diseases, 2011, 70, 1511-1512.	0.9	86
116	Common Variable Immunodeficiency. Journal of Computer Assisted Tomography, 2010, 34, 395-401.	0.9	29
117	Different molecular behavior of CD40 mutants causing hyper-IgM syndrome. Blood, 2010, 116, 5867-5874.	1.4	29
118	Implications of gluten exposure period, CD clinical forms, and HLA typing in the association between celiac disease and dental enamel defects in children. A caseâ€“control study. International Journal of Paediatric Dentistry, 2010, 20, 119-124.	1.8	24
119	The transmembrane activator TACI triggers immunoglobulin class switching by activating B cells through the adaptor MyD88. Nature Immunology, 2010, 11, 836-845.	14.5	295
120	Defect of regulatory T cells in patients with Omenn syndrome. Journal of Allergy and Clinical Immunology, 2010, 125, 209-216.	2.9	83
121	Poor health-related quality of life and abnormal psychosocial adjustment in Italian children with perinatal HIV infection receiving highly active antiretroviral treatment. AIDS Care - Psychological and Socio-Medical Aspects of AIDS/HIV, 2010, 22, 858-865.	1.2	37
122	A HomozygousCARD9Mutation in a Family with Susceptibility to Fungal Infections. New England Journal of Medicine, 2009, 361, 1727-1735.	27.0	733
123	Hereditary Deficiency of gp91 ^{phox} Is Associated With Enhanced Arterial Dilatation. Circulation, 2009, 120, 1616-1622.	1.6	123
124	Allele *1 of HS1.2 Enhancer Associates with Selective IgA Deficiency and IgM Concentration. Journal of Immunology, 2009, 183, 8280-8285.	0.8	19
125	The Quality of Life of Children and Adolescents with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2009, 29, 501-507.	3.8	34
126	Immunization after the elective end of antineoplastic chemotherapy in children. Pediatric Blood and Cancer, 2009, 52, 165-168.	1.5	16

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127	Immunoglobulin D enhances immune surveillance by activating antimicrobial, proinflammatory and B cellâ€“stimulating programs in basophils. Nature Immunology, 2009, 10, 889-898.	14.5	362
128	Unrelated hematopoietic stem cell transplantation for Cernunnosâ€XLF deficiency. Pediatric Transplantation, 2009, 13, 785-789.	1.0	30
129	Molecular characterization of a large cohort of patients with Chronic Granulomatous Disease and identification of novel CYBB mutations: An Italian multicenter study. Molecular Immunology, 2009, 46, 1935-1941.	2.2	36
130	Differentiating PFAPA Syndrome From Monogenic Periodic Fevers. Pediatrics, 2009, 124, e721-e728.	2.1	138
131	MULTIPLE RELAPSES OF VISCERAL LEISHMANIASIS IN AN ADOLESCENT WITH IDIOPATHIC CD4+ LYMPHOCYTOPENIA ASSOCIATED WITH NOVEL IMMUNOPHENOTYPIC AND MOLECULAR FEATURES. Pediatric Infectious Disease Journal, 2009, 28, 161-163.	2.0	6
132	Relevance of biallelic versus monoallelic TNFRSF13B mutations in distinguishing disease-causing from risk-increasing TNFRSF13B variants in antibody deficiency syndromes. Blood, 2009, 113, 1967-1976.	1.4	254
133	Prospective Study on CVID Patients with Adverse Reactions to Intravenous or Subcutaneous IgG Administration. Journal of Clinical Immunology, 2008, 28, 263-267.	3.8	33
134	Autosomal recessive agammaglobulinemia: Novel insights from mutations in Ig-beta. Current Allergy and Asthma Reports, 2008, 8, 404-408.	5.3	10
135	The EUROclass trial: defining subgroups in common variable immunodeficiency. Blood, 2008, 111, 77-85.	1.4	722
136	Clinical features, long-term follow-up and outcome of a large cohort of patients with Chronic Granulomatous Disease: An Italian multicenter study. Clinical Immunology, 2008, 126, 155-164.	3.2	293
137	Sensorineural Hearing Loss in Primary Antibody Deficiency Disorders. Journal of Pediatrics, 2008, 153, 293-296.	1.8	14
138	Toll-Like Receptor Stimulation Induces Higher TNF-α Secretion in Peripheral Blood Mononuclear Cells from Patients with Hyper IgE Syndrome. International Archives of Allergy and Immunology, 2008, 146, 190-194.	2.1	14
139	Viral Double-Stranded RNA Triggers Ig Class Switching by Activating Upper Respiratory Mucosa B Cells through an Innate TLR3 Pathway Involving BAFF. Journal of Immunology, 2008, 181, 276-287.	0.8	105
140	Characterization of Ig Gene Somatic Hypermutation in the Absence of Activation-Induced Cytidine Deaminase. Journal of Immunology, 2008, 181, 1299-1306.	0.8	27
141	CpG Drives Human Transitional B Cells to Terminal Differentiation and Production of Natural Antibodies. Journal of Immunology, 2008, 180, 800-808.	0.8	209
142	IgÎ² deficiency in humans. Current Opinion in Allergy and Clinical Immunology, 2008, 8, 515-519.	2.3	4
143	Mutations of the IgÎ² gene cause agammaglobulinemia in man. Journal of Experimental Medicine, 2007, 204, 2047-2051.	8.5	87
144	Genetic Causes of Bronchiectasis: Primary Immune Deficiencies and the Lung. Respiration, 2007, 74, 264-275.	2.6	46

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145	Diversion of the Fecal Stream Resolves Ulcerative Colitis Complicating Chronic Granulomatous Disease in an Adult Patient. Journal of Clinical Gastroenterology, 2007, 41, 491-493.	2.2	3
146	Pulmonary and Sinusal Changes in 45 Patients With Primary Immunodeficiencies. Journal of Computer Assisted Tomography, 2007, 31, 620-628.	0.9	32
147	Intestinal Bacteria Trigger T Cell-Independent Immunoglobulin A2 Class Switching by Inducing Epithelial-Cell Secretion of the Cytokine APRIL. Immunity, 2007, 26, 812-826.	14.3	656
148	Memory B-cell subsets as a predictive marker of outcome in hypogammaglobulinemia during infancy. Journal of Allergy and Clinical Immunology, 2007, 120, 474-476.	2.9	19
149	A novel immunodeficiency characterized by the exclusive presence of transitional B cells unresponsive to CpG. Immunology, 2007, 121, 183-188.	4.4	23
150	Long-Term Follow-Up and Outcome of a Large Cohort of Patients with Common Variable Immunodeficiency. Journal of Clinical Immunology, 2007, 27, 308-316.	3.8	465
151	ICOS Deficiency Is Associated with a Severe Reduction of CXCR5+CD4 Germinal Center Th Cells. Journal of Immunology, 2006, 177, 4927-4932.	0.8	349
152	Clinical, Immunological and Molecular Characteristics of 37 Iranian Patients with X-Linked Agammaglobulinemia. International Archives of Allergy and Immunology, 2006, 141, 408-414.	2.1	52
153	Cernunnos, a Novel Nonhomologous End-Joining Factor, Is Mutated in Human Immunodeficiency with Microcephaly. Cell, 2006, 124, 287-299.	28.9	640
154	Linkage of autosomal-dominant common variable immunodeficiency to chromosome 4q. European Journal of Human Genetics, 2006, 14, 867-875.	2.8	46
155	Mutational Analysis of Human BLyS in Patients with Common Variable Immunodeficiency. Journal of Clinical Immunology, 2006, 26, 396-399.	3.8	13
156	Analysis of families with common variable immunodeficiency (CVID) and IgA deficiency suggests linkage of CVID to chromosome 16q. Human Genetics, 2006, 118, 725-729.	3.8	31
157	Combined decrease of defined B and T cell subsets in a group of common variable immunodeficiency patients. Clinical Immunology, 2006, 121, 203-214.	3.2	52
158	Neutrophils from patients withTNFRSF1A mutations display resistance to tumor necrosis factorâ€“induced apoptosis: Pathogenetic and clinical implications. Arthritis and Rheumatism, 2006, 54, 998-1008.	6.7	138
159	X-linked susceptibility to mycobacteria is caused by mutations in NEMO impairing CD40-dependent IL-12 production. Journal of Experimental Medicine, 2006, 203, 1745-1759.	8.5	264
160	Hyper immunoglobulin M syndrome due to CD40 deficiency: clinical, molecular, and immunological features. Immunological Reviews, 2005, 203, 48-66.	6.0	176
161	Re-immunisation schedule in leukaemic children after intensive chemotherapy: a possible strategy. European Journal of Haematology, 2005, 74, 20-23.	2.2	41
162	Reply to Cesaro and Ek et al.: Vaccination after the end of chemotherapy for acute lymphoblastic leukaemia in children. European Journal of Haematology, 2005, 75, 179-180.	2.2	3

#	Article	IF	Citations
163	Letter to the editor: Humoral immunodeficiencies in Down syndrome: Serum IgG subclass and antibody response to hepatitis B vaccine. American Journal of Medical Genetics Part A, 2005, 37, 231-233.	2.4	18
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