Raffaele Badolato

List of Publications by Year
in descending order

169	7,943	43973 48	58464 82
papers	citations	h-index	g-index

173	173	173	10767
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Children living with HIV in Europe: do migrants have worse treatment outcomes?. HIV Medicine, 2022, 23, 186-196.	1.0	2
2	Immunological Evaluation of Patients Affected with Jacobsen Syndrome Reveals Profound Not Age-Related Lymphocyte Alterations. Journal of Clinical Immunology, 2022, 42, 365-374.	2.0	4
3	Off-label use of combined antiretroviral therapy, analysis of data collected by the Italian Register for HIV-1 infection in paediatrics in a large cohort of children. BMC Infectious Diseases, 2022, 22, 55.	1.3	2
4	When a Nontuberculous Mycobacterial Infection Reveals an Error of Immunity. Pediatric Infectious Disease Journal, 2022, Publish Ahead of Print, .	1.1	0
5	Incidence of acute respiratory infections in preschool children in an outpatient setting before and during Covid-19 pandemic in Lombardy Region, Italy. Italian Journal of Pediatrics, 2022, 48, 18.	1.0	14
6	Identical <scp>EP300</scp> variant leading to Rubinsteinâ€“Taybi syndrome with different clinical and immunologic phenotype. American Journal of Medical Genetics, Part A, 2022, 188, 2129-2134.	0.7	2
7	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.	1.4	2
8	Pathogenesis of Autoimmune Cytopenias in Inborn Errors of Immunity Revealing Novel Therapeutic Targets. Frontiers in Immunology, 2022, 13, 846660.	2.2	3
9	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.	2.0	21
10	The Effective Management of Fever in Pediatrics and Insights on Remote Management: Experts' Consensus Using a Delphi Approach. Frontiers in Pediatrics, 2022, 10, 834673.	0.9	1
11	Coronavirus disease 2019 in patients with inborn errors of immunity: An international study. Journal of Allergy and Clinical Immunology, 2021, 147, 520-531.	1.5	278
12	Absent B cells, agammaglobulinemia, and hypertrophic cardiomyopathy in folliculin-interacting protein 1 deficiency. Blood, 2021, 137, 493-499.	0.6	26
13	Plasmacytoid Dendritic Cells Depletion and Elevation of IFN-Î³ Dependent Chemokines CXCL9 and CXCL10 in Children With Multisystem Inflammatory Syndrome. Frontiers in Immunology, 2021, 12, 654587.	2.2	39
14	Impact of a diagnostic therapeutic educational pathway program for asthma management in preschool children. Italian Journal of Pediatrics, 2021, 47, 60.	1.0	1
15	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.	2.0	56
16	Case Report: Hypomorphic Function and Somatic Reversion in DOCK8 Deficiency in One Patient With Two Novel Variants and Sclerosing Cholangitis. Frontiers in Immunology, 2021, 12, 673487.	2.2	5
17	IFN-Î± levels in ruxolitinib-treatead Aicardi-GoutiÃ¨res patient during SARS-CoV-2 infection: A case report. Clinical Immunology, 2021, 227, 108743.	1.4	1
18	Genetic, Immunological, and Clinical Features of 32 Patients with Autosomal Recessive STAT1 Deficiency. Journal of Immunology, 2021, 207, 133-152.	0.4	33

#	Article	IF	Citations
19	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.	1.4	3
20	Alternative Approach to Varicella Zoster Virus Prevention in a Child with Lymphopenia. Journal of Clinical Immunology, 2021, 41, 1681-1682.	2.0	0
21	Case Report: Analysis of Inflammatory Cytokines IL-6, CCL2/MCP1, CCL5/RANTES, CXCL9/MIG, and CXCL10/IP10 in a Cystic Fibrosis Patient Cohort During the First Wave of the COVID-19 Pandemic. Frontiers in Pediatrics, 2021, 9, 645063.	0.9	3
22	Paranasal mucoceles in children with cystic fibrosis: Management of a not so rare clinical condition. American Journal of Otolaryngology - Head and Neck Medicine and Surgery, 2021, 42, 103107.	0.6	1
23	Case Report: The JAK-Inhibitor Ruxolitinib Use in Aicardi-Goutieres Syndrome Due to ADAR1 Mutation. Frontiers in Pediatrics, 2021, 9, 725868.	0.9	9
24	Factors Associated With Severe Gastrointestinal Diagnoses in Children With SARS-CoV-2 Infection or Multisystem Inflammatory Syndrome. JAMA Network Open, 2021, 4, e2139974.	2.8	24
25	Novel and emerging treatments for Aicardi-GoutiÃ¨res syndrome. Expert Review of Clinical Immunology, 2020, 16, 189-198.	1.3	27
26	More than an â€˜atypicalâ€™ phenotype: dual molecular diagnosis of autoimmune lymphoproliferative syndrome and Becker muscular dystrophy. British Journal of Haematology, 2020, 191, 291-294.	1.2	4
27	Activated Phosphoinositide 3-Kinase Delta Syndrome 1: Clinical and Immunological Data from an Italian Cohort of Patients. Journal of Clinical Medicine, 2020, 9, 3335.	1.0	23
28	Paediatric MAS/HLH caused by a novel monoallelic activating mutation in p110Î´. Clinical Immunology, 2020, 219, 108543.	1.4	8
29	The Italian Registry for Primary Immunodeficiencies (Italian Primary Immunodeficiency Network;) Tj ETQq1 1 0.784314 rgBT /Overlock 1	2.0	15
30	Heightened Circulating Interferon-Inducible Chemokines, and Activated Pro-Cytolytic Th1-Cell Phenotype Features Covid-19 Aggravation in the Second Week of Illness. Frontiers in Immunology, 2020, 11, 580987.	2.2	46
31	Successful Hematopoietic Stem Cell Transplantation in a Patient with Complete IFN-Î³ Receptor 2 Deficiency: a Case Report and Literature Review. Journal of Clinical Immunology, 2020, 40, 1191-1195.	2.0	7
32	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.	1.5	12
33	Immunologic characterization of a immunosuppressed multiple sclerosis patient that recovered from SARS-CoV-2 infection. Journal of Neuroimmunology, 2020, 345, 577282.	1.1	20
34	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.	1.5	59
35	Transient Decrease of Circulating and Tissular Dendritic Cells in Patients With Mycobacterial Disease and With Partial Dominant IFNÎ³R1 Deficiency. Frontiers in Immunology, 2020, 11, 1161.	2.2	5
36	Prevalence of Immunological Defects in a Cohort of 97 Rubinsteinâ€“Taybi Syndrome Patients. Journal of Clinical Immunology, 2020, 40, 851-860.	2.0	19

#	Article	IF	Citations
37	Immune response in children with COVIDâ€19 is characterized by lower levels of Tâ€cell activation than infected adults. European Journal of Immunology, 2020, 50, 1412-1414.	1.6	40
38	Mild encephalitis/encephalopathy with reversible splenial lesion (MERS) in twin sisters with two CD36 frameshift mutations. Neurological Sciences, 2020, 41, 2271-2274.	0.9	2
39	Two Xâ€linked agammaglobulinemia patients develop pneumonia as COVIDâ€19 manifestation but recover. Pediatric Allergy and Immunology, 2020, 31, 565-569.	1.1	298
40	Evaluation of a Diagnostic Therapeutic Educational Pathway for Asthma Management in Children and Adolescents. Frontiers in Pediatrics, 2020, 8, 39.	0.9	5
41	Asymptomatic case of Covid-19 in an infant with cystic fibrosis. Journal of Cystic Fibrosis, 2020, 19, e18.	0.3	29
42	Two X-linked agammaglobulinemia patients develop pneumonia as COVID-19 manifestation but recover. , 2020, 31, 565.		1
43	Multicentre Italian study of SARS-CoV-2 infection in children and adolescents, preliminary data as at 10 April 2020. Eurosurveillance, 2020, 25, .	3.9	222
44	From Natural Killer Cell Receptor Discovery to Characterization of Natural Killer Cell Defects in Primary Immunodeficiencies. Frontiers in Immunology, 2019, 10, 1757.	2.2	2
45	Hermansky-Pudlak syndrome type II and lethal hemophagocytic lymphohistiocytosis: Case description and review of the literature. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2476-2478.e5.	2.0	15
46	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.	2.0	40
47	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.	2.0	10
48	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.	2.2	41
49	Tregs and Th17 lymphocytes in human DYRK1A haploinsufficiency. Immunology Letters, 2019, 214, 52-54.	1.1	1
50	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.	2.0	27
51	The hidden hypothesis: A disseminated tuberculosis case. International Journal of Infectious Diseases, 2019, 85, 88-91.	1.5	0
52	Multisystem autoimmune disease caused by increased STAT3 phosphorylation and dysregulated gene expression. Haematologica, 2019, 104, e322-e325.	1.7	15
53	Cerebellar involvement in warts Hypogammaglobulinemia immunodeficiency myelokathexis patients: neuroimaging and clinical findings. Orphanet Journal of Rare Diseases, 2019, 14, 61.	1.2	4
54	Progressive severe B cell and NK cell deficiency with T cell senescence in adult CD40L deficiency. Clinical Immunology, 2018, 190, 11-14.	1.4	14

#	Article	IF	Citations
55	Vaccination in immunocompromised host: Recommendations of Italian Primary Immunodeficiency Network Centers (IPINET). Vaccine, 2018, 36, 3541-3554.	1.7	29
56	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.	1.4	33
57	Perinatally HIV-Infected Youths After Transition from Pediatric to Adult Care, a Single-Center Experience from Northern Italy. AIDS Research and Human Retroviruses, 2018, 34, 241-243.	0.5	12
58	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.	1.4	24
59	Metagenomic Discovery of 83 New Human Papillomavirus Types in Patients with Immunodeficiency. MSphere, 2018, 3, .	1.3	75
60	Strategies for Prevention of Mother-to-Child Transmission Adopted in the â€œReal-Worldâ€•Setting: Data From the Italian Register for HIV-1 Infection in Children. Journal of Acquired Immune Deficiency Syndromes (1999), 2018, 79, 54-61.	0.9	3
61	Sine causa tetraparesis. Medicine (United States), 2018, 97, e13893.	0.4	9
62	Heterozygous Mutation in Adenosine Deaminase Gene in a Patient With Severe Lymphopenia Following Corticosteroid Treatment of Autoimmune Hemolytic Anemia. Frontiers in Pediatrics, 2018, 6, 272.	0.9	2
63	A novel <scp>EP</scp>300 mutation associated with Rubinsteinâ€Taybi syndrome type 2 presenting as combined immunodeficiency. Pediatric Allergy and Immunology, 2018, 29, 776-781.	1.1	4
64	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.	1.5	58
65	Long-Term Survival After Hematopoietic Stem Cell Transplantation for Complete STAT1 Deficiency. Journal of Clinical Immunology, 2017, 37, 701-706.	2.0	21
66	Signal transducer and activator of transcription gain-of-function primary immunodeficiency/immunodysregulation disorders. Current Opinion in Pediatrics, 2017, 29, 711-717.	1.0	31
67	STAT mutations as program switchers: turning primary immunodeficiencies into autoimmune diseases. Journal of Leukocyte Biology, 2017, 101, 29-38.	1.5	59
68	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.	1.5	30
69	CXCL12 Mediates Aberrant Costimulation of B Lymphocytes in Warts, Hypogammaglobulinemia, Infections, Myelokathexis Immunodeficiency. Frontiers in Immunology, 2017, 8, 1068.	2.2	13
70	How I treat warts, hypogammaglobulinemia, infections, and myelokathexis syndrome. Blood, 2017, 130, 2491-2498.	0.6	46
71	Diagnostics of Primary Immunodeficiencies through Next-Generation Sequencing. Frontiers in Immunology, 2016, 7, 466.	2.2	80
72	Novel <scp>STAT</scp>1 gainâ€ofâ€function mutation and suppurative infections. Pediatric Allergy and Immunology, 2016, 27, 220-223.	1.1	14

#	Article	IF	Citations
73	Impairment of dendritic cell functions in patients with adaptor protein-3 complex deficiency. Blood, 2016, 127, 3382-3386.	0.6	11
74	Atypical presentation of autoimmune lymphoproliferative syndrome due to CASP10 mutation. Immunology Letters, 2016, 177, 22-24.	1.1	14
75	Clinical and immunological data of nine patients with chronic mucocutaneous candidiasis disease. Data in Brief, 2016, 7, 311-315.	0.5	10
76	Clinical heterogeneity of dominant chronic mucocutaneous candidiasis disease: presenting as treatment-resistant candidiasis and chronic lung disease. Clinical Immunology, 2016, 164, 1-9.	1.4	27
77	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.	1.5	16
78	Interferon-Î± Production by Plasmacytoid Dendritic Cells Is Dispensable for an Effective Anti-Cytomegalovirus Response in Adaptor Protein-3-Deficient Mice. Journal of Interferon and Cytokine Research, 2015, 35, 232-238.	0.5	4
79	Failure of interferon-Î³ pre-treated mesenchymal stem cell treatment in a patient with Crohnâ€™s disease. World Journal of Gastroenterology, 2015, 21, 4379.	1.4	21
80	Primary immunodeficiences â€“ options for the future. Pediatric Allergy and Immunology, 2014, 25, 27-29.	1.1	1
81	Bruton tyrosine kinase mediates TLR9-dependent human dendritic cell activation. Journal of Allergy and Clinical Immunology, 2014, 133, 1644-1650.e4.	1.5	62
82	Primary immunodeficiencies appearing as combined lymphopenia, neutropenia, and monocytopenia. Immunology Letters, 2014, 161, 222-225.	1.1	15
83	Activin A as a Mediator of NKâ€“Dendritic Cell Functional Interactions. Journal of Immunology, 2014, 192, 1241-1248.	0.4	27
84	Autosomal Recessive Agammaglobulinemia: The Third Case of IgÎ² Deficiency Due to a Novel Non-sense Mutation. Journal of Clinical Immunology, 2014, 34, 425-427.	2.0	11
85	Expansion of CCR4+ activated T cells is associated with memory B cell reduction in DOCK8-deficient patients. Clinical Immunology, 2014, 152, 164-170.	1.4	11
86	Î±- and Î²-Papillomavirus infection in a young patient with an unclassified primary T-cell immunodeficiency and multiple mucosal and cutaneous lesions. Journal of the American Academy of Dermatology, 2014, 71, 108-115.e1.	0.6	22
87	Combined DOCK8 and CLEC7A mutations causing immunodeficiency in 3 brothers with diarrhea, eczema, and infections. Journal of Allergy and Clinical Immunology, 2013, 131, 594-597.e3.	1.5	22
88	Clinical, laboratory and molecular signs of immunodeficiency in patients with partial oculo-cutaneous albinism. Orphanet Journal of Rare Diseases, 2013, 8, 168.	1.2	70
89	Disseminated Mycobacterium genavense infection after immunosuppressive therapy shows underlying new composite heterozygous mutations of Î²1 subunit of IL-12 receptor gene. Journal of Allergy and Clinical Immunology, 2013, 131, 607-610.	1.5	8
90	Defects of leukocyte migration in primary immunodeficiencies. European Journal of Immunology, 2013, 43, 1436-1440.	1.6	27

#	Article	IF	Citations
91	The CXCR4 mutations in WHIM syndrome impair the stability of the T-cell immunologic synapse. Blood, 2013, 122, 666-673.	0.6	59
92	Occurrence of Nodular Lymphocyte-Predominant Hodgkin Lymphoma in Hermansky-Pudlak Type 2 Syndrome Is Associated to Natural Killer and Natural Killer T Cell Defects. PLoS ONE, 2013, 8, e80131.	1.1	34
93	From Bone Marrow Transplantation to Cellular Therapies: Possible Therapeutic Strategies in Managing Autoimmune Disorders. Current Pharmaceutical Design, 2012, 18, 5776-5781.	0.9	5
94	Tetralogy of Fallot is an Uncommon Manifestation of Warts, Hypogammaglobulinemia, Infections, and Myelokathexis Syndrome. Journal of Pediatrics, 2012, 161, 763-765.	0.9	37
95	A novel compound heterozygous TACI mutation in an autosomal recessive common variable immunodeficiency (CVID) family. Human Immunology, 2012, 73, 836-839.	1.2	16
96	B Cell Responses to CpG Correlate with CXCL16 Expression Levels in Common Variable Immunodeficiency. Scientific World Journal, The, 2012, 2012, 1-9.	0.8	5
97	Exome sequencing reveals a pallidin mutation in a Hermansky-Pudlakâ€“like primary immunodeficiency syndrome. Blood, 2012, 119, 3185-3187.	0.6	76
98	Antiretroviral use in Italian children with perinatal HIV infection over a 14â€year period. Acta Paediatrica, International Journal of Paediatrics, 2012, 101, e287-95.	0.7	5
99	First-line antiretroviral therapy with a protease inhibitor versus non-nucleoside reverse transcriptase inhibitor and switch at higher versus low viral load in HIV-infected children: an open-label, randomised phase 2/3 trial. Lancet Infectious Diseases, The, 2011, 11, 273-283.	4.6	123
100	Psychosocial Issues in Children and Adolescents With HIV Infection Evaluated With a World Health Organization Age-Specific Descriptor System. Journal of Developmental and Behavioral Pediatrics, 2011, 32, 52-55.	0.6	13
101	Clinical heterogeneity and diagnostic delay of autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy syndrome. Clinical Immunology, 2011, 139, 6-11.	1.4	49
102	Severe impairment of IFN-Î³ and IFN-Î± responses in cells of a patient with a novel STAT1 splicing mutation. Blood, 2011, 118, 1806-1817.	0.6	84
103	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.	1.6	26
104	Clinical and Genetic Features of Warts, Hypogammaglobulinemia, Infections and Myelokathexis (WHIM) Syndrome. Current Molecular Medicine, 2011, 11, 317-325.	0.6	74
105	Risk of Perinatal HIV Infection in Infants Born in Italy to Immigrant Mothers. Clinical Infectious Diseases, 2011, 53, 310-313.	2.9	22
106	Patient-centred screening for primary immunodeficiency, a multi-stage diagnostic protocol designed for non-immunologists: 2011 update. Clinical and Experimental Immunology, 2011, 167, 108-119.	1.1	143
107	Defect of plasmacytoid dendritic cells in warts, hypogammaglobulinemia, infections, myelokathexis (WHIM) syndrome patients. Blood, 2010, 116, 4870-4873.	0.6	59
108	Italian consensus statement on paediatric HIV infection. Infection, 2010, 38, 301-319.	2.3	8

#	Article	IF	Citations
109	PTX3 genetic variations affect the risk of Pseudomonas aeruginosa airway colonization in cystic fibrosis patients. Genes and Immunity, 2010, 11, 665-670.	2.2	81
110	Uncovering an ILâ€10â€dependent NFâ€KB recruitment to the ILâ€1ra promoter that is impaired in STAT3 functionally defective patients. FASEB Journal, 2010, 24, 1365-1375.	0.2	45
111	Defect of regulatory T cells in patients with Omenn syndrome. Journal of Allergy and Clinical Immunology, 2010, 125, 209-216.	1.5	83
112	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.	0.6	37
113	Leukocyte trafficking in primary immunodeficiencies. Journal of Leukocyte Biology, 2009, 85, 335-343.	1.5	28
114	Is the Interruption of Antiretroviral Treatment During Pregnancy an Additional Major Risk Factor for Motherâ€toâ€Child Transmission of HIV Type 1?. Clinical Infectious Diseases, 2009, 48, 1310-1317.	2.9	25
115	OR.67. Lack of iNKT Cells and Defects on Differentiation of Dendritic Cells in Patients with Hermansky-Pudlak Type 2 Syndrome (HPS2). Clinical Immunology, 2009, 131, S28.	1.4	0
116	Five-year follow-up of children with perinatal HIV-1 infection receiving early highly active antiretroviral therapy. BMC Infectious Diseases, 2009, 9, 140.	1.3	41
117	Clinical and genetic diagnosis of warts, hypogammaglobulinemia, infections, and myelokathexis syndrome in 10 patients. Journal of Allergy and Clinical Immunology, 2009, 123, 1170-1173.e3.	1.5	52
118	Type I interferon-dependent gene MxA in perinatal HIV-infected patients under antiretroviral therapy as marker for therapy failure and blood plasmacytoid dendritic cells depletion. Journal of Translational Medicine, 2008, 6, 49.	1.8	10
119	Decreased Type I Interferon Receptor-Soluble Isoform in Antiretroviral-Treated HIV-Positive Children. Journal of Interferon and Cytokine Research, 2008, 28, 181-189.	0.5	6
120	Editorial Commentary:Immunological Nonresponse to Highly Active Antiretroviral Therapy in HIVâ€Infected Subjects: Is the Bone Marrow Impairment Causing CD4 Lymphopenia?. Clinical Infectious Diseases, 2008, 46, 1911-1912.	2.9	9
121	Epigenetic Regulation of Protein-Coding and MicroRNA Genes by the Gfi1-Interacting Tumor Suppressor PRDM5. Molecular and Cellular Biology, 2007, 27, 6889-6902.	1.1	79
122	G-CSF treatment of severe congenital neutropenia reverses neutropenia but does not correct the underlying functional deficiency of the neutrophil in defending against microorganisms. Blood, 2007, 109, 4716-4723.	0.6	80
123	Response: ELA2 genotype-phenotype correlation to be expected in patients with severe congenital neutropenia. Blood, 2007, 110, 2773-2774.	0.6	0
124	Novel insights from adaptor protein 3 complexÂdeficiency. Journal of Allergy and Clinical Immunology, 2007, 120, 735-741.	1.5	51
125	A robust immunoassay for anti-interferon autoantibodies that is highly specific for patients with autoimmune polyglandular syndrome type 1. Clinical Immunology, 2007, 125, 131-137.	1.4	43
126	Role of dendritic cell-derived CXCL13 in the pathogenesis of Bartonella henselae B-rich granuloma. Blood, 2006, 107, 454-462.	0.6	65

#	Article	IF	Citations
127	Innate immunity defects in Hermansky-Pudlak type 2 syndrome. Blood, 2006, 107, 4857-4864.	0.6	136
128	Combined decrease of defined B and T cell subsets in a group of common variable immunodeficiency patients. Clinical Immunology, 2006, 121, 203-214.	1.4	52
129	Virologic, immunologic, and clinical benefits from early combined antiretroviral therapy in infants with perinatal HIV-1 infection. Aids, 2006, 20, 1789.	1.0	0
130	Virologic, immunologic, and clinical benefits from early combined antiretroviral therapy in infants with perinatal HIV-1 infection.. Aids, 2006, 20, 207-215.	1.0	88
131	Immunodeficiencies with Autoimmune Consequences. Advances in Immunology, 2006, 89, 321-370.	1.1	64
132	Neutropenia-Associated Mutations of PFAAP5, a Novel Protein Mediating Transcriptional Repressor Interaction between Gfi1 and Neutrophil Elastase.. Blood, 2006, 108, 501-501.	0.6	0
133	Genotyping for Guiding Drug Choice in Human Immunodeficiency Virus-Infected Children Failing Multiple Antiretroviral Treatment Regimens. Pediatric Infectious Disease Journal, 2005, 24, 747-749.	1.1	8
134	A new case of IPEX receiving bone marrow transplantation. Bone Marrow Transplantation, 2005, 35, 1033-1034.	1.3	66
135	Hyper immunoglobulin M syndrome due to CD40 deficiency: clinical, molecular, and immunological features. Immunological Reviews, 2005, 203, 48-66.	2.8	176
136	IL-10 Enhances CCL2 Release and Chemotaxis Induced by CCL16 in Human Monocytes. International Journal of Immunopathology and Pharmacology, 2005, 18, 339-349.	1.0	28
137	Mechanisms of WHIM syndrome. Drug Discovery Today Disease Mechanisms, 2005, 2, 479-485.	0.8	1
138	AIRE deficiency in thymus of 2 patients with Omenn syndrome. Journal of Clinical Investigation, 2005, 115, 728-732.	3.9	146
139	AIRE deficiency in thymus of 2 patients with Omenn syndrome. Journal of Clinical Investigation, 2005, 115, 728-732.	3.9	69
140	Toll Receptor-Mediated Regulation of NADPH Oxidase in Human Dendritic Cells. Journal of Immunology, 2004, 173, 5749-5756.	0.4	131
141	Leukocyte circulation: one-way or round-trip? Lessons from primary immunodeficiency patients. Journal of Leukocyte Biology, 2004, 76, 1-6.	1.5	29
142	Toll-like receptor-4 genotype in children with respiratory infections. Allergy: European Journal of Allergy and Clinical Immunology, 2004, 59, 1018-1019.	2.7	5
143	Congenital neutropenia: advances in diagnosis and treatment. Current Opinion in Allergy and Clinical Immunology, 2004, 4, 513-521.	1.1	39
144	Altered leukocyte response to CXCL12 in patients with warts hypogammaglobulinemia, infections, myelokathexis (WHIM) syndrome. Blood, 2004, 104, 444-452.	0.6	172

#	Article	IF	Citations
145	Autoimmune Polyendocrinopathy-Candidiasis-Ectodermal Dystrophy Syndrome: Time to Review Diagnostic Criteria?. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 3146-3148.	1.8	75
146	Development of systemic lupus erythematosus in a young child affected with chronic granulomatous disease following withdrawal of treatment with interferon-gamma. British Journal of Rheumatology, 2003, 42, 804-805.	2.5	20
147	Functional defects of dendritic cells in patients with CD40 deficiency. Blood, 2003, 102, 4099-4106.	0.6	50
148	X-chromosome inactivation analysis in a female carrier of FOXP3 mutation. Clinical and Experimental Immunology, 2002, 130, 127-130.	1.1	88
149	Defective migration of monocyte-derived dendritic cells in LAD-1 immunodeficiency. Journal of Leukocyte Biology, 2002, 72, 650-6.	1.5	19
150	Monocytes from Wiskott-Aldrich patients differentiate in functional mature dendritic cells with a defect in CD83 expression. European Journal of Immunology, 2001, 31, 3413-3421.	1.6	23
151	Mutations of CD40 gene cause an autosomal recessive form of immunodeficiency with hyper IgM. Proceedings of the National Academy of Sciences of the United States of America, 2001, 98, 12614-12619.	3.3	347
152	Interaction of Bartonella henselae with the Murine Macrophage Cell Line J774: Infection and Proinflammatory Response. Infection and Immunity, 2001, 69, 5974-5980.	1.0	51
153	Serum amyloid A is an activator of PMN antimicrobial functions: induction of degranulation, phagocytosis, and enhancement of anti-Candida activity. Journal of Leukocyte Biology, 2000, 67, 381-386.	1.5	79
154	Complete genomic organization of the human JAK3 gene and mutation analysis in severe combined immunodeficiency by single-strand conformation polymorphism. Human Genetics, 2000, 106, 73-79.	1.8	38
155	Combined Immunodeficiencies Due to Defects in Signal Transduction: Defects of the Î³c-JAK3 Signaling Pathway as a Model. Immunobiology, 2000, 202, 106-119.	0.8	28
156	New insights into the biology of the acute phase response. Journal of Clinical Immunology, 1999, 19, 203-214.	2.0	329
157	Prenatal molecular diagnosis of Wiskott-Aldrich syndrome by direct mutation analysis. , 1999, 19, 36-40.		15
158	Expression of Inducible Nitric Oxide Synthase in Human Granulomas and Histiocytic Reactions. American Journal of Pathology, 1999, 154, 145-152.	1.9	108
159	Ureteral obstruction in a patient with chronic granulomatous disease, receiving combined prophylaxis with IFN-Î³ and antibiotics. European Journal of Pediatrics, 1998, 157, 352-353.	1.3	1
160	Molecular and biochemical characterization of JAK3 deficiency in a patient with severe combined immunodeficiency over 20 years after bone marrow transplantation: implications for treatment. British Journal of Haematology, 1998, 102, 1363-1366.	1.2	19
161	Interleukin-15 Activates Proinflammatory and Antimicrobial Functions in Polymorphonuclear Cells. Infection and Immunity, 1998, 66, 2640-2647.	1.0	76
162	Structural and Functional Basis for JAK3-Deficient Severe Combined Immunodeficiency. Blood, 1997, 90, 3996-4003.	0.6	138

#	Article	IF	Citations
163	Interleukin-15 (IL-15) Induces IL-8 and Monocyte Chemotactic Protein 1 Production in Human Monocytes. Blood, 1997, 90, 2804-2809.	0.6	118
164	Expression of growth hormone receptor by peripheral blood lymphocytes in children: evaluation in clinical conditions of impaired growth. Clinical Endocrinology, 1997, 47, 329-335.	1.2	23
165	Leishmania major:Infection of Human Monocytes Induces Expression of IL-8 and MCAF. Experimental Parasitology, 1996, 82, 21-26.	0.5	64
166	Role of cytokines, acute-phase proteins, and chemokinesin the progression of rheumatoid arthritis. Seminars in Arthritis and Rheumatism, 1996, 26, 526-538.	1.6	145
167	Leishmania promastigotes selectively inhibit interleukin 12 induction in bone marrow-derived macrophages from susceptible and resistant mice.. Journal of Experimental Medicine, 1996, 183, 515-526.	4.2	318
168	Serum amyloid A is a chemoattractant: induction of migration, adhesion, and tissue infiltration of monocytes and polymorphonuclear leukocytes.. Journal of Experimental Medicine, 1994, 180, 203-209.	4.2	464
169	Differential expression of surface membrane growth hormone receptor on human peripheral blood lymphocytes detected by dual fluorochrome flow cytometry. Journal of Clinical Endocrinology and Metabolism, 1994, 79, 984-990.	1.8	59