Peter Arkwright

List of Publications by Year in descending order

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166 papers 12,920 citations

64 h-index 24961 109 g-index

170 all docs

170 docs citations

times ranked

170

14466 citing authors

#	Article	IF	CITATIONS
1	Surveillance and control of meningococcal disease in the COVID-19 era: A Global Meningococcal Initiative review. Journal of Infection, 2022, 84, 289-296.	1.7	26
2	An Integrated Taxonomy for Monogenic Inflammatory Bowel Disease. Gastroenterology, 2022, 162, 859-876.	0.6	37
3	Striving for Evidence-Based Management of Food Allergies. Journal of Allergy and Clinical Immunology: in Practice, 2022, 10, 56-58.	2.0	1
4	Clinical Outcome and Underlying Genetic Cause of Functional Terminal Complement Pathway Deficiencies in a Multicenter UK Cohort. Journal of Clinical Immunology, 2022, , 1.	2.0	1
5	Keratinocyte <scp>EGF</scp> signaling dominates in Atopic Dermatitis lesions: a comparative <scp>RNAseq</scp> analysis. Experimental Dermatology, 2022, , .	1.4	2
6	Neutrophil dysfunction triggers inflammatory bowel disease in G6PC3 deficiency. Journal of Leukocyte Biology, 2021, 109, 1147-1154.	1.5	14
7	Introducing a New Epoch in Inborn Errors of Immunity. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 660-662.	2.0	1
8	Staphylococcus aureus second immunoglobulin-binding protein drives atopic dermatitis via IL-33. Journal of Allergy and Clinical Immunology, 2021, 147, 1354-1368.e3.	1.5	35
9	Dupilumab provides favourable longâ€ŧerm safety and efficacy in children aged ≥ 6 to < 12 years with uncontrolled severe atopic dermatitis: results from an open″abel phase lla study and subsequent phase lll open″abel extension study. British Journal of Dermatology, 2021, 184, 857-870.	1.4	45
10	TCF3 Dominant Negative Variant Causes an Early Block in B-Lymphopoiesis and Agammaglobulinemia. Journal of Clinical Immunology, 2021, 41, 1391-1394.	2.0	6
11	The Risk of Allergic Reaction to SARS-CoV-2 Vaccines and Recommended Evaluation and Management: A Systematic Review, Meta-Analysis, GRADE Assessment, and International Consensus Approach. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 3546-3567.	2.0	152
12	Genetic, Immunological, and Clinical Features of 32 Patients with Autosomal Recessive STAT1 Deficiency. Journal of Immunology, 2021, 207, 133-152.	0.4	33
13	Laboratory Safety of Dupilumab in Patients Aged 6–11 Years with Severe Atopic Dermatitis: Results from a Phase III Clinical Trial. Paediatric Drugs, 2021, 23, 515-527.	1.3	15
14	Simple Measurement of IgA Predicts Immunity and Mortality in Ataxia-Telangiectasia. Journal of Clinical Immunology, 2021, 41, 1878-1892.	2.0	9
15	Dupilumab in adolescents with uncontrolled moderateâ€toâ€severe atopic dermatitis: results from a phase <scp>ll</scp> openâ€label trial and subsequent phase <scp>lll</scp> openâ€label extension. British Journal of Dermatology, 2020, 182, 85-96.	1.4	111
16	Vertebral, pelvic, and hip fracture risk in adults with severe atopic dermatitis. Journal of Allergy and Clinical Immunology, 2020, 145, 487-488.	1.5	8
17	A Rapid Shift from Chronic Hyperoxia to Normoxia Induces Systemic Anaphylaxis via Transient Receptor Potential Ankyrin 1 Channels on Mast Cells. Journal of Immunology, 2020, 205, 2959-2967.	0.4	7
18	Antibody persistence following meningococcal ACWY conjugate vaccine licensed in the European Union by age group and vaccine. Expert Review of Vaccines, 2020, 19, 745-754.	2.0	2

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19	Insulin hypersensitivity in type 1 diabetes: investigation and treatment with immunodepletion. Practical Diabetes, 2020, 37, 59.	0.1	4
20	Genomic profiling of acute myeloid leukaemia associated with ataxia telangiectasia identifies a complex karyotype with wildâ€type ⟨i>TP53⟨/i> and mutant ⟨i>KRAS, G3BP1⟨/i> and ⟨i>IL7R⟨/i>. Pediatric Blood and Cancer, 2020, 67, e28354.	0.8	4
21	Hereditary Alpha-Tryptasemia: UK Prevalence and Variability in Disease Expression. Journal of Allergy and Clinical Immunology: in Practice, 2020, 8, 3549-3556.	2.0	70
22	Efficacy and safety of dupilumab with concomitant topical corticosteroids in children 6 to 11Âyears old with severe atopic dermatitis: A randomized, double-blinded, placebo-controlled phase 3 trial. Journal of the American Academy of Dermatology, 2020, 83, 1282-1293.	0.6	214
23	Impact on quality of life and safety of sublingual and subcutaneous immunotherapy in children with severe house dust mite and pollen-associated allergic rhinoconjunctivitis. Clinical and Translational Allergy, 2020, 10, 10.	1.4	10
24	Type 2 immunity in the skin and lungs. Allergy: European Journal of Allergy and Clinical Immunology, 2020, 75, 1582-1605.	2.7	304
25	Infant Alveolar Macrophages Are Unable to Effectively Contain Mycobacterium tuberculosis. Frontiers in Immunology, 2020, 11, 486.	2.2	15
26	Impaired lymphocyte function and differentiation in CTPS1-deficient patients result from a hypomorphic homozygous mutation. JCI Insight, 2020, 5, .	2.3	29
27	Refractory very early-onset inflammatory bowel disease associated with cytosolic isoleucyl-tRNA synthetase deficiency: A case report. World Journal of Gastroenterology, 2020, 26, 1841-1846.	1.4	6
28	Immune Response and Safety of Viral Vaccines in Children with Autoimmune Diseases on Immune Modulatory Drug Therapy. Expert Review of Vaccines, 2020, 19, 1115-1127.	2.0	12
29	Fennel as a cause of immediate hypersensitivity to toothpaste. Annals of Allergy, Asthma and Immunology, 2020, 125, 99-100.	0.5	2
30	Hematopoietic stem cell transplantation for cytidine triphosphate synthase 1 (CTPS1) deficiency. Bone Marrow Transplantation, 2019, 54, 130-133.	1.3	13
31	Life-threatening pulmonary interstitial lung disease complicating pediatric nonhumoral immunodeficiencies. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 2456-2458.e4.	2.0	O
32	Immunodeficiency, autoimmune thrombocytopenia and enterocolitis caused by autosomal recessive deficiency of <i>PIK3CD</i> -encoded phosphoinositide 3-kinase δ. Haematologica, 2019, 104, e483-e486.	1.7	26
33	Mast cell disorders: From infancy to maturity. Allergy: European Journal of Allergy and Clinical Immunology, 2019, 74, 53-63.	2.7	44
34	Hypomorphic caspase activation and recruitment domain 11 (CARD11) mutations associated with diverse immunologic phenotypes with or without atopic disease. Journal of Allergy and Clinical Immunology, 2019, 143, 1482-1495.	1.5	116
35	Hematopoietic stem cell transplant effectively rescues lymphocyte differentiation and function in DOCK8-deficient patients. JCl Insight, 2019, 4, .	2.3	23
36	Allergy testing in predicting outcome of open food challenge to peanut. Journal of Allergy and Clinical Immunology, 2018, 141, 457-458.	1.5	10

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37	Bronchiectasis and deteriorating lung function in agammaglobulinaemia despite immunoglobulin replacement therapy. Clinical and Experimental Immunology, 2018, 191, 212-219.	1.1	30
38	Home-based program of maintaining unresponsiveness in children with allergic reactions to larger amounts of peanuts. Annals of Allergy, Asthma and Immunology, 2018, 120, 539-540.	0.5	7
39	Increased Risk for Malignancies in 131 Affected CTLA4 Mutation Carriers. Frontiers in Immunology, 2018, 9, 2012.	2.2	79
40	The United Kingdom Primary Immune Deficiency (UKPID) registry 2012 to 2017. Clinical and Experimental Immunology, 2018, 192, 284-291.	1.1	57
41	Phenotype, penetrance, and treatment of 133 cytotoxic T-lymphocyte antigen 4–insufficient subjects. Journal of Allergy and Clinical Immunology, 2018, 142, 1932-1946.	1.5	344
42	Severity and threshold of peanut reactivity during hospitalâ€based open oral food challenges: An international multicenter survey. Pediatric Allergy and Immunology, 2018, 29, 754-761.	1.1	34
43	Biallelic <i>RIPK1</i> mutations in humans cause severe immunodeficiency, arthritis, and intestinal inflammation. Science, 2018, 361, 810-813.	6.0	181
44	Clinical and laboratory features of seventy-eight UK patients with Good's syndrome (thymoma and) Tj ETQq	0 0 0 rgBT	/Oygrlock 10
45	Inherited p40phox deficiency differs from classic chronic granulomatous disease. Journal of Clinical Investigation, 2018, 128, 3957-3975.	3.9	99
46	Factors that determine parents' perception of their child's risk of life-threatening food-induced anaphylaxis. Allergy and Asthma Proceedings, 2017, 38, 44-53.	1.0	14
47	Human Adaptive Immunity Rescues an Inborn Error of Innate Immunity. Cell, 2017, 168, 789-800.e10.	13.5	68
48	Identifying functional defects in patients with immune dysregulation due to LRBA and CTLA-4 mutations. Blood, 2017, 129, 1458-1468.	0.6	102
49	Chromatin-remodeling factor SMARCD2 regulates transcriptional networks controlling differentiation of neutrophil granulocytes. Nature Genetics, 2017, 49, 742-752.	9.4	87
50	Dedicator of cytokinesis 8–deficient CD4 + TÂcells are biased to a T H 2 effector fate at the expense of T H 1 and T H 17Âcells. Journal of Allergy and Clinical Immunology, 2017, 139, 933-949.	1.5	69
51	14 Years after Discovery: Clinical Follow-up on 15 Patients with Inducible Co-Stimulator Deficiency. Frontiers in Immunology, 2017, 8, 964.	2.2	57
52	Mast cell hyperactivity underpins the development of oxygen-induced retinopathy. Journal of Clinical Investigation, 2017, 127, 3987-4000.	3.9	24
53	Unique and shared signaling pathways cooperate to regulate the differentiation of human CD4+ T cells into distinct effector subsets. Journal of Experimental Medicine, 2016, 213, 1589-1608.	4.2	77
54	Hematopoietic stem cell transplantation for CTLA4 deficiency. Journal of Allergy and Clinical Immunology, 2016, 138, 615-619.e1.	1.5	88

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55	Bone marrow transplantation for MHC class I deficiency corrects T-cell immunity but dissociates natural killer cell repertoire formation from function. Journal of Allergy and Clinical Immunology, 2016, 138, 1733-1736.e2.	1.5	7
56	Hyperactive mTOR pathway promotes lymphoproliferation and abnormal differentiation in autoimmune lymphoproliferative syndrome. Blood, 2016, 128, 227-238.	0.6	77
57	Elevated basal serum tryptase identifies a multisystem disorder associated with increased TPSAB1 copy number. Nature Genetics, 2016, 48, 1564-1569.	9.4	279
58	Clinical and laboratory correlates of lung disease and cancer in adults with idiopathic hypogammaglobulinaemia. Clinical and Experimental Immunology, 2016, 184, 73-82.	1.1	24
59	Whole-exome sequencing to analyze population structure, parental inbreeding, and familial linkage. Proceedings of the National Academy of Sciences of the United States of America, 2016, 113, 6713-6718.	3.3	53
60	Defective Leukocyte Adhesion and Chemotaxis Contributes to Combined Immunodeficiency in Humans with Autosomal Recessive MST1 Deficiency. Journal of Clinical Immunology, 2016, 36, 117-122.	2.0	63
61	AIRE is not essential for the induction of human tolerogenic dendritic cells. Autoimmunity, 2016, 49, 211-218.	1.2	2
62	Skin pH Is the Master Switch of Kallikrein 5-Mediated Skin Barrier Destruction in a Murine Atopic Dermatitis Model. Journal of Investigative Dermatology, 2016, 136, 127-135.	0.3	92
63	Reply. Journal of Allergy and Clinical Immunology: in Practice, 2015, 3, 828-829.	2.0	0
64	The extended clinical phenotype of 64 patients with dedicator of cytokinesis 8 deficiency. Journal of Allergy and Clinical Immunology, 2015, 136, 402-412.	1.5	163
65	Spectrum and Management of Complement Immunodeficiencies (Excluding Hereditary Angioedema) Across Europe. Journal of Clinical Immunology, 2015, 35, 199-205.	2.0	40
66	Multicenter experience in hematopoietic stem cell transplantation for serious complications of common variable immunodeficiency. Journal of Allergy and Clinical Immunology, 2015, 135, 988-997.e6.	1.5	123
67	Monogenic mutations differentially affect the quantity and quality of T follicular helper cells in patients with human primary immunodeficiencies. Journal of Allergy and Clinical Immunology, 2015, 136, 993-1006.e1.	1.5	181
68	Impairment of immunity to <i>Candida</i> and <i>Mycobacterium</i> in humans with bi-allelic <i>RORC</i> mutations. Science, 2015, 349, 606-613.	6.0	366
69	The value of microarray-based comparative genomic hybridisation (aCGH) testing in the paediatric clinic. Archives of Disease in Childhood, 2015, 100, 728-731.	1.0	4
70	Marked variability in clinical presentation and outcome of patients with C1q immunodeficiency. Journal of Autoimmunity, 2015, 62, 39-44.	3.0	33
71	The deep fascia of the thigh forms an impenetrable barrier to fluid injected subcutaneously by autoinjectors. Journal of Allergy and Clinical Immunology: in Practice, 2015, 3, 297-299.	2.0	20
72	STAT3 is a critical cell-intrinsic regulator of human unconventional T cell numbers and function. Journal of Experimental Medicine, 2015, 212, 855-864.	4.2	70

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73	Astute Clinician Report: A Novel 10Âbp Frameshift Deletion in Exon 2 of ICOS Causes a Combined Immunodeficiency Associated with an Enteritis and Hepatitis. Journal of Clinical Immunology, 2015, 35, 598-603.	2.0	30
74	Hyperactive mTOR Pathway Promotes Lymphoproliferation and Abnormal Differentiation in Human Autoimmune Lymphoproliferative Syndrome. Blood, 2015, 126, 1020-1020.	0.6	1
75	Atopic Dermatitis in Children. Journal of Allergy and Clinical Immunology: in Practice, 2014, 2, 388-395.	2.0	13
76	CTP synthase 1 deficiency in humans reveals its central role in lymphocyte proliferation. Nature, 2014, 510, 288-292.	13.7	174
77	Successful cure of C1q deficiency in human subjects treated with hematopoietic stem cell transplantation. Journal of Allergy and Clinical Immunology, 2014, 133, 265-267.	1.5	69
78	Signal transducer and activator of transcription 3 (STAT3) mutations underlying autosomal dominant hyper-lgE syndrome impair human CD8+ T-cell memory formation and function. Journal of Allergy and Clinical Immunology, 2013, 132, 400-411.e9.	1.5	63
79	IL-21 signalling via STAT3 primes human na \tilde{A} ve B cells to respond to IL-2 to enhance their differentiation into plasmablasts. Blood, 2013, 122, 3940-3950.	0.6	121
80	Management of Difficult-to-Treat Atopic Dermatitis. Journal of Allergy and Clinical Immunology: in Practice, 2013, 1, 142-151.	2.0	143
81	G6PC3 mutations cause non-syndromic severe congenital neutropenia. Molecular Genetics and Metabolism, 2013, 108, 138-141.	0.5	16
82	Anatomical and Anthropometric Determinants of Intramuscular Versus Subcutaneous Administration in Children with Epinephrine Auto-Injectors. Journal of Allergy and Clinical Immunology, 2013, 131, AB199.	1.5	0
83	Anatomic and anthropometric determinants of intramuscular versus subcutaneous administration in children with epinephrine autoinjectors. Journal of Allergy and Clinical Immunology: in Practice, 2013, 1, 692-694.	2.0	12
84	lgE Sensitization to the Nonspecific Lipid-Transfer Protein Ara h 9 and Peanut-Associated Bronchospasm. BioMed Research International, 2013, 2013, 1-9.	0.9	10
85	Naive and memory human B cells have distinct requirements for STAT3 activation to differentiate into antibody-secreting plasma cells. Journal of Experimental Medicine, 2013, 210, 2739-2753.	4.2	158
86	DOCK8 is critical for the survival and function of NKT cells. Blood, 2013, 122, 2052-2061.	0.6	68
87	The United Kingdom Primary Immune Deficiency (UKPID) Registry: report of the first 4 years' activity 2008–2012. Clinical and Experimental Immunology, 2013, 175, 68-78.	1.1	85
88	Factors determining the effectiveness of oral ciclosporin in the treatment of severe childhood atopic dermatitis. Journal of Dermatological Treatment, 2012, 23, 318-322.	1.1	9
89	Functional STAT3 deficiency compromises the generation of human T follicular helper cells. Blood, 2012, 119, 3997-4008.	0.6	267
90	The use of adrenaline autoinjectors by children and teenagers. Clinical and Experimental Allergy, 2012, 42, 284-292.	1.4	116

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91	STAT1 Hyperphosphorylation and Defective IL12R/IL23R Signaling Underlie Defective Immunity in Autosomal Dominant Chronic Mucocutaneous Candidiasis. PLoS ONE, 2011, 6, e29248.	1.1	101
92	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.	0.6	268
93	Onset of autoimmune lymphoproliferative syndrome (ALPS) in humans as a consequence of genetic defect accumulation. Journal of Clinical Investigation, 2011, 121, 106-112.	3.9	110
94	Ten warning signs of primary immunodeficiency: a new paradigm is needed for the 21st century. Annals of the New York Academy of Sciences, 2011, 1238, 7-14.	1.8	102
95	Differential cytokine secretion results from p65 and c-Rel NF-κB subunit signaling in peripheral blood mononuclear cells of TNF receptor-associated periodic syndrome patients. Cellular Immunology, 2011, 268, 55-59.	1.4	24
96	Clinical Features That Identify Children With Primary Immunodeficiency Diseases. Pediatrics, 2011, 127, 810-816.	1.0	149
97	DOCK8 deficiency impairs CD8 T cell survival and function in humans and mice. Journal of Experimental Medicine, 2011, 208, 2305-2320.	4.2	175
98	Prevalence, outcome and pre-hospital management of anaphylaxis by first aiders and paramedical ambulance staff in Manchester, UK. Resuscitation, 2010, 81, 653-657.	1.3	39
99	Fas stimulation of T lymphocytes promotes rapid intercellular exchange of death signals via membrane nanotubes. Cell Research, 2010, 20, 72-88.	5.7	96
100	Autoimmunity and recurrent infections in partial complement C3 immunodeficiency. Rheumatology, 2010, 49, 1017-1019.	0.9	4
101	Autoantibodies against IL-17A, IL-17F, and IL-22 in patients with chronic mucocutaneous candidiasis and autoimmune polyendocrine syndrome type I. Journal of Experimental Medicine, 2010, 207, 291-297.	4.2	663
102	B cell–intrinsic signaling through IL-21 receptor and STAT3 is required for establishing long-lived antibody responses in humans. Journal of Experimental Medicine, 2010, 207, 155-171.	4.2	346
103	C1q Deficiency Leads to the Defective Suppression of IFN-α in Response to Nucleoprotein Containing Immune Complexes. Journal of Immunology, 2010, 185, 4738-4749.	0.4	190
104	Mutations in STAT3 and diagnostic guidelines for hyper-lgE syndrome. Journal of Allergy and Clinical Immunology, 2010, 125, 424-432.e8.	1.5	247
105	Impaired TH17 responses in patients with chronic mucocutaneous candidiasis with and without autoimmune polyendocrinopathy–candidiasis–ectodermal dystrophy. Journal of Allergy and Clinical Immunology, 2010, 126, 1006-1015.e4.	1.5	52
106	Clinical Features and Outcome of Patients With IRAK-4 and MyD88 Deficiency. Medicine (United States), 2010, 89, 403-425.	0.4	366
107	Effects of age, gender, and immunosuppressive agents on in vivo toll-like receptor pathway responses. Human Immunology, 2010, 71, 372-376.	1.2	19
108	Onset of Autoimmunity In ALPS as a Consequence of Genetic Defects Accumulation. Blood, 2010, 116, 278-278.	0.6	1

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109	Pattern recognition receptor expression is not impaired in patients with chronic mucocutanous candidiasis with or without autoimmune polyendocrinopathy candidiasis ectodermal dystrophy. Clinical and Experimental Immunology, 2009, 156, 40-51.	1.1	22
110	Automatic epinephrine device use in children with food allergies. Journal of Allergy and Clinical Immunology, 2009, 123, 267-268.	1.5	8
111	Anti-CD20 or anti-lgE therapy for severe chronic autoimmune urticaria. Journal of Allergy and Clinical Immunology, 2009, 123, 510-511.	1.5	52
112	FAS-L, IL-10, and double-negative CD4â^'CD8â^' TCR $\hat{l} \pm \hat{l}^2 + T$ cells are reliable markers of autoimmune lymphoproliferative syndrome (ALPS) associated with FAS loss of function. Blood, 2009, 113, 3027-3030.	0.6	134
113	Abnormal tumor necrosis factor receptor I cell surface expression and NF›B activation in tumor necrosis factor receptor–associated periodic syndrome. Arthritis and Rheumatism, 2008, 58, 273-283.	6.7	75
114	Impaired dendritic cell maturation and cytokine production in patients with chronic mucocutanous candidiasis with or without APECED. Clinical and Experimental Immunology, 2008, 154, 406-414.	1.1	48
115	Factors predicting anaphylaxis to peanuts and tree nuts in patients referred to a specialist center. Journal of Allergy and Clinical Immunology, 2008, 121, 632-638.e2.	1.5	161
116	Autoantibodies against Type I Interferons as an Additional Diagnostic Criterion for Autoimmune Polyendocrine Syndrome Type I. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 4389-4397.	1.8	176
117	Recently identified factors predisposing children to infectious diseases. Current Opinion in Infectious Diseases, 2008, 21, 217-222.	1.3	9
118	Mevalonic Aciduria Cured by Bone Marrow Transplantation. New England Journal of Medicine, 2007, 357, 1350-1350.	13.9	38
119	Clinical variability and characteristic autoantibody profile in primary C1q complement deficiency. Rheumatology, 2007, 46, 1612-1614.	0.9	31
120	The 23-valent pneumococcal polysaccharide vaccine does not provide additional serotype antibody protection in children who have been primed with two doses of heptavalent pneumococcal conjugate vaccine. Vaccine, 2007, 25, 6321-6325.	1.7	14
121	Blinded side-to-side comparison of topical corticosteroid and tacrolimus ointment in children with moderate to severe atopic dermatitis. Clinical and Experimental Dermatology, 2007, 32, 145-147.	0.6	11
122	Total and serotype-specific pneumococcal antibody titres in children with normal and abnormal humoral immunity. Vaccine, 2006, 24, 5637-5644.	1.7	34
123	Factors determining the ability of parents to effectively administer intramuscular adrenaline to food allergic children. Pediatric Allergy and Immunology, 2006, 17, 227-229.	1.1	84
124	Polymorphisms of the Bcl-2 family member bfl-1 in children with atopic dermatitis. Pediatric Allergy and Immunology, 2006, 17, 578-582.	1.1	6
125	Effect of childhood eczema and asthma on parental sleep and well-being: a prospective comparative study. British Journal of Dermatology, 2006, 154, 514-518.	1.4	120
126	Killed Mycobacterium vaccae suspension in children with moderate-to-severe atopic dermatitis: a randomized, double-blind, placebo-controlled trial. Clinical and Experimental Allergy, 2006, 36, 1115-1121.	1.4	35

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127	Memory switched B cell percentage and not serum immunoglobulin concentration is associated with clinical complications in children and adults with specific antibody deficiency and common variable immunodeficiency. Clinical Immunology, 2006, 120, 310-318.	1.4	106
128	Novel STAT1 Alleles in Otherwise Healthy Patients with Mycobacterial Disease. PLoS Genetics, 2006, 2, e131.	1.5	171
129	Human Complete Stat-1 Deficiency Is Associated with Defective Type I and II IFN Responses In Vitro but Immunity to Some Low Virulence Viruses In Vivo. Journal of Immunology, 2006, 176, 5078-5083.	0.4	191
130	Recurrent hemiplegia associated with cerebral vasculopathy following third trimester maternal herpes zoster infection. Developmental Medicine and Child Neurology, 2006, 48, 991.	1.1	15
131	Effect of Mycobacterium vaccae on cytokine responses in children with atopic dermatitis. Clinical and Experimental Immunology, 2005, 140, 101-108.	1.1	17
132	Mycobacterium vaccae Reduces Scratching Behavior but not the Rash in NC Mice with Eczema: A Randomized, Blinded, Placebo-Controlled Trial. Journal of Investigative Dermatology, 2005, 124, 140-143.	0.3	6
133	Highly glycosylated $\hat{l}\pm 1$ -acid glycoprotein is synthesized in myelocytes, stored in secondary granules, and released by activated neutrophils. Journal of Leukocyte Biology, 2005, 78, 462-470.	1.5	45
134	Treatment of Epstein-Barr-virus-associated primary CNS B cell lymphoma with allogeneic T-cell immunotherapy and stem-cell transplantation. Lancet Oncology, The, 2005, 6, 344-346.	5.1	51
135	CD4+CD25+ T-regulatory cells are decreased in patients with autoimmune polyendocrinopathy candidiasis ectodermal dystrophy. Journal of Allergy and Clinical Immunology, 2005, 116, 1158-1159.	1.5	65
136	A new analytical system for quantification scratching behaviour in mice. British Journal of Dermatology, 2004, 150, 33-38.	1.4	38
137	Childhood linear IgA disease in association with autoimmune lymphoproliferative syndrome. British Journal of Dermatology, 2004, 150, 578-580.	1.4	20
138	Autoimmune Lymphoproliferative Syndrome with SomaticFasMutations. New England Journal of Medicine, 2004, 351, 1409-1418.	13.9	276
139	SAP mediates specific cytotoxic T-cell functions in X-linked lymphoproliferative disease. Blood, 2004, 103, 3821-3827.	0.6	104
140	Effect of Mycobacterium vaccae on atopic dermatitis in children of different ages. British Journal of Dermatology, 2003, 149, 1029-1034.	1.4	47
141	Association between novel GM-CSF gene polymorphisms and the frequency and severity of atopic dermatitis. Journal of Allergy and Clinical Immunology, 2003, 112, 593-598.	1.5	43
142	End-Organ Dysfunction in Cystic Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2003, 167, 384-389.	2.5	66
143	Autoimmunity in human primary immunodeficiency diseases. Blood, 2002, 99, 2694-2702.	0.6	163
144	Age-Related Prevalence and Antibiotic Resistance of Pathogenic Staphylococci and Streptococci in Children With Infected Atopic Dermatitis at a Single-Specialty Center. Archives of Dermatology, 2002, 138, 939-41.	1.7	52

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145	Hyper IgD syndrome (HIDS) associated with in vitro evidence of defective monocyte TNFRSF1A shedding and partial response to TNF receptor blockade with etanercept. Clinical and Experimental Immunology, 2002, 130, 484-488.	1.1	60
146	Intradermal administration of a killed Mycobacterium vaccae suspension (SRL 172) is associated with improvement in atopic dermatitis in children with moderate-to-severe disease. Journal of Allergy and Clinical Immunology, 2001, 107, 531-534.	1.5	164
147	Atopic dermatitis is associated with a low-producer transforming growth factor \hat{l}^21 cytokine genotype. Journal of Allergy and Clinical Immunology, 2001, 108, 281-284.	1.5	104
148	V(D)J recombination defects in lymphocytes due to RAG mutations: severe immunodeficiency with a spectrum of clinical presentations. Blood, 2001, 97, 81-88.	0.6	324
149	Children with atopic dermatitis who carry toxin-positive Staphylococcus aureus strains have an expansion of blood CD5â ⁻² B lymphocytes without an increase in disease severity. Clinical and Experimental Immunology, 2001, 125, 184-189.	1.1	11
150	Cytokine promoter gene polymorphisms and idiopathic recurrent pregnancy loss. Journal of Reproductive Immunology, 2001, 51, 21-27.	0.8	82
151	Cytomegalovirus infection in infants with autoimmune lymphoproliferative syndrome (ALPS). Clinical and Experimental Immunology, 2000, 121, 353-357.	1.1	12
152	Atopic eczema is associated with delayed maturation of the antibody response to Pneumococcal vaccine. Clinical and Experimental Immunology, 2000, 122, 16-19.	1.1	57
153	Clinical course of patients with major histocompatibility complex class II deficiency. Archives of Disease in Childhood, 2000, 83, 356-359.	1.0	59
154	TGF-beta 1 genotype and accelerated decline in lung function of patients with cystic fibrosis. Thorax, 2000, 55, 459-462.	2.7	192
155	X linked lymphoproliferative disease in a United Kingdom family. Archives of Disease in Childhood, 1998, 79, 52-55.	1.0	14
156	Dietary management of atopic eczema: is this justified?. British Journal of Hospital Medicine, 1998, 59, 690-2.	0.3	0
157	Infantile nephrotic syndrome and atopy. Pediatric Nephrology, 1996, 10, 509-510.	0.9	7
158	Suppression of allogeneic reactivity in vitm by the syncytiotrophoblast membrane glycocalyx of the human term placenta is carbohydrate dependent. Glycobiology, 1994, 4, 39-47.	1.3	22
159	Pre-eclampsia is associated with an increase in trophoblast glycogen content and glycogen synthase activity, similar to that found in hydatidiform moles Journal of Clinical Investigation, 1993, 91, 2744-2753.	3.9	71
160	Glycoprotein glycosylation and the immunosuppressive effects of human pregnancy serum. Journal of Reproductive Immunology, 1992, 21, 97-102.	0.8	8
161	Syncytiotrophoblast membrane protein glycosylation patterns in normal human pregnancy and changes with gestational age and parturition. Placenta, 1991, 12, 637-651.	0.7	18
162	Free and Sulfate-Conjugated Catecholamines during Exercise in Man*. Journal of Clinical Endocrinology and Metabolism, 1984, 58, 415-418.	1.8	32

#	Article	IF	CITATIONS
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