

Yu Lung Lau

List of Publications by Year in descending order

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Version: 2024-02-01

315
papers

16,628
citations

14655

66
h-index

23533

111
g-index

325
all docs

325
docs citations

325
times ranked

22342
citing authors

#	ARTICLE	IF	CITATIONS
1	Vaccine breakthrough hypoxemic COVID-19 pneumonia in patients with auto-Abs neutralizing type I IFNs. <i>Science Immunology</i> , 2023, 8, .	11.9	35
2	Second dose of COVID-19 vaccination in immediate reactions to the first BNT162b2. <i>Pediatric Allergy and Immunology</i> , 2022, 33, e13683.	2.6	8
3	A global effort to dissect the human genetic basis of resistance to SARS-CoV-2 infection. <i>Nature Immunology</i> , 2022, 23, 159-164.	14.5	41
4	Epidemiology of Acute Myocarditis/Pericarditis in Hong Kong Adolescents Following Comirnaty Vaccination. <i>Clinical Infectious Diseases</i> , 2022, 75, 673-681.	5.8	88
5	Human genetic and immunological determinants of critical COVID-19 pneumonia. <i>Nature</i> , 2022, 603, 587-598.	27.8	216
6	Adolescents' attitudes to the COVID-19 vaccination. <i>Vaccine</i> , 2022, 40, 967-969.	3.8	26
7	Exosomes derived from γ -T cells synergize with radiotherapy and preserve antitumor activities against nasopharyngeal carcinoma in immunosuppressive microenvironment. , 2022, 10, e003832.		24
8	Bacillus Calmette-Guérin Scar erythema in a 14-year-old girl post-BNT162b2 vaccination. <i>Pediatrics International</i> , 2022, 64, e15090.	0.5	1
9	Antibody responses to 2 doses of mRNA COVID-19 vaccine in pediatric patients with kidney diseases. <i>Kidney International</i> , 2022, 101, 1069-1072.	5.2	13
10	Myocarditis Following COVID-19 BNT162b2 Vaccination Among Adolescents in Hong Kong. <i>JAMA Pediatrics</i> , 2022, 176, 612.	6.2	46
11	Comprehensive analysis of recessive carrier status using exome and genome sequencing data in 1543 Southern Chinese. <i>Npj Genomic Medicine</i> , 2022, 7, 23.	3.8	6
12	Safety and reactogenicity of a liquid formulation of human rotavirus vaccine (porcine) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 307 Td (circ 2184-2190.	3.8	3
13	Survey data on the attitudes of adolescents in Hong Kong towards the COVID-19 vaccination. <i>Data in Brief</i> , 2022, 42, 108069.	1.0	0
14	Inborn Errors of Immunity in Algerian Children and Adults: A Single-Center Experience Over a Period of 13 Years (2008-2021). <i>Frontiers in Immunology</i> , 2022, 13, 900091.	4.8	4
15	Fatal SARS in X-Linked Lymphoproliferative Disease Type 1: A Case Report. <i>Frontiers in Pediatrics</i> , 2022, 10, 794110.	1.9	3
16	COVID-19 vaccine acceptance and hesitancy among ethnic minorities in Hong Kong. <i>Human Vaccines and Immunotherapeutics</i> , 2022, 18, 1-6.	3.3	3
17	The risk of COVID-19 death is much greater and age dependent with type I IFN autoantibodies. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2022, 119, e2200413119.	7.1	110
18	Impact of a focus education in Zoom on COVID-19 vaccine hesitancy in Hong Kong parents of the preschoolers. <i>Human Vaccines and Immunotherapeutics</i> , 2022, 18, .	3.3	3

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19	Recessive inborn errors of type I IFN immunity in children with COVID-19 pneumonia. <i>Journal of Experimental Medicine</i> , 2022, 219, .	8.5	59
20	Severity of SARS-CoV-2 Omicron BA.2 infection in unvaccinated hospitalized children: comparison to influenza and parainfluenza infections. <i>Emerging Microbes and Infections</i> , 2022, 11, 1742-1750.	6.5	43
21	Immunogenicity and reactogenicity of SARS-CoV-2 vaccines BNT162b2 and CoronaVac in healthy adolescents. <i>Nature Communications</i> , 2022, 13, .	12.8	42
22	Glucose metabolism controls human γ T-cell-mediated tumor immunosurveillance in diabetes. , 2022, 19, 944-956.		8
23	Excessive deubiquitination of NLRP3-R779C variant contributes to very-early-onset inflammatory bowel disease development. <i>Journal of Allergy and Clinical Immunology</i> , 2021, 147, 267-279.	2.9	38
24	Hemophagocytic Lymphohistiocytosis in Children with Chronic Granulomatous Disease—Single-Center Experience from North India. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2021, 9, 771-782.e3.	3.8	7
25	Actionable secondary findings in 1116 Hong Kong Chinese based on exome sequencing data. <i>Journal of Human Genetics</i> , 2021, 66, 637-641.	2.3	3
26	Liver Abscess in Chronic Granulomatous Disease—Two Decades of Experience from a Tertiary Care Centre in North-West India. <i>Journal of Clinical Immunology</i> , 2021, 41, 552-564.	3.8	7
27	Genome-wide association study on Northern Chinese identifies <i>KLF2</i> , <i>DOT1L</i> and <i>STAB2</i> associated with systemic lupus erythematosus. <i>Rheumatology</i> , 2021, 60, 4407-4417.	1.9	16
28	Saliva viral load better correlates with clinical and immunological profiles in children with coronavirus disease 2019. <i>Emerging Microbes and Infections</i> , 2021, 10, 235-241.	6.5	21
29	Clinical, Immunological, and Molecular Profile of Chronic Granulomatous Disease: A Multi-Centric Study of 236 Patients From India. <i>Frontiers in Immunology</i> , 2021, 12, 625320.	4.8	31
30	Identification of 38 novel loci for systemic lupus erythematosus and genetic heterogeneity between ancestral groups. <i>Nature Communications</i> , 2021, 12, 772.	12.8	128
31	Rare versus common diseases: a false dichotomy in precision medicine. <i>Npj Genomic Medicine</i> , 2021, 6, 19.	3.8	14
32	Actionable pharmacogenetic variants in Hong Kong Chinese exome sequencing data and projected prescription impact in the Hong Kong population. <i>PLoS Genetics</i> , 2021, 17, e1009323.	3.5	17
33	A thematic study: impact of COVID-19 pandemic on rare disease organisations and patients across ten jurisdictions in the Asia Pacific region. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 119.	2.7	22
34	Wiskott Aldrich Syndrome: A Multi-Institutional Experience From India. <i>Frontiers in Immunology</i> , 2021, 12, 627651.	4.8	16
35	Clinical Characteristics and Transmission of COVID-19 in Children and Youths During 3 Waves of Outbreaks in Hong Kong. <i>JAMA Network Open</i> , 2021, 4, e218824.	5.9	48
36	Phenotypic and Functional Characteristics of a Novel Influenza Virus Hemagglutinin-Specific Memory NK Cell. <i>Journal of Virology</i> , 2021, 95, .	3.4	8

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37	Invasive cerebral phaeohyphomycosis in a Chinese boy with CARD9 deficiency and showing unique radiological features, managed with surgical excision and antifungal treatment. <i>International Journal of Infectious Diseases</i> , 2021, 107, 59-61.	3.3	7
38	Shared genetic study gives insights into the shared and distinct pathogenic immunity components of IgA nephropathy and SLE. <i>Molecular Genetics and Genomics</i> , 2021, 296, 1017-1026.	2.1	4
39	A Fetus with Congenital Microcephaly, Microphthalmia and Cataract Was Detected with Biallelic Variants in the OCLN Gene: A Case Report. <i>Diagnostics</i> , 2021, 11, 1576.	2.6	0
40	Perception of personalized medicine, pharmacogenomics, and genetic testing among undergraduates in Hong Kong. <i>Human Genomics</i> , 2021, 15, 54.	2.9	9
41	Hospital mortality in patients with rare diseases during pandemics: lessons learnt from the COVID-19 and SARS pandemics. <i>Orphanet Journal of Rare Diseases</i> , 2021, 16, 361.	2.7	10
42	Autoantibodies neutralizing type I IFNs are present in ~4% of uninfected individuals over 70 years old and account for ~20% of COVID-19 deaths. <i>Science Immunology</i> , 2021, 6, .	11.9	357
43	X-linked recessive TLR7 deficiency in ~1% of men under 60 years old with life-threatening COVID-19. <i>Science Immunology</i> , 2021, 6, .	11.9	267
44	Epidemiology and Trends of Infective Meningitis in Neonates and Infants Less than 3 Months Old in Hong Kong. <i>International Journal of Infectious Diseases</i> , 2021, 111, 288-294.	3.3	5
45	A Novel X-Linked Inhibitor of Apoptosis Deficient Variant Showing Attenuated Epstein-Barr Virus Response. <i>Journal of the Pediatric Infectious Diseases Society</i> , 2021, 10, 345-348.	1.3	3
46	Human Complement C4B Allotypes and Deficiencies in Selected Cases With Autoimmune Diseases. <i>Frontiers in Immunology</i> , 2021, 12, 739430.	4.8	11
47	HLA alleles associated with asparaginase hypersensitivity in Chinese children. <i>Journal of Hematology and Oncology</i> , 2021, 14, 182.	17.0	2
48	The estimated age-group specific influenza vaccine coverage rates in Hong Kong and the impact of the school outreach vaccination program. <i>Human Vaccines and Immunotherapeutics</i> , 2021, , 1-5.	3.3	5
49	NLRP3 Inflammasome Contributes to Host Defense Against <i>Talaromyces marneffe</i> Infection. <i>Frontiers in Immunology</i> , 2021, 12, 760095.	4.8	8
50	Phenomic Analysis of Chronic Granulomatous Disease Reveals More Severe Integumentary Infections in X-Linked Compared With Autosomal Recessive Chronic Granulomatous Disease. <i>Frontiers in Immunology</i> , 2021, 12, 803763.	4.8	3
51	Assessment of SARS-CoV-2 Immunity in Convalescent Children and Adolescents. <i>Frontiers in Immunology</i> , 2021, 12, 797919.	4.8	13
52	Accelerated Immunodeficiency-associated Vaccine-derived Poliovirus Serotype 3 Sequence Evolution Rate in an 11-week-old Boy With X-linked Agammaglobulinemia and Perinatal Human Immunodeficiency Virus Exposure. <i>Clinical Infectious Diseases</i> , 2020, 70, 132-135.	5.8	2
53	Evaluating impact of school outreach vaccination programme in Hong Kong influenza season 2018-2019. <i>Human Vaccines and Immunotherapeutics</i> , 2020, 16, 823-826.	3.3	14
54	Genetic Approaches for Definitive Diagnosis of Agammaglobulinemia in Consanguineous Families. <i>Journal of Clinical Immunology</i> , 2020, 40, 96-104.	3.8	3

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55	Identification of Regulatory Modules That Stratify Lupus Disease Mechanism through Integrating Multi-Omics Data. <i>Molecular Therapy - Nucleic Acids</i> , 2020, 19, 318-329.	5.1	10
56	Exosomes derived from VÎ2-T cells control Epstein-Barr virus-associated tumors and induce T cell antitumor immunity. <i>Science Translational Medicine</i> , 2020, 12, .	12.4	48
57	Functional analysis and evaluation of respiratory cilia in healthy Chinese children. <i>Respiratory Research</i> , 2020, 21, 259.	3.6	10
58	Haematological and immunological data of Chinese children infected with coronavirus disease 2019. <i>Data in Brief</i> , 2020, 31, 105953.	1.0	15
59	Monoallelic Mutations in <i>CC2D1A</i> Suggest a Novel Role in Human Heterotaxy and Ciliary Dysfunction. <i>Circulation Genomic and Precision Medicine</i> , 2020, 13, e003000.	3.6	4
60	CHARGE syndrome patient with novel CHD7 mutation presenting with severe laryngomalacia and feeding difficulty. <i>BMJ Case Reports</i> , 2020, 13, e233037.	0.5	0
61	Coexistence of paternally-inherited ABCC8 mutation and mosaic paternal uniparental disomy 11p hyperinsulinism. <i>International Journal of Pediatric Endocrinology (Springer)</i> , 2020, 2020, 13.	1.6	4
62	Current Perspectives and Unmet Needs of Primary Immunodeficiency Care in Asia Pacific. <i>Frontiers in Immunology</i> , 2020, 11, 1605.	4.8	13
63	Hyper IgE Syndrome Associated With Warts: A First Case of Dedicator of Cytokinesis 8 Deficiency in the Philippines. <i>Frontiers in Pediatrics</i> , 2020, 8, 604725.	1.9	2
64	A case report of complement C4B deficiency in a patient with steroid and IVIG-refractory anti-NMDA receptor encephalitis. <i>BMC Neurology</i> , 2020, 20, 339.	1.8	5
65	Rapid whole-exome sequencing facilitates precision medicine in paediatric rare disease patients and reduces healthcare costs. <i>The Lancet Regional Health - Western Pacific</i> , 2020, 1, 100001.	2.9	40
66	COVID-19 in children across three Asian cosmopolitan regions. <i>Emerging Microbes and Infections</i> , 2020, 9, 2588-2596.	6.5	21
67	A Comparison Between Chinese Children Infected with Coronavirus Disease-2019 and with Severe Acute Respiratory Syndrome 2003. <i>Journal of Pediatrics</i> , 2020, 224, 30-36.	1.8	25
68	Independent Replication on Genome-Wide Association Study Signals Identifies IRF3 as a Novel Locus for Systemic Lupus Erythematosus. <i>Frontiers in Genetics</i> , 2020, 11, 600.	2.3	9
69	Host DNA released by NETosis in neutrophils exposed to seasonal H1N1 and highly pathogenic H5N1 influenza viruses. <i>Respiratory Research</i> , 2020, 21, 160.	3.6	14
70	Cost-effectiveness analysis of chromosomal microarray as a primary test for prenatal diagnosis in Hong Kong. <i>BMC Pregnancy and Childbirth</i> , 2020, 20, 109.	2.4	7
71	NFÎ2 mutation as a novel cause for inherited thrombocytopenia. <i>British Journal of Haematology</i> , 2020, 189, e41-e44.	2.5	3
72	Evaluating the Clinical Utility of Genome Sequencing for Cytogenetically Balanced Chromosomal Abnormalities in Prenatal Diagnosis. <i>Frontiers in Genetics</i> , 2020, 11, 620162.	2.3	4

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73	Clinical, Immunological, and Molecular Features of Severe Combined Immune Deficiency: A Multi-Institutional Experience From India. <i>Frontiers in Immunology</i> , 2020, 11, 619146.	4.8	31
74	Clinical and Genetic Profile of X-Linked Agammaglobulinemia: A Multicenter Experience From India. <i>Frontiers in Immunology</i> , 2020, 11, 612323.	4.8	16
75	Risk factors for drug allergies in Chinese children. <i>Asian Pacific Journal of Allergy and Immunology</i> , 2020, 38, 271-278.	0.4	0
76	Application of Flow Cytometry in the Diagnostics Pipeline of Primary Immunodeficiencies Underlying Disseminated <i>Talaromyces marneffei</i> Infection in HIV-Negative Children. <i>Frontiers in Immunology</i> , 2019, 10, 2189.	4.8	30
77	Genetic studies on systemic lupus erythematosus in East Asia point to population differences in disease susceptibility. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2019, 181, 262-268.	1.6	8
78	Clinical and genetic characteristics of Chinese pediatric patients with chronic granulomatous disease. <i>Pediatric Allergy and Immunology</i> , 2019, 30, 378-386.	2.6	27
79	Cell lineage-specific genome-wide DNA methylation analysis of patients with paediatric-onset systemic lupus erythematosus. <i>Epigenetics</i> , 2019, 14, 341-351.	2.7	28
80	HLA-IMPUTER: an easy to use web application for HLA imputation and association analysis using population-specific reference panels. <i>Bioinformatics</i> , 2019, 35, 1244-1246.	4.1	4
81	Health professionals'™ involvement and information provision in genetic counseling following prenatal diagnosis of sex chromosome aneuploidy in Hong Kong. <i>International Journal of Gynecology and Obstetrics</i> , 2019, 144, 314-316.	2.3	0
82	Cross-reactivity pattern of a rare presentation of generalized delayed-type hypersensitivity to local anaesthetics. <i>Asian Pacific Journal of Allergy and Immunology</i> , 2019, 37, 179-182.	0.4	3
83	A case of prenatal isolated talipes and 22q11.2 deletion syndrome"an important chromosomal disorder missed by noninvasive prenatal screening. <i>Prenatal Diagnosis</i> , 2018, 38, 376-378.	2.3	6
84	Identification of <i>ST3AGL4</i> , <i>MFHAS1</i> , <i>CSNK2A2</i> and <i>CD226</i> as loci associated with systemic lupus erythematosus (SLE) and evaluation of SLE genetics in drug repositioning. <i>Annals of the Rheumatic Diseases</i> , 2018, 77, 1078-1084.	0.9	34
85	RASGRP1 mutation in autoimmune lymphoproliferative syndrome-like disease. <i>Journal of Allergy and Clinical Immunology</i> , 2018, 142, 595-604.e16.	2.9	44
86	Uncompromised NK cell activation is essential for virus-specific CTL activity during acute influenza virus infection. <i>Cellular and Molecular Immunology</i> , 2018, 15, 827-837.	10.5	29
87	Infectious and non-infectious complications in primary immunodeficiency disorders: an autopsy study from North India. <i>Journal of Clinical Pathology</i> , 2018, 71, 425-435.	2.0	8
88	PD-1/PD-L1 Pathway Mediates the Alleviation of Pulmonary Fibrosis by Human Mesenchymal Stem Cells in Humanized Mice. <i>American Journal of Respiratory Cell and Molecular Biology</i> , 2018, 58, 684-695.	2.9	73
89	Healthcare burden of rare diseases in Hong Kong " adopting ORPHAcodes in ICD-10 based healthcare administrative datasets. <i>Orphanet Journal of Rare Diseases</i> , 2018, 13, 147.	2.7	39
90	Type I and III Interferon Productions Are Impaired in X-Linked Agammaglobulinemia Patients Toward Poliovirus but Not Influenza Virus. <i>Frontiers in Immunology</i> , 2018, 9, 1826.	4.8	9

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91	Clinical and molecular features of X-linked hyper IgM syndrome – An experience from North India. <i>Clinical Immunology</i> , 2018, 195, 59-66.	3.2	16
92	Detecting Small Inversions Using SRinversion. <i>Methods in Molecular Biology</i> , 2018, 1833, 107-114.	0.9	0
93	Meta-analysis of GWAS – Both Chinese and European populations identifies GPR173 as a novel X chromosome susceptibility gene for SLE. <i>Arthritis Research and Therapy</i> , 2018, 20, 92.	3.5	19
94	Exome sequencing for paediatric-onset diseases: impact of the extensive involvement of medical geneticists in the diagnostic odyssey. <i>Npj Genomic Medicine</i> , 2018, 3, 19.	3.8	11
95	Spinal Muscular Atrophy With Respiratory Distress Type 1 – A Child With Atypical Presentation. <i>Child Neurology Open</i> , 2018, 5, 2329048X1876981.	1.1	2
96	The unmet provision of allergy services in Hong Kong impairs capability for allergy prevention – implications for the Asia Pacific region. <i>Asian Pacific Journal of Allergy and Immunology</i> , 2018, 37, 1-8.	0.4	17
97	Prenatal Tobacco Exposure Shortens Telomere Length in Children. <i>Nicotine and Tobacco Research</i> , 2017, 19, 111-118.	2.6	32
98	Chronic Mucocutaneous Candidiasis. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2017, 5, 1119-1121.	3.8	6
99	Confirmation of five novel susceptibility loci for Systemic Lupus Erythematosus (SLE) and integrated network analysis of 82 SLE susceptibility loci. <i>Human Molecular Genetics</i> , 2017, 26, ddx026.	2.9	47
100	Homozygous transcription factor 3 gene (TCF3) mutation is associated with severe hypogammaglobulinemia and B-cell acute lymphoblastic leukemia. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 1191-1194.e4.	2.9	38
101	Emerging Infections and Pertinent Infections Related to Travel for Patients with Primary Immunodeficiencies. <i>Journal of Clinical Immunology</i> , 2017, 37, 650-692.	3.8	6
102	Wiskott-Aldrich syndrome protein regulates autophagy and inflammasome activity in innate immune cells. <i>Nature Communications</i> , 2017, 8, 1576.	12.8	50
103	Use of clinical chromosomal microarray in Chinese patients with autism spectrum disorder – implications of a copy number variation involving DPP10. <i>Molecular Autism</i> , 2017, 8, 31.	4.9	16
104	Sclerosing cholangitis and intracranial lymphoma in a child with classical Wiskott – Aldrich syndrome. <i>Pediatric Blood and Cancer</i> , 2017, 64, 106-109.	1.5	12
105	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
106	Cellular and Molecular Defects Underlying Invasive Fungal Infections – Revelations from Endemic Mycoses. <i>Frontiers in Immunology</i> , 2017, 8, 735.	4.8	57
107	Family History of Early Infant Death Correlates with Earlier Age at Diagnosis But Not Shorter Time to Diagnosis for Severe Combined Immunodeficiency. <i>Frontiers in Immunology</i> , 2017, 8, 808.	4.8	34
108	Genome-Wide DNA Methylation Analysis of Chinese Patients with Systemic Lupus Erythematosus Identified Hypomethylation in Genes Related to the Type I Interferon Pathway. <i>PLoS ONE</i> , 2017, 12, e0169553.	2.5	40

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109	Prevalence of and Risk Factors for Childhood Asthma, Rhinitis, and Eczema in Hong Kong: Proposal for a Cross-Sectional Survey. <i>JMIR Research Protocols</i> , 2017, 6, e106.	1.0	5
110	Genome-wide association meta-analysis in Chinese and European individuals identifies ten new loci associated with systemic lupus erythematosus. <i>Nature Genetics</i> , 2016, 48, 940-946.	21.4	283
111	Effects of Complement <i>C4</i> Gene Copy Number Variations, Size Dichotomy, and <i>C4A</i> Deficiency on Genetic Risk and Clinical Presentation of Systemic Lupus Erythematosus in East Asian Populations. <i>Arthritis and Rheumatology</i> , 2016, 68, 1442-1453.	5.6	58
112	Smoke-free legislation reduces hospital admissions for childhood lower respiratory tract infection. <i>Tobacco Control</i> , 2016, 25, e90-e94.	3.2	16
113	X-Linked Agammaglobulinemia in a Large Series of North African Patients: Frequency, Clinical Features and Novel BTK Mutations. <i>Journal of Clinical Immunology</i> , 2016, 36, 187-194.	3.8	28
114	Association of <i>MTOR</i> Mutations With Developmental Brain Disorders, Including Megalencephaly, Focal Cortical Dysplasia, and Pigmentary Mosaicism. <i>JAMA Neurology</i> , 2016, 73, 836.	9.0	234
115	Rotavirus vaccine effectiveness in Hong Kong children. <i>Vaccine</i> , 2016, 34, 4935-4942.	3.8	18
116	X-linked agammaglobulinemia. <i>Annals of Allergy, Asthma and Immunology</i> , 2016, 117, 405-411.	1.0	22
117	SRinversion: a tool for detecting short inversions by splitting and re-aligning poorly mapped and unmapped sequencing reads. <i>Bioinformatics</i> , 2016, 32, 3559-3565.	4.1	7
118	Human oropharynx as natural reservoir of <i>Streptobacillus hongkongensis</i> . <i>Scientific Reports</i> , 2016, 6, 24419.	3.3	11
119	Incidence of rotavirus gastroenteritis by age in African, Asian and European children: Relevance for timing of rotavirus vaccination. <i>Human Vaccines and Immunotherapeutics</i> , 2016, 12, 2406-2412.	3.3	36
120	Genome-wide search followed by replication reveals genetic interaction of <i>CD80</i> and <i>ALOX5AP</i> associated with systemic lupus erythematosus in Asian populations. <i>Annals of the Rheumatic Diseases</i> , 2016, 75, 891-898.	0.9	28
121	Gene-Based Meta-Analysis of Genome-Wide Association Study Data Identifies Independent Single Nucleotide Polymorphisms in <i>ANXA6</i> as Being Associated With Systemic Lupus Erythematosus in Asian Populations. <i>Arthritis and Rheumatology</i> , 2015, 67, 2966-2977.	5.6	14
122	Compound heterozygous mutations in <i>TTC7A</i> cause familial multiple intestinal atresias and severe combined immunodeficiency. <i>Clinical Genetics</i> , 2015, 88, 542-549.	2.0	27
123	The Therapeutic Effect of Pamidronate on Lethal Avian Influenza A H7N9 Virus Infected Humanized Mice. <i>PLoS ONE</i> , 2015, 10, e0135999.	2.5	12
124	Tricho-hepato-enteric syndrome (THE-S): two cases and review of the literature. <i>European Journal of Pediatrics</i> , 2015, 174, 1405-1411.	2.7	15
125	Meta-analysis of two Chinese populations identifies an autoimmune disease risk allele in 22q11.21 as associated with systemic lupus erythematosus. <i>Arthritis Research and Therapy</i> , 2015, 17, 67.	3.5	6
126	Lethal Coinfection of Influenza Virus and <i>Streptococcus pneumoniae</i> Lowers Antibody Response to Influenza Virus in Lung and Reduces Numbers of Germinal Center B Cells, T Follicular Helper Cells, and Plasma Cells in Mediastinal Lymph Node. <i>Journal of Virology</i> , 2015, 89, 2013-2023.	3.4	23

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127	Functional polymorphisms of the CCL2 and MBL genes cumulatively increase susceptibility to severe acute respiratory syndrome coronavirus infection. <i>Journal of Infection</i> , 2015, 71, 101-109.	3.3	78
128	HaploShare: identification of extended haplotypes shared by cases and evaluation against controls. <i>Genome Biology</i> , 2015, 16, 92.	8.8	7
129	HLAreporter: a tool for HLA typing from next generation sequencing data. <i>Genome Medicine</i> , 2015, 7, 25.	8.2	62
130	Prevalence of BTK mutations in male Algerian patients with agammaglobulinemia and severe B cell lymphopenia. <i>Clinical Immunology</i> , 2015, 161, 286-290.	3.2	5
131	X-linked hyper-IgM syndrome with CD40LG mutation: Two case reports and literature review in Taiwanese patients. <i>Journal of Microbiology, Immunology and Infection</i> , 2015, 48, 113-118.	3.1	25
132	Meta-analysis of GWAS on two Chinese populations followed by replication identifies novel genetic variants on the X chromosome associated with systemic lupus erythematosus. <i>Human Molecular Genetics</i> , 2015, 24, 274-284.	2.9	35
133	Solving the genetic puzzle of systemic lupus erythematosus. <i>Pediatric Nephrology</i> , 2015, 30, 1735-1748.	1.7	9
134	A patient with mosaic neurofibromatosis type 2 presenting with early onset meningioma. <i>BMJ Case Reports</i> , 2014, 2014, bcr2014203919-bcr2014203919.	0.5	3
135	Inferring Influenza Infection Attack Rate from Seroprevalence Data. <i>PLoS Pathogens</i> , 2014, 10, e1004054.	4.7	46
136	Three SNPs in chromosome 11q23.3 are independently associated with systemic lupus erythematosus in Asians. <i>Human Molecular Genetics</i> , 2014, 23, 524-533.	2.9	29
137	Targeted Activation of Human V α 39V β 2-T Cells Controls Epstein-Barr Virus-Induced B Cell Lymphoproliferative Disease. <i>Cancer Cell</i> , 2014, 26, 565-576.	16.8	115
138	Under-recognition of 22q11.2 deletion in adult Chinese patients with conotruncal anomalies: Implications in transitional care. <i>European Journal of Medical Genetics</i> , 2014, 57, 306-311.	1.3	29
139	BCG vaccination in patients with severe combined immunodeficiency: Complications, risks, and vaccination policies. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 1134-1141.	2.9	212
140	IL-10-producing regulatory B cells induced by IL-33 (BregIL-33) effectively attenuate mucosal inflammatory responses in the gut. <i>Journal of Autoimmunity</i> , 2014, 50, 107-122.	6.5	158
141	<i>Penicillium marneffei</i> infection and impaired IFN- γ immunity in humans with autosomal-dominant gain-of-phosphorylation STAT1 mutations. <i>Journal of Allergy and Clinical Immunology</i> , 2014, 133, 894-896.e5.	2.9	69
142	Recurrent abdominal pain as the presentation of tumor necrosis factor receptor-associated periodic syndrome (TRAPS) in an Asian girl: A case report and review of the literature. <i>Journal of Microbiology, Immunology and Infection</i> , 2014, 47, 550-554.	3.1	7
143	Influenza Virus-Induced Lung Inflammation Was Modulated by Cigarette Smoke Exposure in Mice. <i>PLoS ONE</i> , 2014, 9, e86166.	2.5	24
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