

Wen-I Lee

List of Publications by Year in descending order

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Version: 2024-02-01

33
papers

764
citations

687363

13
h-index

526287

27
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36
all docs

36
docs citations

36
times ranked

1172
citing authors

#	ARTICLE	IF	CITATIONS
1	Molecular analysis of a large cohort of patients with the hyper immunoglobulin M (IgM) syndrome. <i>Blood</i> , 2005, 105, 1881-1890.	1.4	193
2	Incidence of severe combined immunodeficiency through newborn screening in a Chinese population. <i>Journal of the Formosan Medical Association</i> , 2015, 114, 12-16.	1.7	68
3	Anti-IFN- γ autoantibodies are strongly associated with HLA-DR*15:02/16:02 and HLA-DQ*05:01/05:02 across Southeast Asia. <i>Journal of Allergy and Clinical Immunology</i> , 2016, 137, 945-948.e8.	2.9	63
4	Akhil Kakroo. <i>Journal of Clinical Immunology</i> , 2005, 25, 162-173.	3.8	61
5	Novel Mutations of the Tetratricopeptide Repeat Domain 7A Gene and Phenotype/Genotype Comparison. <i>Frontiers in Immunology</i> , 2017, 8, 1066.	4.8	39
6	Patients with inhibitory and neutralizing auto-antibodies to interferon- β resemble the sporadic adult-onset phenotype of Mendelian Susceptibility to Mycobacterial Disease (MSMD) lacking Bacille Calmette-Guérin (BCG)-induced diseases. <i>Immunobiology</i> , 2013, 218, 762-771.	1.9	35
7	Chinese Patients with Defective IL-12/23-Interferon- β Circuit in Taiwan: Partial Dominant Interferon- β Receptor 1 Mutation Presenting as Cutaneous Granuloma and IL-12 Receptor β 1 Mutation as Pneumatocele. <i>Journal of Clinical Immunology</i> , 2009, 29, 238-245.	3.8	31
8	Distribution, Infections, Treatments and Molecular Analysis in a Large Cohort of Patients with Primary Immunodeficiency Diseases (PIDs) in Taiwan. <i>Journal of Clinical Immunology</i> , 2006, 26, 274-283.	3.8	30
9	Clinical Features and Genetic Analysis of Taiwanese Patients With the Hyper IgM Syndrome Phenotype. <i>Pediatric Infectious Disease Journal</i> , 2013, 32, 1010-1016.	2.0	24
10	Inducible CO-stimulator molecule, a candidate gene for defective isotype switching, is normal in patients with hyper-IgM syndrome of unknown molecular diagnosis. <i>Journal of Allergy and Clinical Immunology</i> , 2003, 112, 958-964.	2.9	21
11	Identifying Mutations of the Tetratricopeptide Repeat Domain 37 (TTC37) Gene in Infants With Intractable Diarrhea and a Comparison of Asian and Non-Asian Phenotype and Genotype. <i>Medicine (United States)</i> , 2016, 95, e2918.	1.0	15
12	Biomarkers associating endothelial dysregulation in pediatric-onset systemic lupus erythematosus. <i>Pediatric Rheumatology</i> , 2019, 17, 69.	2.1	15
13	Cytotoxic Function and Cytokine Production of Natural Killer Cells and Natural Killer T-Like Cells in Systemic Lupus Erythematosus Regulation with Interleukin-15. <i>Mediators of Inflammation</i> , 2019, 2019, 1-12.	3.0	15
14	Diminished toll-like receptor response in febrile infection-related epilepsy syndrome (FIRES). <i>Biomedical Journal</i> , 2020, 43, 293-304.	3.1	15
15	Analysis of genetic defects in patients with the common variable immunodeficiency phenotype in a single Taiwanese tertiary care hospital. <i>Annals of Allergy, Asthma and Immunology</i> , 2007, 99, 433-442.	1.0	14
16	Persistent improper upregulation of Th17 and T Reg cells in patients with juvenile idiopathic arthritis. <i>Journal of Microbiology, Immunology and Infection</i> , 2016, 49, 402-408.	3.1	14
17	Current Perspectives and Unmet Needs of Primary Immunodeficiency Care in Asia Pacific. <i>Frontiers in Immunology</i> , 2020, 11, 1605.	4.8	13
18	Serial cytokine expressions in infants with incontinentia pigmenti. <i>Immunobiology</i> , 2013, 218, 772-779.	1.9	11

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19	A Nationwide Study of Severe and Protracted Diarrhoea in Patients with Primary Immunodeficiency Diseases. <i>Scientific Reports</i> , 2017, 7, 3669.	3.3	11
20	A Novel CD3G Mutation in a Taiwanese Patient With Normal T Regulatory Function Presenting With the CVID Phenotype Free of Autoimmunity—Analysis of all Genotypes and Phenotypes. <i>Frontiers in Immunology</i> , 2019, 10, 2833.	4.8	11
21	Immunologic Difference between Hypersensitivity to Mosquito Bite and Hemophagocytic Lymphohistiocytosis Associated with Epstein-Barr Virus Infection. <i>PLoS ONE</i> , 2013, 8, e76711.	2.5	10
22	Immunologic Analysis of HIV-Uninfected Taiwanese Children with BCG-Induced Disease. <i>Journal of Clinical Immunology</i> , 2009, 29, 319-329.	3.8	9
23	Applying T-cell receptor excision circles and immunoglobulin δ -deleting recombination excision circles to patients with primary immunodeficiency diseases. <i>Annals of Medicine</i> , 2014, 46, 555-565.	3.8	8
24	Dupilumab for STAT3-Hyper-IgE Syndrome With Refractory Intestinal Complication. <i>Pediatrics</i> , 2021, 148, .	2.1	8
25	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
26	Clinical Features of Female Taiwanese Carriers with X-linked Chronic Granulomatous Disease from 2004 to 2019. <i>Journal of Clinical Immunology</i> , 2021, 41, 1303-1314.	3.8	6
27	Comparison of [3H]-Thymidine, Carboxyfluorescein Diacetate Succinimidyl Ester and Ki-67 in Lymphocyte Proliferation. <i>Frontiers in Pediatrics</i> , 2022, 10, 638549.	1.9	6
28	Identifying patients with neutrophil elastase (ELANE) mutations from patients with a presumptive diagnosis of autoimmune neutropenia. <i>Immunobiology</i> , 2013, 218, 828-833.	1.9	5
29	Pulmonary function evaluation in pediatric patients with primary immunodeficiency complicated by bronchiectasis. <i>Journal of Microbiology, Immunology and Infection</i> , 2020, 53, 1014-1020.	3.1	2
30	Laser therapy for twin-twin transfusion syndrome in a dizygotic monochorionic twin pregnancy: A case report. <i>Taiwanese Journal of Obstetrics and Gynecology</i> , 2021, 60, 791-793.	1.3	2
31	The influence of clinical features mimicking primary immunodeficiency diseases (mPID) on children with Langerhans cell histiocytosis (LCH) — Four with mPID among 39 LCH children from one referral center during 18-year period. <i>Immunobiology</i> , 2020, 225, 151877.	1.9	1
32	Lipopolysaccharide stimulation test on cultured PBMCs assists the discrimination of cryopyrin-associated periodic syndrome from systemic juvenile idiopathic arthritis. <i>Scientific Reports</i> , 2021, 11, 11903.	3.3	1
33	Pneumonia and empyema caused by <i>Methylobacterium</i> in a patient with X-linked chronic granulomatous disease. <i>Pediatrics and Neonatology</i> , 2019, 60, 584-586.	0.9	0