Tiphanie P Vogel

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2805232/publications.pdf

Version: 2024-02-01

38 papers

2,114 citations

567281 15 h-index 330143 37 g-index

38 all docs 38 docs citations

38 times ranked

3824 citing authors

#	Article	IF	Citations
1	Early-onset lymphoproliferation and autoimmunity caused by germline STAT3 gain-of-function mutations. Blood, 2015, 125, 591-599.	1.4	436
2	Activation-Specific Metabolic Requirements for NK Cell IFN- \hat{l}^3 Production. Journal of Immunology, 2015, 194, 1954-1962.	0.8	227
3	Jakinibs for the treatment of immune dysregulation in patients with gain-of-function signal transducer and activator of transcription 1 (STAT1) or STAT3 mutations. Journal of Allergy and Clinical Immunology, 2018, 142, 1665-1669.	2.9	196
4	Multisystem inflammatory syndrome in children and adults (MIS-C/A): Case definition & amp; guidelines for data collection, analysis, and presentation of immunization safety data. Vaccine, 2021, 39, 3037-3049.	3.8	175
5	Clinical Aspects of STAT3 Gain-of-Function Germline Mutations: A Systematic Review. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 1958-1969.e9.	3.8	144
6	Genetic and mechanistic diversity in pediatric hemophagocytic lymphohistiocytosis. Blood, 2018, 132, 89-100.	1.4	139
7	The Ying and Yang of STAT3 in Human Disease. Journal of Clinical Immunology, 2015, 35, 615-623.	3.8	130
8	Heterozygous Truncating Variants in POMP Escape Nonsense-Mediated Decay and Cause a Unique Immune Dysregulatory Syndrome. American Journal of Human Genetics, 2018, 102, 1126-1142.	6.2	128
9	Somatic mutations and clonal hematopoiesis in congenital neutropenia. Blood, 2018, 131, 408-416.	1.4	91
10	Loss- or Gain-of-Function Mutations in ACOX1 Cause Axonal Loss via Different Mechanisms. Neuron, 2020, 106, 589-606.e6.	8.1	71
11	Inflammatory syndromes associated with SARS-CoV-2 infection: dysregulation of the immune response across the age spectrum. Journal of Clinical Investigation, 2020, 130, 6194-6197.	8.2	71
12	Dominant-negative mutations in human <i>IL6ST</i> underlie hyper-lgE syndrome. Journal of Experimental Medicine, 2020, 217, .	8.5	64
13	Inflammatory caspase regulation: maintaining balance between inflammation and cell death in health and disease. FEBS Journal, 2019, 286, 2628-2644.	4.7	49
14	Up, Down, and All Around: Diagnosis and Treatment of Novel STAT3 Variant. Frontiers in Pediatrics, 2017, 5, 49.	1.9	25
15	Successful Treatment of Interstitial Lung Disease in STAT3 Gain-of-Function Using JAK Inhibitors. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 893-897.	5.6	25
16	Successful treatment of chronic atypical neutrophilic dermatosis with lipodystrophy and elevated temperature (CANDLE) syndrome with tofacitinib. Pediatric Dermatology, 2021, 38, 528-529.	0.9	17
17	STAT3 gain of function: a new aetiology of severe rheumatic disease. Rheumatology, 2019, 58, 365-367.	1.9	12
18	Multisystem inflammatory syndrome in children associated with SARS-CoV-2 in a solid organ transplant recipient. American Journal of Transplantation, 2021, 21, 2596-2599.	4.7	12

#	Article	IF	CITATIONS
19	Immune Dysregulation Mimicking Systemic Lupus Erythematosus in a Patient With Lysinuric Protein Intolerance: Case Report and Review of the Literature. Frontiers in Pediatrics, 2021, 9, 673957.	1.9	12
20	STAT3 Gain-of-Function Mutations Underlie Deficiency in Human Nonclassical CD16+ Monocytes and CD141+ Dendritic Cells. Journal of Immunology, 2021, 207, 2423-2432.	0.8	11
21	A Remarkable Response of Granulomatous Hypophysitis to Infliximab in a Patient With a Background of Crohn's Diseaseâ€"A Case Report. Frontiers in Endocrinology, 2020, 11, 350.	3.5	8
22	Azathioprine-Associated Complete NK Cell Deficiency. Journal of Clinical Immunology, 2017, 37, 514-516.	3.8	7
23	A Novel STAT3 Mutation in a Qatari Patient With Hyper-IgE Syndrome. Frontiers in Pediatrics, 2019, 7, 130.	1.9	7
24	Vasoplegic Shock Represents a Dominant Hemodynamic Profile of Multisystem Inflammatory Syndrome Following COVID-19 in Children and Adolescents. Pediatric Critical Care Medicine, 2022, 23, e295-e299.	0.5	7
25	Outcomes After SARS-CoV-2 Vaccination Among Children With a History of Multisystem Inflammatory Syndrome. JAMA Network Open, 2022, 5, e224750.	5.9	7
26	Complicated Diagnosis and Treatment of HA20 due to Contiguous Gene Deletions involving 6q23.3. Journal of Clinical Immunology, 2021, 41, 1420-1423.	3.8	6
27	Genetic errors of immunity distinguish pediatric nonmalignant lymphoproliferative disorders. Journal of Allergy and Clinical Immunology, 2022, 149, 758-766.	2.9	6
28	Immune thrombocytopenia following multisystem inflammatory syndrome in children (MIS-C) – a case series. Pediatric Hematology and Oncology, 2021, 38, 1-8.	0.8	5
29	Comparison of Laboratory and Hemodynamic Time Series Data Across Original, Alpha, and Delta Variants in Patients With Multisystem Inflammatory Syndrome in Children. Pediatric Critical Care Medicine, 2022, 23, e372-e381.	0.5	5
30	The Heart and Pediatric Rheumatology. Rheumatic Disease Clinics of North America, 2014, 40, 61-85.	1.9	4
31	A Zebra at the Rodeo: Dyspnea, Hematuria, and a Family History of Arthritis. Arthritis Care and Research, 2022, 74, 165-170.	3.4	4
32	Pulmonary manifestations and outcomes in paediatric ANCA–associated vasculitis: a single-centre experience. Rheumatology, 2020, 60, 3199-3208.	1.9	3
33	Lung biopsy in the diagnosis of pediatric ANCAâ€associated vasculitis. Pediatric Pulmonology, 2021, 56, 145-152.	2.0	3
34	Pulmonary Histopathology Findings in Patients With STAT3 Gain of Function Syndrome. Pediatric and Developmental Pathology, 2021, 24, 227-234.	1.0	3
35	Case Report: Hyper IgE, but Not the Usual Suspects–Kimura Disease in an Adolescent Female. Frontiers in Pediatrics, 2021, 9, 674317.	1.9	2
36	Genetic Disorders Should Be Considered Prior to Diagnosing Interstitial Pneumonia With Autoimmune Features: Comment on the Review by Wilfong et al. Arthritis and Rheumatology, 2019, 71, 2132-2133.	5.6	1

#	Article	IF	CITATIONS
37	Extracorporeal Membrane Oxygenation Support for Antineutrophil Cytoplasmic Antibody-associated Vasculitides. ASAIO Journal, 2021, Publish Ahead of Print, .	1.6	1
38	Genomic Characterization of a Pediatric Cohort with Non-Malignant Lymphoproliferative Disorders. Blood, 2019, 134, 83-83.	1.4	0