

THURSDAY POSTER PRESENTATIONS

Poster Number	Poster Title	Poster Presenter	Organization
1	Clinical Roadmap for Implementing Results from Electronic Health Records Queries	Aaron T. Chin, MD	UCLA: Division of Allergy, Immunology, and Rheumatology
2	Comprehensive Analysis Reveals a Neuroimmunological Association Induced by Oncogenic Hepatitis Viruses	Adriel L. Nobile	Sao Paulo University - USP
3	Aggressive Immunosuppressive Therapy Is Beneficial Early on in Reducing Autoreactive T Cell Proliferation in Omenn Syndrome/Severe Combined Immunodeficiency	Afia B. Adu-Gyamfi, MD	Mayo Clinic, Pediatrics and Adolescent Medicine
4	Clingen Framework for PIK3CD Variant Classification: Use of Adapted ACMG/AMP Guidelines	Alejandro U. Nieto-Patlán, PhD	Baylor College of Medicine
5	RECQL4 Mutation Is Associated with a Hematopoietic-cell-intrinsic Severe T Cell Deficiency and Is Amenable to Treatment with Unconditioned, Unmanipulated Hematopoietic Stem Cell Transplant	Alexandra E. Grier, MD PhD	Children's Hospital of Pennsylvania
6	Identification of TNFA-related Biomarkers in Patients and Carriers with Adenosine Deaminase 2 Deficiency (DADA2)	Alison L. Brittain, DO, PhD	Nationwide Children's Hospital
7	Oral Pathology in STAT3DN Hyper IgE Syndrome	Amanda K. Urban, CRNP	Clinical Research Directorate, Frederick National Laboratory for Cancer Research
8	A Quantitative Approach for Identifying Health Disparities Driving Diagnostic and Treatment Delays in VEO-IBD	Amanda Salih, MD, MPH	Baylor College of Medicine/Texas Children's Hospital
9	Novel CARD14 Variant: A Case of Autoinflammatory Disorder in a 5-year-old with Therapeutic Insights	Amer Khojah, MD	Umm Al-Qura University
10	Enteropathy in Patients with Common Variable Immunodeficiency: A Dutch Cohort Study	Amir Abdelmoumen, MD	University Medical Center Utrecht, Center of Translational Immunology, Utrecht, The Netherlands
11	Novel Pathogenic MAGT1 Variant Identified in an Adult Patient with XMEN Disease	Anna Lang, MD, PhD	University of Washington
12	Integrative Characterization of the Neuroimmunological Interactions of Major Depressive Disorder and Its Intersection with COVID-19	Anny S. Adri	Universidade de São Paulo
13	A Case of Eosinophilic Fasciitis Successfully Managed with Benralizumab in a Patient with Hypogammaglobulinemia and B-cell Aplasia	Arabelle Abellard, MD	Rush University Medical Center

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14	Good Syndrome: A Case of Bowel Perforation, Chronic Diarrhea and Clostridium Difficile Colitis Leading to Detection of a Thymoma	Arabelle Abellard, MD	Rush University Medical Center
15	Assessing MEFV Variant Pathogenicity by ASC-Specks Flow Detection	Blandine A. Monjarret, MSc	Centre de Recherche Azrieli du CHU Sainte-Justine
16	Expanded phenotypic presentations in CADINS disease associated with novel CARD11 dominant interfering variants that impact NF-kB and AP-1 signaling	Bradly M. Bauman, PhD	Department of Pharmacology & Molecular Therapeutics, Uniformed Services University of the Health Sciences
17	DHR-based Flow Cytometry Beyond CGD	Chaitanya Maddukuri, DO	Cleveland Clinic
18	Novel SAMD9 Variant Leading to MIRAGE Syndrome Treated with Subcutaneous Immunoglobulin: A Case Report	Christopher T. Peek, MD, PhD	Baylor College of Medicine
19	Patient with Adenosine Deaminase Severe Combined Immunodeficiency (ADA-SCID) and Glutaric Aciduria Type 1 (GA1) Successfully Treated with Delayed Hematopoietic Stem Cell Transplant (HSCT)	David M. Roth, MD, MSPH	UPMC Children's Hospital of Pediatrics
20	CMV-specific Adaptive Immune Response in U.S. Mennonite Patients with Hypomorphic RAG1 or RAG2 Mutations Presenting with Clinical Variability	David Potts, MSc	Department of Pediatrics, Division of Allergy & Immunology, University of South Florida
21	Diagnostic Approach in a Young Adult with RAG1 Variants Initially Diagnosed with Specific Antibody Deficiency and Bronchiectasis	David Potts, MSc	Department of Pediatrics, Division of Allergy & Immunology, University of South Florida
22	Diagnostic Approach to Complex Common Variable Immunodeficiency Patient with Phosphoinositide 3-kinase Catalytic Domain Variant of Uncertain Significance	Rahim Z. Miller, BS	University of South Florida
23	Autoimmune Lymphoproliferative Syndrome or Autoimmune Lymphoproliferative Syndrome-like? Why the Genetic Distinction Matters	Delena Nguyen, MD	University Hospitals Cleveland Medical Center
24	Case Study: IgG deficient patient receiving immune globulin intravenous, human-slra 10% demonstrates improvement in quality of life	Derek Blake, RN, BSN, IgCN	Soleo Health
25	Transient Severe T Cell Lymphopenia in a Patient with Cornelia De Lange Syndrome Captured by TREC Screening	Devyn L. Rohlfs Rivera, MD	Louisiana State University Health Science Center New Orleans
26	Neutralizing Interferon Lambda 1 Autoantibodies in a Pediatric Patient with Chronic Granulomatous Disease and Colitis	Elizabeth N. Karanja, BS	National Institutes of Health

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27	Human ITCH E3 Ubiquitin Protein Ligase (ITCH) Deficiency Syndrome Successfully Treated with Hematopoietic Cell Transplant (HCT)	Emily Neal	University of Pittsburgh School of Medicine
28	Prospective Identification of Inborn Errors of Immunity in a 104-patient Autoimmune Cytopenia Cohort	Emma Westermann-Clark, MD, MA	University of South Florida
29	Looking for ALPS: 10 Years Experience of a Combined Assessment of Serum FASL Levels and Circulating Double Negative T Cells at the CHU Sainte-justine	Fabien Touzot, MD, PhD	Université de Montréal
30	Increased dosage of elapegamase-1v1r improved metabolic and immunologic function in a patient with late-onset adenosine deaminase deficiency and neutralizing anti-drug antibodies	Fathima A. Mohamed, PhD	University of Minnesota Medical School
31	The Neuroimmune Signature Induced by HTLV Infections	Fernando Y N. Vale	Department of Clinical and Toxicological Analyses, University of São Paulo
32	A Case of Chronic Rhinosinusitis with Nasal Polyps in a Patient with a Single Heterozygous RNU4ATAC Pathogenic Variant	Gabrielle Robertson, MD	Division of Allergy & Immunology, Cohen Children's Medical Center/Long Island Jewish Medical Center
33	Spatial Mapping of Immune Cells in Barrier Tissues of Immunocompromised Patients Affected by Human Papillomavirus-related Disease	Glennys V. Reynoso	National Institutes of Health
34	Characterizing a Unique B-cell Precursor Population as a Potential Diagnostic Tool for WHIM Syndrome and APDS	Grace Blair, BS	USF Health Department of Pediatric Allergy and Immunology
35	Traversing Immunodeficiency Complexity: Profound Ureaplasma Infection in Agammaglobulinemia with TCF3 Mutation	Haggar Elbashir, MBBS, MRCP, MSc	Department of Allergy & Clinical Immunology, Lancashire Teaching Hospitals NHS Foundation Trust, Preston, United Kingdom
36	Evolving Clinical Presentation in an Infant with Thrombocytopenia: Insights from a Hispanic Male Baby with Wiskott-Aldrich Syndrome	Hanadys Ale, MD	Memorial Healthcare System
37	Phenotypic Differences in Monochorionic Diamniotic Twins with Chronic Granulomatous Disease	Hannah Shin, MD	University of California San Diego
38	Clinical and Immunologic Phenotype of Prolidase Deficiency	Iris Martin, BA	Laboratory of Clinical Immunology and Microbiology, National Institute of Allergy and Infectious Diseases, NIH

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39	PRF1-Related Isolated CNS Hemophagocytic Lymphohistiocytosis Successfully Treated with Ruxolitinib Monotherapy	Jamie Pruitt, DO	University of Michigan
40	Autoimmunity and Neoplasia in a Cohort of Adult Patients with Inborn Errors of Immunity and the Implications of Having a Positive Molecular Diagnosis	Javier Arroyo-Rodenas, MSc	12 de Octubre University Hospital, Madrid, Spain; 12 de Octubre Research Institute (imas12), Madrid, Spain
41	Neurodevelopmental Outcomes (ND) in Patients with Severe Combined Immunodeficiency (SCID) Following Hematopoietic Cell Transplantation (HCT) in the Era of Newborn Screening. a PIDTC Study	Jennifer M. Puck, MD	University of California, San Francisco
42	Differentiating IPEX- like Syndrome from Other Causes of Autoimmune Enteropathy	Jenny Chen, MD	Duke University Medical Center
43	Biologic Use and Outcomes in CVID Enteropathy	John Hsi-en Ho, MD	Icahn School of Medicine at Mount Sinai
44	JAK1-selective inhibitor upadacitinib induced dynamic immunologic alterations and clinical response in treatment-resistant CVID gastrointestinal disease	John Hsi-en Ho, MD	Icahn School of Medicine at Mount Sinai
45	Relative Location of Prime-boost Immunization Determines Memory B Cell Fate	John S. Barber, MD	Duke University Hospital
46	Seronegative Autoimmune Hepatitis Complicated by Severe Aplastic Anemia	Jonathan Lim, DO	Cleveland Clinic Children's
47	Demyelinating CNS Disease: An Unusual Complication of XLA	Jose G. Calderon, MD	Nicklaus Children's Hospital
48	A Primary Neurological Presentation of CASP-8 Deficiency State	Karl Mueller, KM	Mayo Clinic
49	Identification of a Novel XIAP Pathogenic Variant Associated with XIAP Deficiency Presenting as VEOIBD	Katheryn Schloss, MD, MPH	UPMC
50	Hematopoietic Stem Cell Transplantation for Very Early Onset Inflammatory Bowel Disease Caused by IL10RA Deficiency – Preparation Is the Key to Success	Kavitha Ganesan, MD	Apollo hospitals , Chennai
51	Assessment and Characterization of Tbet+ B Cells in Various Inborn Errors of Immunity (IEIs)	Kranthi Nomula, MD	Division of Pediatric Allergy/Immunology, University of South Florida at Johns Hopkins All Children's Hospital, St. Petersburg, FL, USA.
52	A Subtle Clinical Presentation of LAD1 Deficiency	Latoya Jeanpierre, MD	Children's Hospital of Philadelphia
53	A Peculiar Presentation: STAT3 Loss of Function with Recurrent Osteomyelitis and Review of the Literature	Laura Bou-Maroun, MD	University of Michigan

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54	Case of Mitochondrial Encephalomyopathy Secondary to COVID-19 in a Pediatric Case of SIFD Syndrome with a Novel TRNT1 Mutation	Madeline Schutt, MD	Ann & Robert H. Lurie Children's Hospital
55	Comprehensive Characterization of Innate and Adaptive Immune Profile of COVID-19 Early Convalescent Children	Magda Carneiro-Sampaio, MD, PhD	Faculdade de Medicina da USP/ CoBEII
56	Digeorge Syndrome with an Atypical Presentation: Lymphoproliferation and Malignancy	Magda Carneiro-Sampaio, MD, PhD	Faculdade de Medicina da USP/ CoBEII
57	Resolving Variants of Unknown Significance (VUS) in PIK3CD and PIK3R1	Manish Butte, MD, PhD	UCLA
58	Ocrelizumab Induced B-cell Depletion in a Newborn Male	Maria A. Sacta, MD, PhD	Children's Hospital of Philadelphia
59	Characterization of CD4+ T lymphocytes in STAT3-DN hyper-IgE syndrome patients	Maria Carrabba, MD, PhD	Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico of Milan, Italy
60	Immune response after anti-SARS-CoV-2 mRNA repeated boosters vaccination in patients with Common Variable Immunodeficiency	Maria Carrabba, MD, PhD	Fondazione IRCCS Ca' Granda Ospedale Maggiore Policlinico of Milan, Italy
61	The Challenges of Recognition and Diagnosis of APDS2 in a Family with Novel PIK3R1 Variant	Maria F. Villavicencio, MD	University of Kansas School of Medicine - Wichita
62	Expansion of Tbet+ T and B Cells in the Lung of Hypomorphic Rag1 Mice	Marta Toth, PhD	Department of Pediatrics, Division of Allergy & Immunology, University of South Florida
63	Age-associated Distribution of Th Subsets in Blood of Locid vs CVID Patients	Martin Perez-Andres, PhD	Department of Medicine, Cancer Research Centre (IBMCC, USAL-CSIC), Cytometry Service (NUCLEUS), University of Salamanca (USAL)
64	Celiac Disease in Selective IgA Deficiency Patients Is Associated with T-cell Defects	Martin Perez-Andres, PhD	Department of Medicine, Cancer Research Centre (IBMCC, USAL-CSIC), Cytometry Service (NUCLEUS), University of Salamanca (USAL)
65	Late Onset and Long Lasting Neutropenias: Preliminary Data on WES Analysis	Maurizio Miano, MD	Hematology Unit- IRCCS Istituto Giannina Gaslini. Genoa, Italy
66	Provider Perspectives of Long-term Follow-up Care of Patients with Severe Combined Immunodeficiency	Morna Dorsey, MD, MMSc	University of California, San Francisco
67	Incidental Diagnosis of NFKB2 Mutation in Patient with ACTH Deficiency and Low IgA	Navya Kartha, DO	Akron Children's Hospital

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68	Inherited Pomp-related Autoinflammation and Immune Dysregulation Disease Treated with Baricitinib Prior to Hematopoietic Stem Cell Transplant	Nicholas Hartog, MD	Corewell Health Allergy and Immunology
69	Thymus Hypoplasia in 22q11.2DS (DiGeorge): From Mechanism to Restoration	Nicolai van Oers, PhD	UT Southwestern Medical Center
70	Chronic Granulomatous Disease: A Clinical or a Molecular Diagnosis?	Nikhil Chowdary Peddi, MD	St Barnabas Hospital, SBH Health System
71	Patient with Trisomy 21 and Congenital Chylothorax with Secondary Hypogammaglobulinemia	Nikhil Chowdary Peddi, MD	St Barnabas Hospital, SBH Health System
72	A Quality Improvement Approach for Transition of Immunodeficiency Patients to Adult Healthcare	Nikita Raje, MBBS, MSc	Children's Mercy Kansas City
73	Off-label Treatment with the Selective PI3K δ Inhibitor Leniolisib in 2 Pediatric Patients with Activated Phosphoinositide 3-kinase Delta Syndrome 2 (APDS2)	Olaf Neth, MD	Hospital Infantil Universitario Virgen del Rocio Instituto de Biomedicina de Sevilla (IBiS)
74	Exploring Molecular Landscapes and Immunological Dynamics in Glioblastoma: Insights from Oncolytic Virotherapy	Pedro M. Barcelos	Department of Clinical and Toxicological Analyses, University of São Paulo
75	Unique Presentation of Mycobacterium Haemophilum, Associated with T-cell Dysfunction After Nelarabine Administration	Pooja Kapadia, MD	Division of Allergy & Immunology, Cohen Children's Medical Center/Long Island Jewish Medical Center
76	Disseminated Cryptococcal Lymphadenitis in a Patient with stat1 Gain-of-function: Diagnostic and Therapeutic Challenges	Roberta Romano, MD	University of Naples Federico II, Translational medical sciences, Section of Pediatrics, Naples, Italy
77	Use of Ruxolitinib for Patients with Hyperinflammatory Syndromes - A Real-world Experience Pilot Study	Rohith Jesudas, MD	St. Jude Children's Research Hospital
78	Novel Hypomorphic BTK Variant in X-linked Agammaglobulinemia of a Kindred	Ruth S. Cajuste, BS	USF Health Department of Pediatric Allergy and Immunology
79	SARS-CoV-2 Breakthrough Infection Increases the Adaptive Immune Responses of Vaccinated Immunosuppressed Children	Samuel Sassine, MD, MSc	Université de Montréal
80	From Pancytopenia to Hyperleukocytosis, an Unexpected Presentation of Immune Reconstitution Inflammatory Syndrome in an Infant with Methylmalonic Acidemia	Samuel Sassine, MD, MSc	Université de Montréal
81	Neutropenia and Myelodysplasia Without Syndromic Features in a 5-year-old Boy with Novel SAMD9 Variant	Sarah Chamseddine, MD	Boston Children's Hospital
82	Disseminated Histoplasmosis in an Adult with STAT1 Mutation	Sarah Nimri, MD	University of Iowa Hospitals and Clinics.

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83	Anti-IL12 autoantibodies in a teenage girl with multiple recurrent abscesses	Saul O. Lugo Reyes, MD, MS	National Institute of Pediatrics
84	Two siblings with autoinflammation and combined immunodeficiency due to autosomal recessive RNF31-loss of function	Saul O. Lugo Reyes, MD, MS	National Institute of Pediatrics
85	Complex Acquired Angioedema Case in a 46-year-old Emirati Female with a History of Breast Cancer	Shuayb Elkhailifa, MBBS, MRCP, FRCPath	Cleveland Clinic Abu Dhabi and University of Manchester
86	The Use of Tofacitinib and Upadacitinib in Very Early Onset IBD	Smridhi Mahajan, MD	University of Texas Southwestern Medical Center
87	Long Term Management of Transplanted Patients with Chronic Granulomatous Disease, Wiskott-aldrich Syndrome, and Primary Immune Regulatory Disorders: A PIDTC Survey	Stephanie Si Lim, MD, MBA	University of Hawaii Cancer Center
88	Helper T Cell Immunity in Humans with Inherited CD4 Deficiency	Stuart G Tangye, PhD	Garvan Institute
89	Mycobacteria Chelonae Panniculitis Presenting as Refractory Cellulitis in Patient with Hypogammaglobulinemia	Sushmitha Boppana, MD	Cleveland Clinic Akron General
90	Subtle Presentation of Fungal Disease in X-CGD	Talia Arceri, MD	Children's Hospital of Philadelphia
91	A Non-consanguineous Family with Hepatic Veno-occlusive Disease and Immunodeficiency	Thais Moura	University of Sao Paulo
92	Treatable Acute Neuroinflammatory Disease Associated with Complement Factor I Loss-of-function in the Plain Community	Whitney Reid, MD	Children's Hospital of Philadelphia
93	Otulin-related Conditions: Report of a New Case and Review of the Literature Using Genia	Xiao P. Peng, MD, PhD	Director of the Genetics of Blood and Immunity Clinic; Department of Genetic Medicine, Johns Hopkins University School of Medicine
94	Biallelic Splice Variants in NHEJ1 Deficiency Causing Primary Hematologic and Oncologic Manifestations: A Tale of Two Patients	Xiao P. Peng, MD, PhD	Director of the Genetics of Blood and Immunity Clinic; Department of Genetic Medicine, Johns Hopkins University School of Medicine
95	The Neuroimmunological Network in Cancer-induced HPV Infections	Yohan Lucas Gonçalves Corrêa	Universidade de São Paulo
96	Follow up of Cell-immune Response Using Covidcell DTH Skin Test in Kidney Transplanted Patients	Yvelise Barrios, MD, PhD	Universidad de La Laguna