Congress poster

Registration code	Abstract Title		Author	
RN 10001	Clinical features and molecular characteristics of seven patients with chronic mucocutaneous	Gao		Hong
	candidiasis caused by autosomal dominant STAT1 gene mutations			
RN 10002	Inducible nitric oxide synthase and systemic lupus erythematosus	Pan		Lu
RN 10003	A single center analysis of henoch-schonlein purpura nephritis above grade III in children	Feng		Dan
RN 10004	Variation in IL-21-secreting T follicular helper cells in Kawasaki disease	Xu		Meng
RN 10005	Clinical and Molecular Features of Fifteen Children with Hyper-IgE Syndrome in Mainland China	Xiang		Qing qing
RN 10007	Nine cases of Inflammatory Bowel Disease in 5 pedigrees with IL10RA mutaitons	Wang		Wei
RN 10010	Treatment of leukocyte adhesion deficiency type I with umbilical cord blood transplantation: report of two cases and review of literatures	Zhai	Xiao	wen
RN 10011	The role of IL-17 in lung inflammation infected by hMPV	Chen		Xin
RN 10012	A retrospective single-center study: Clinical analysis of macrophage activation syndrome in pediatric patients with autoimmune diseases	He	Sheng	nan
RN 10588	Four patients with Atypical UNC13D-Related Hemophagocytic Lymphohistiocytosis Characterized by humoral immunodeficiency and nervous system Involvement	Zhao		Qin
RN 15060	Lymphnoditis and ARDS after BCG vaccination: 1 case report	Zhang		Wei
RN 16514	Chronic Granulomatous Disease presenting as repeated fever in a 11-months-old child	chen	hai	хіа
RN 16634	Gastrointestinal tract injury and clinical characteristics in 172 children with Henoch-Schonlein purpura checked by gastroscope	zeng	hua	song
RN 16862	Primary IqG1 and IqG3 subclass deficiency and JRA in a 40 -month-old Vietnamese girl: a case report	Nguyen	Thuv	Duong
RN 18640	Gene therapy for X-linked severe combined immunodeficiency disease	Hu		Mingfeng
RN 19173	The clinical analysis of Kawasasi Disease associated with macrophage activation syndrome and the diagnostic criteria exploration	He	Sheng	nan
RN 28490	A case of Interferon y receptor 1 heterozygous mutation complicated with recurrent mycobacteria infection	Jiang	Criorig	Min
RN 31480	Respiratory Pathogens Analysis Among Children Hospitalized of Chongqing with Acute Asthma	zhang		yao
RN 33576	Hyperuricemia and Primary Hypertension in Children	Di	Ya	zhen
RN 34763	Diagnostic approaches for primary immunodeficiency disease from Sanger DNA sequencing to whole exome sequencing	Chung Yin		Wong
RN 36354	Clinical characteristics of childhood onset systemic lupus erythematosus at different onset ages	wang		li
RN 37359	THE ROAD LESS TRAVELLED: A CASE-REPORT ON INTESTINAL BEHCET'S DISEASE	Glaiza	Metrillo	
	*		Menino	Madulara
RN 37493 RN 37708	Felty's syndrome in a child: a case report	Song		Lei
	Reduced B cell Maturation Antigen (BCMA) Expression in B-cells of CVID Patients	Reza	V:	Yazdani
RN 37713	Advances in research of high mobility group box1 in systemic lupus erythematosus	Ruan	Xiao	xia
RN 41283	A case of autoimmune lymphoproliferative syndrome in children	lin		qiang
RN 44822	Study of RSV and RV detection rate, clinical characteristics and follow-up of 457 outpatient children with acute lower respiratory tract infection	Gao		Yu
RN 47101	Hypersensitive Pneumonitis: An Initial Presentation of Chronic Granulomatous Disease in a Child	Liu		Hui
RN 48740	Clinical, Genetic and Immunological Characteristics of 18 Chinese Patients with X-linked lymphoproliferative Syndrome	Xu		Tao
RN 53517	First Report of Primary Immunodeficiency Diseases from a Resource Limited Setting in Hills of North-India	Avinash	_	Sharma
RN 54126	Novel NLR-family CARD-containing protein 4 (NLRC4) mutation causes Familial Cold Autoinflammatory Syndrome (FCAS) in a pediatric patient	Yu	Zhong	xun
RN 54979	A meta analysis of drug therapy for Henoch-Schonlein purpura in children	Zhao	Meng	Jia
RN 57878	Profile of fungal infections in children with chronic granulomatous disease (CGD) at Chandigarh, North India	Dharmaga		Bhattarai
RN 59193	Dock8 regulates BCR signaling and activation of memory B cells via WASP and CD19	Sun	Xiao	yu
RN 61260	A Case report of child with STING associated vasculopathy with onset in infancy (SAVI)	chen		li
RN 66048	A Boy with Recurrent Severe Pneumonia	AYE MYA	SANDAR	
RN 66830	Adjunctive vitamin D for treatment of active juvenile idiopathic arthritis: an open-label, randomized controlled trial	tang		tao
RN 69492	A report of X-linked Agammaglobulinemia with severe hand-foot-mouth disease	MA		Yan
RN 72879	Human CTL-based functional analysis demonstrates the reliability of a munc13-4 protein expression assay for FHL3 diagnosis	Ryuta		Nishikomor
RN 77573	Mutation of IL-12RB1 in 2 children with disseminated BCG infection	Wei	Qi	jiao
RN 78012	CHRONIC EPSTEIN-BARR VIRUS (EBV) RELATED ACUTE-ON-CHRONIC LIVER FAILURE: A CASE STUDY	Ling		Yun
RN 78276	Immunodeficiency, Centromeric Instability, and Facial Anomalies Syndrome (DNMT3B deficiency) with clinical presentations mimicking Omenn syndrome	Athipat		Athipongar
RN 82605	Clinical Analysis of HLA-B27-positive Enthesitis Related Arthritis in 70 Cases	Wen	Qiu	Yue
RN 83474	SCID Case Sharing	Wang		Jia
RN 85920	Good Syndrome, Bad outcome: A case of thymoma with immunodeficiency	Kristy	Monte	Pinoy
RN 86556	A boy with recurrent abscess	SU	LINN	HLAING
RN 95289	Novel CD40L gene variants in single centre cohort of X-linked Hyper IgM patients from North India	BABU	MATHEW	PANNERS
RN 96496	High IgM syndrome with pulmonary infection in one pediatric cases	Fan		Liu
RN 97133	A report and literature review of one case of primary immunodeficiency disease caused by POLE1 gene mutation	Liang		Ming
RN 98100	Epidemiological characteristics of nasopharyngeal Streptococcus pneumonia strains among children with lower respiratory tract infection in Chongqing, China	Yu	Yi	yi

