Amit Rawat

List of Publications by Year
in descending order

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261 papers	3,252 citations	218662 26 h-index	243610 44 g-index
275 all docs	275 docs citations	275 times ranked	4653 citing authors

#	Article	IF	Citations
1	T Cell Abnormalities in X-Linked Agammaglobulinaemia: an Updated Review. Clinical Reviews in Allergy and Immunology, 2023, 65, 31-42.	6.5	3
2	Utility of Immunohistochemistry and Immunofluorescence in Determining the Pathogenic Variants of Chronic Granulomatous Disease. Journal of Clinical Immunology, 2022, 42, 85-93.	3.8	1
3	Clinical and Genetic Spectrum of Inborn Errors of Immunity in a Tertiary Care Center in Southern India. Indian Journal of Pediatrics, 2022, 89, 233-242.	0.8	4
4	Features of nephrotic syndrome in infants with severe combined immunodeficiency. Journal of Allergy and Clinical Immunology: in Practice, 2022, 10, 356-357.	3.8	0
5	Microalbuminuria and Urinary Neutrophil Gelatinase-associated Lipocalin (uNGAL) in human immunodeficiency virus infected children. Indian Journal of Nephrology, 2022, 32, 22.	0.5	0
6	Recurrent pneumonia in a child: Knitting clinical and radiological features to clinch the diagnosis. , 2022, 2, 61.		0
7	Immunoglobulin Profile and Lymphocyte Subsets in Preterm Neonates. Indian Pediatrics, 2022, 59, 214-217.	0.4	1
8	Atypical Wiskottâ€“Aldrich syndrome without thrombocytopenia partially responding to omalizumab therapy. Clinical and Experimental Dermatology, 2022, , .	1.3	1
9	Unusual clinical manifestations and predominant stopgain ATM gene variants in a single centre cohort of ataxia telangiectasia from North India. Scientific Reports, 2022, 12, 4036.	3.3	1
10	An Autopsy Case of Wiskott-Aldrich Syndrome Revealing â€œFDC-Only Lymphoid Folliclesâ€•in Lymphoid Tissue: A Morphologic Correlate of Defective Immune Synapse. Pediatric and Developmental Pathology, 2022, , 109352662110583.	1.0	1
11	Cutaneous involvement in <scp>DOCK8</scp> â€related immunodeficiency syndrome responding to thalidomide. Dermatologic Therapy, 2022, 35, e15491.	1.7	1
12	Novel TBXAS1 variants in two Indian children with Ghosal hematodiaphyseal dysplasia: A concise report. European Journal of Medical Genetics, 2022, 65, 104498.	1.3	2
13	Pediatric systemic lupus erythematosus: phagocytic defect and oxidase activity of neutrophils. Pediatric Research, 2022, 92, 1535-1542.	2.3	2
14	Immunoglobulin Profile and Lymphocyte Subsets in Preterm Neonates.. Indian Pediatrics, 2022, , .	0.4	0
15	Lymphoproliferation in Inborn Errors of Immunity: The Eye Does Not See What the Mind Does Not Know. Frontiers in Immunology, 2022, 13, .	4.8	5
16	Deficiency of Human Adenosine Deaminase Type 2 â€“ A Diagnostic Conundrum for the Hematologist. Frontiers in Immunology, 2022, 13, 869570.	4.8	13
17	Mechanisms of Immune Dysregulation in COVID-19 Are Different From SARS and MERS: A Perspective in Context of Kawasaki Disease and MIS-C. Frontiers in Pediatrics, 2022, 10, .	1.9	6
18	Association of SNP (rs1042579) in thrombomodulin gene and plasma thrombomodulin level in North Indian children with Kawasaki disease. Molecular Biology Reports, 2022, 49, 7399-7407.	2.3	2

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19	063â€... Anti-MOG and anti AQP4 antibodies in pediatric acquired memyelinating syndromes (ADS): an Indian cohort. Journal of Neurology, Neurosurgery and Psychiatry, 2022, 93, A32.1-A32.	1.9	0
20	Utility of targeted next generation sequencing for inborn errors of immunity at a tertiary care centre in North India. Scientific Reports, 2022, 12, .	3.3	7
21	Adipocytokine profile in children with Kawasaki disease at a mean follow-up period of 5.5 years: A study from North India. World Journal of Clinical Pediatrics, 2022, 11, 360-368.	2.1	2
22	Healing With Complication: An Unusual Case of Nasal Tip Ulceration in Leukocyte Adhesion Deficiency Type 1. Journal of Allergy and Clinical Immunology: in Practice, 2022, , .	3.8	0
23	Macrophage activation syndrome in children with Kawasaki disease: an experience from a tertiary care hospital in northwest India. Rheumatology, 2021, 60, 3413-3419.	1.9	20
24	Poor allograft outcome in Indian patients with post-transplant C3 glomerulopathy. CKJ: Clinical Kidney Journal, 2021, 14, 291-300.	2.9	2
25	LINEZOLID-INDUCED MITOCHONDRIAL TOXICITY PRESENTING AS RETINAL NERVE FIBER LAYER MICROCYSTS AND OPTIC AND PERIPHERAL NEUROPATHY IN A PATIENT WITH CHRONIC GRANULOMATOUS DISEASE. Retinal Cases and Brief Reports, 2021, 15, 224-229.	0.6	12
26	A young female with early onset arthritis, uveitis, hepatic, and renal granulomas: a clinical tryst with Blau syndrome over 20Âyears and case-based review. Rheumatology International, 2021, 41, 173-181.	3.0	22
27	Haemolytic Uremic Syndrome Associated with Citrobacter freundii in a Young Boy with X-Linked Agammaglobulinemia. Journal of Clinical Immunology, 2021, 41, 227-229.	3.8	1
28	A 5â€yearâ€old boy with Kawasaki disease shock syndrome, myocarditis and macrophage activation syndrome. Journal of Paediatrics and Child Health, 2021, 57, 1312-1315.	0.8	3
29	A neonate with absent lymphocytes: Is this severe combined immunodeficiency?. Pediatric Hematology Oncology Journal, 2021, 6, 57-59.	0.1	0
30	Hemophagocytic Lymphohistiocytosis in Children with Chronic Granulomatous Diseaseâ€"Single-Center Experience from North India. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 771-782.e3.	3.8	7
31	Novel SERPING1 gene mutations and clinical experience of type 1 hereditary angioedema from North India. Pediatric Allergy and Immunology, 2021, 32, 599-611.	2.6	12
32	Infection Due to Serratia sp. in Chronic Granulomatous Diseaseâ€"Is the Incidence Low in Tropical Countries?. Journal of Clinical Immunology, 2021, 41, 486-490.	3.8	2
33	Sensitization to AÂfumigatus in subjects with nonâ€cystic fibrosis bronchiectasis. Mycoses, 2021, 64, 412-419.	4.0	12
34	Chronic Granulomatous Disease: A Perspective from a Developing Nation. International Archives of Allergy and Immunology, 2021, 182, 360-364.	2.1	2
35	Cutaneous IgA vasculitisâ€"presenting manifestation of a novel mutation in the IKZF1 gene. Rheumatology, 2021, 60, e101-e103.	1.9	3
36	Deforming Polyarthritis in a North Indian Familyâ€"Clinical Expansion of STING-Associated Vasculopathy with Onset in Infancy (SAVI). Journal of Clinical Immunology, 2021, 41, 209-211.	3.8	3

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37	Splice-site mutation in COPA gene and familial arthritis â€“ a new frontier. Rheumatology, 2021, 60, e7-e9.	1.9	5
38	Catch the thief by its marks: inverse Gottron papules, interstitial lung disease, anti MDA-5 antibody positivity in juvenile dermatomyositis. Rheumatology, 2021, 60, e56-e58.	1.9	2
39	Epsteinâ€Barr virusâ€associated lymphocytic cholangitis in a child with Xâ€linked lymphoproliferative syndrome. Scandinavian Journal of Immunology, 2021, 93, e12975.	2.7	1
40	Pericardial effusion in anti-complement factor H antibody-associated atypical hemolytic uremic syndrome: two case reports. CEN Case Reports, 2021, 10, 255-260.	0.9	0
41	Aspergillus fumigatus Skull Bone Osteomyelitis and Native Valve Endocarditis in a Young Boy: an Unusual Presentation of Chronic Granulomatous Disease. Journal of Clinical Immunology, 2021, 41, 814-816.	3.8	4
42	Liver Abscess in Chronic Granulomatous Diseaseâ€”Two Decades of Experience from a Tertiary Care Centre in North-West India. Journal of Clinical Immunology, 2021, 41, 552-564.	3.8	7
43	Infection triggered anti complement factor H (CFH) positive atypical Hemolytic Uremic Syndrome in children: Âlessons for the clinical nephrologist. Journal of Nephrology, 2021, 34, 943-947.	2.0	1
44	Pathophysiology of Hereditary Angioedema (HAE) Beyond the SERPING1 Gene. Clinical Reviews in Allergy and Immunology, 2021, 60, 305-315.	6.5	19
45	Autoinflammatory disorders. , 2021, , 245-290.		0
46	Importance of Morphology in the Era of Molecular Biology: Lesson Learnt from a Case of Chediakâ€“Higashi Syndrome. Indian Journal of Hematology and Blood Transfusion, 2021, 37, 517-519.	0.6	2
47	Clinical, Immunological, and Molecular Profile of Chronic Granulomatous Disease: A Multi-Centric Study of 236 Patients From India. Frontiers in Immunology, 2021, 12, 625320.	4.8	31
48	Extensive Molluscum Contagiosum in X-Linked Agammaglobulinemia. Journal of Allergy and Clinical Immunology: in Practice, 2021, 9, 985.	3.8	3
49	Clinical Profile of Hyper-IgE Syndrome in India. Frontiers in Immunology, 2021, 12, 626593.	4.8	13
50	Clinical and Molecular Findings in Mendelian Susceptibility to Mycobacterial Diseases: Experience From India. Frontiers in Immunology, 2021, 12, 631298.	4.8	36
51	X-Linked Agammaglobulinemia With Chronic Meningoencephalitis: A Diagnostic Challenge. Indian Pediatrics, 2021, 58, 169-175.	0.4	2
52	Human leukocyte antigen B27 and B57 alleles in HIV-infected long-term nonprogressor children. Aids, 2021, 35, 703-705.	2.2	0
53	Monocyte platelet aggregates in children with Kawasaki disease- a preliminary study from a tertiary care centre in North-West India. Pediatric Rheumatology, 2021, 19, 25.	2.1	4
54	Spectrum of Systemic Auto-Inflammatory Diseases in India: A Multi-Centric Experience. Frontiers in Immunology, 2021, 12, 630691.	4.8	11

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55	The Spectrum of Clinical, Immunological, and Molecular Findings in Familial Hemophagocytic Lymphohistiocytosis: Experience From India. Frontiers in Immunology, 2021, 12, 612583.	4.8	7
56	Autoantibody Profile of Children with Juvenile Dermatomyositis. Indian Journal of Pediatrics, 2021, 88, 1170-1173.	0.8	6
57	Wiskott Aldrich Syndrome: A Multi-Institutional Experience From India. Frontiers in Immunology, 2021, 12, 627651.	4.8	16
58	Skewed TCR Alpha, but not Beta, Gene Rearrangements and Lymphoma Associated with a Pathogenic TRAC Variant. Journal of Clinical Immunology, 2021, 41, 1395-1399.	3.8	4
59	Congenital Rubella: A Salient Cause of Congenital Heart Defects in Infants. Journal of Tropical Pediatrics, 2021, 67, .	1.5	4
60	False-positive HIV serology, Candida lusitaniae pneumonia, and a novel mutation in the CYBB gene. Immunobiology, 2021, 226, 152110.	1.9	2
61	Achromobacter xylosoxidans Pneumonia in a Young Child with Chronic Granulomatous Diseaseâ€"a Case-Based Review. Journal of Clinical Immunology, 2021, 41, 1686-1692.	3.8	1
62	Immunoglobulins and Lymphocyte Subsets in Children with Infantile Tremor Syndrome. Indian Journal of Pediatrics, 2021, 88, 1139-1141.	0.8	0
63	Autoimmunity in Wiskottâ€"Aldrich Syndrome: Updated Perspectives. The Application of Clinical Genetics, 2021, Volume 14, 363-388.	3.0	15
64	Multidrug-Resistant Nontyphoidal Salmonella Associated with Invasive Disease in an Immunocompetent Child. Indian Journal of Pediatrics, 2021, 88, 1266-1266.	0.8	3
65	Hematologically important mutations: X-linked chronic granulomatous disease (fourth update). Blood Cells, Molecules, and Diseases, 2021, 90, 102587.	1.4	22
66	An updated review on Mendelian susceptibility to mycobacterial diseasesâ€" a silver jubilee celebration of its first genetic diagnosis. Expert Review of Clinical Immunology, 2021, 17, 1103-1120.	3.0	7
67	Reticular dysgenesis exacerbated by hemophagocytic lymphohistiocytosis and the presence of unusual histiocyte-like cells in bone marrow. Immunobiology, 2021, 226, 152143.	1.9	1
68	Hematologically important mutations: The autosomal forms of chronic granulomatous disease (third) Tj ETQq0 0 0 rgBT /Overlock 10 Tf	1.4	22
69	Clinical Characteristics of Children With Celiac Disease Not Responding to Hepatitis B Vaccination in India. JPGN Reports, 2021, 2, e046.	0.4	1
70	Mystery of a Family with Recurrent Male Infant Deaths- Solved by Autopsy and Molecular Tests. Indian Journal of Pediatrics, 2021, 88, 257-262.	0.8	0
71	Serial urinary neutrophil gelatinase associated lipocalin in pediatric diabetic ketoacidosis with acute kidney injury. Clinical Diabetes and Endocrinology, 2021, 7, 20.	2.7	4
72	Expression of CD40 Ligand on T Cells and Soluble CD40 Ligand in Children With Kawasaki Disease. Journal of Clinical Rheumatology, 2021, 27, 194-200.	0.9	3

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73	Hemophagocytic Lymphohistiocytosis in a PICU of a Developing Economy: Clinical Profile, Intensive Care Needs, Outcome, and Predictors of Mortality. Pediatric Critical Care Medicine, 2021, 22, e44-e57.	0.5	6
74	Hair microscopy: an easy adjunct to diagnosis of systemic diseases in children. Applied Microscopy, 2021, 51, 18.	1.4	4
75	Persistent Pneumonia in an Infant. Indian Pediatrics, 2021, 58, 1067-1073.	0.4	1
76	Early diagnosis of Wiskott-Aldrich syndrome in the neonatal period and successful haematopoietic stem cell transplant in infancy. Current Medicine Research and Practice, 2021, 11, 288.	0.1	0
77	Phenomic Analysis of Chronic Granulomatous Disease Reveals More Severe Integumentary Infections in X-Linked Compared With Autosomal Recessive Chronic Granulomatous Disease. Frontiers in Immunology, 2021, 12, 803763.	4.8	3
78	Clinical profile, long-term follow-up and outcome of juvenile systemic scleroderma: 25 years of clinical experience from North-West India. Clinical and Experimental Rheumatology, 2021, 39 Suppl 131, 149-156.	0.8	0
79	Persistent Pneumonia in an Infant. Indian Pediatrics, 2021, 58, 1067-1073.	0.4	0
80	Clinical profile, long-term follow-up and outcome of juvenile systemic scleroderma: 25 years of clinical experience from North-West India. Clinical and Experimental Rheumatology, 2021, 39, 149-156.	0.8	1
81	Association of ITPKC gene polymorphisms rs28493229 and rs2290692 in North Indian children with Kawasaki disease. Pediatric Research, 2021, , .	2.3	2
82	Scrotal and Penile Ulcer in Juvenile Dermatomyositis. Journal of Clinical Rheumatology, 2020, 26, e7-e8.	0.9	2
83	Kikuchi-Fujimoto Disease: An Under Recognized Cause of Fever with Lymphadenopathy. Indian Journal of Pediatrics, 2020, 87, 85-85.	0.8	3
84	Current status and prospects of primary immunodeficiency diseases in Asia. Genes and Diseases, 2020, 7, 3-11.	3.4	25
85	Leukocyte adhesion defect: Where do we stand circa 2019?. Genes and Diseases, 2020, 7, 107-114.	3.4	34
86	Platelets in Kawasaki disease: Is this only a numbers game or something beyond?. Genes and Diseases, 2020, 7, 62-66.	3.4	25
87	An updated review on phenocopies of primary immunodeficiency diseases. Genes and Diseases, 2020, 7, 12-25.	3.4	13
88	An updated review on activated PI3 kinase delta syndrome (APDS). Genes and Diseases, 2020, 7, 67-74.	3.4	30
89	Recent advances in elucidating the genetics of common variable immunodeficiency. Genes and Diseases, 2020, 7, 26-37.	3.4	37
90	Recent advances in chronic granulomatous disease. Genes and Diseases, 2020, 7, 84-92.	3.4	40

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91	Genetics of severe combined immunodeficiency. Genes and Diseases, 2020, 7, 52-61.	3.4	42
92	An update on the genetics and pathogenesis of hereditary angioedema. Genes and Diseases, 2020, 7, 75-83.	3.4	43
93	Catastrophes due to missing complements: C1q deficiency lupus with Kikuchi-Fujimoto disease and macrophage activation syndrome. Rheumatology, 2020, 59, 1778-1780.	1.9	2
94	0.9% saline versus Plasma-Lyte as initial fluid in children with diabetic ketoacidosis (SPinK trial): a double-blind randomized controlled trial. Critical Care, 2020, 24, 1.	5.8	528
95	Outcome of C3 glomerulopathy patients: largest single-centre experience from South Asia. Journal of Nephrology, 2020, 33, 539-550.	2.0	6
96	Longâ€term outcome in children with juvenile dermatomyositis: A singleâ€center study from north India. International Journal of Rheumatic Diseases, 2020, 23, 392-396.	1.9	15
97	A Neonate With Fungal Lung Nodules Mimicking Pulmonary Malignancy. Pediatric Infectious Disease Journal, 2020, 39, e474-e475.	2.0	0
98	Serum Ferritin Predicts Neither Organ Dysfunction Nor Mortality in Pediatric Sepsis Due to Tropical Infections. Frontiers in Pediatrics, 2020, 8, 607673.	1.9	12
99	Seronegative panencephalitis complicated by viral encephalomyelitis in a case of Goodâ€™s syndrome â€“ a neuropathological report. International Journal of Neuroscience, 2020, , 1-6.	1.6	2
100	Epidermal necrolysis as the presenting manifestation of pediatric lupus. Pediatric Dermatology, 2020, 37, 1119-1124.	0.9	2
101	Anti-complement factor I antibody associated atypical hemolytic uremic syndrome â€“ A new insight for future perspective!. Immunobiology, 2020, 225, 152000.	1.9	8
102	Successful perioperative management of three patients with hereditary angioedema without C1 esterase inhibitor therapy: A developing country perspective. Immunobiology, 2020, 225, 152022.	1.9	3
103	X-Linked Thrombocytopenia and Vanishing White Matter Disease in a Child: Double Tragedy. Journal of Clinical Immunology, 2020, 40, 1176-1180.	3.8	0
104	Nocardiosis Associated with Primary Immunodeficiencies (Nocar-DIP): an International Retrospective Study and Literature Review. Journal of Clinical Immunology, 2020, 40, 1144-1155.	3.8	11
105	Refractory Autoimmune Cytopenia in a Young Boy with a Novel LRBA Mutation Successfully Managed with Sirolimus. Journal of Clinical Immunology, 2020, 40, 1184-1186.	3.8	7
106	Intracranial Aneurysm Biomarker Candidates Identified by a Proteome-Wide Study. OMICS A Journal of Integrative Biology, 2020, 24, 483-492.	2.0	14
107	An uncommon overlap of two common rheumatological disorders. Lupus, 2020, 29, 1121-1125.	1.6	1
108	CT Coronary Angiography Studies After a Mean Follow-up of 3.8 Years in Children With Kawasaki Disease and Spontaneous Defervescence. Frontiers in Pediatrics, 2020, 8, 274.	1.9	11

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109	Immunogenetics of Kawasaki disease. Clinical Reviews in Allergy and Immunology, 2020, 59, 122-139.	6.5	73
110	Revisiting the complement system in systemic lupus erythematosus. Expert Review of Clinical Immunology, 2020, 16, 397-408.	3.0	25
111	Kawasaki Disease in Children Older Than 10 Years: A Clinical Experience From Northwest India. Frontiers in Pediatrics, 2020, 8, 24.	1.9	17
112	A young girl with hypogammaglobulinemia and granulomatous hepatitis caused by a novel mutation in ZBTB24 gene: A case based analysis. Immunobiology, 2020, 225, 151912.	1.9	6
113	Clinico-laboratory profile of Kawasaki disease with arthritis in children. European Journal of Pediatrics, 2020, 179, 875-879.	2.7	11
114	Clinical, Immunological, and Molecular Features of Severe Combined Immune Deficiency: A Multi-Institutional Experience From India. Frontiers in Immunology, 2020, 11, 619146.	4.8	31
115	Clinical and Genetic Profile of X-Linked Agammaglobulinemia: A Multicenter Experience From India. Frontiers in Immunology, 2020, 11, 612323.	4.8	16
116	Transient Erythroblastopenia. Journal of Clinical Rheumatology, 2020, Publish Ahead of Print, .	0.9	1
117	Clinical and Genetic Spectrum of a Large Cohort of Patients With Leukocyte Adhesion Deficiency Type 1 and 3: A Multicentric Study From India. Frontiers in Immunology, 2020, 11, 612703.	4.8	24
118	Case Report: Ceftriaxone-Resistant Invasive Salmonella Enteritidis Infection with Secondary Hemophagocytic Lymphohistiocytosis: A Contrast with Enteric Fever. American Journal of Tropical Medicine and Hygiene, 2020, 103, 2515-2517.	1.4	1
119	Acute Lung Infiltrates in a Child With Chronic Granulomatous Disease: Is It Voriconazole-Induced Lung Injury?. Journal of Pediatric Pharmacology and Therapeutics, 2020, 25, 465-466.	0.5	1
120	An Infant with Suppurative Adenitis, Nonhealing Wound, and Perianal Sinus. Journal of Allergy and Clinical Immunology: in Practice, 2019, 7, 290-291.	3.8	0
121	An unusual cause of deforming erosive arthritis in an adult. Rheumatology, 2019, 59, 602.	1.9	3
122	Myriad Faces of Chronic Granulomatous Disease: All in an Indian Family with Novel CYBB Defect. Journal of Clinical Immunology, 2019, 39, 611-615.	3.8	3
123	Flow Cytometry for Diagnosis of Primary Immune Deficienciesâ€"A Tertiary Center Experience From North India. Frontiers in Immunology, 2019, 10, 2111.	4.8	18
124	Biomarkers for Kawasaki Disease: Clinical Utility and the Challenges Ahead. Frontiers in Pediatrics, 2019, 7, 242.	1.9	46
125	When Transient Lymphopenia Mimics SCID!. Indian Journal of Pediatrics, 2019, 86, 574-575.	0.8	1
126	Recurrent Salmonella typhi Infection and Autoimmunity in a Young Boy with Complete IL-12 Receptor Î²1 Deficiency. Journal of Clinical Immunology, 2019, 39, 358-362.	3.8	9

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127	Revisiting Fatal Granulomatous Disease of Childhood Through an Autopsy: Still Lethal in the Developing World!. Journal of Clinical Immunology, 2019, 39, 241-244.	3.8	2
128	Genetics of COPA syndrome . The Application of Clinical Genetics, 2019, Volume 12, 11-18.	3.0	17
129	THU0539â€...SERUM SOLUBLE CD25: AN USEFUL BIOMARKER OF MACROPHAGE ACTIVATION SYNDROME IN SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS. , 2019, , .		1
130	THU0510â€...NEUTROPHIL FUNCTION IN PEDIATRIC SYSTEMIC LUPUS ERYTHEMATOSUS. , 2019, , .		0
131	AB1028â€...MYOCARDITIS IN PEDIATRIC LUPUS: A CLINICAL CONUNDRUM. , 2019, , .		0
132	FRI0566â€...STUDY ON SERUM DNAASE1 ACTIVITY IN PEDIATRIC ONSET SYSTEMIC LUPUS ERYTHEMATOSUS FROM A TERTIARY CARE CENTRE IN NORTH WEST INDIA. , 2019, , .		0
133	AB1052â€...SYSTEMIC LUPUS ERYTHEMATOSUS IN CONTEXT OF HUMAN IMMUNODEFICIENCY VIRUS INFECTION: A CLINICAL CONUNDRUM. , 2019, , .		0
134	AB1054â€...HEPATIC INVOLVEMENT AS A PRESENTATION IN PEDIATRIC LUPUS: A RETROSPECTIVE STUDY OF 3 CASES. , 2019, , .		0
135	AB1053â€...MACROPHAGE ACTIVATION SYNDROME AS A PRESENTATION IN PEDIATRIC LUPUS: A RETROSPECTIVE STUDY OF 3 CASES. , 2019, , .		0
136	SAT0504â€...IGG4 RELATED DISEASE IN CHILDREN: A SINGLE CENTRE EXPERIENCE FROM NORTH-WEST INDIA. , 2019, , .		0
137	AB1003â€...DELAY IN DIAGNOSIS OF KAWASAKI DISEASE IS THE COMMONEST PROXIMATE REASON FOR DEVELOPMENT OF GIANT CORONARY ARTERY ANEURYSMS- OUR EXPERIENCE AT CHANDIGARH, NORTH INDIA. , 2019, , .		0
138	AB0518â€...- JUVENILE SYSTEMIC LUPUS ERYTHEMATOSUS RELATED PANCREATITIS: AN UNCOMMON MANIFESTATION OF A COMMON DISEASE. , 2019, , .		11
139	AB0949â€...IS PEDIATRIC ONSET LUPUS MORE SEVERE IN BOYS? OUR EXPERIENCE AT A TERTIARY CARE CENTER IN NORTH-WEST INDIA. , 2019, , .		1
140	AB0589â€...ANCA ASSOCIATED VASCULITIS: OUR EXPERIENCE FROM A TERITIARY CARE CENTER OVER 10 YEARS. , 2019, , .		0
141	Fcâ€gamma receptor expression profile in a Northâ€Indian cohort of pediatricâ€onset systemic lupus erythematosus: An observational study. International Journal of Rheumatic Diseases, 2019, 22, 449-457.	1.9	0
142	Monoclonal Gammopathy of Unclear Significance in a Child with Wiskott-Aldrich Syndrome: a Rare Occurrence. Journal of Clinical Immunology, 2019, 39, 7-10.	3.8	3
143	Reduced Natural Killer Cell Subsets in Perinatally Acquired Long-Term Non-Progressor Human Immunodeficiency Virusâ€"Infected Children. AIDS Research and Human Retroviruses, 2019, 35, 437-443.	1.1	2
144	Complement factor H gene polymorphisms and vivax malaria associated thrombotic microangiopathy. Saudi Journal of Kidney Diseases and Transplantation: an Official Publication of the Saudi Center for Organ Transplantation, Saudi Arabia, 2019, 30, 540.	0.3	1

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145	Purine Nucleoside Phosphorylate Deficiency Severe Combined Immunodeficiency in an Infant: Subtle Diagnostic Clues. Indian Pediatrics, 2019, 56, 146.	0.4	0
146	Functional mannose binding lectin levels in patients with pediatric onset systemic lupus erythematosus in remission. International Journal of Rheumatic Diseases, 2018, 21, 710-715.	1.9	4
147	Inverse Gottron papules in juvenile dermatomyositis: an under recognized clinical entity. Rheumatology International, 2018, 38, 1153-1160.	3.0	7
148	295â€ƒRole of CD40 ligand in the pathogenesis of KD: a single centre study from North India. Rheumatology, 2018, 57, .	1.9	0
149	Mystery Case: Tortuous hairs and tortuous blood vessels. Neurology, 2018, 90, e1174-e1176.	1.1	2
150	Infectious and non-infectious complications in primary immunodeficiency disorders: an autopsy study from North India. Journal of Clinical Pathology, 2018, 71, 425-435.	2.0	8
151	Pulmonary presentation of Kawasaki diseaseâ€"A diagnostic challenge. Pediatric Pulmonology, 2018, 53, 103-107.	2.0	35
152	Correlation between fungal sensitisation in childhood persistent asthma and disease severity. Mycoses, 2018, 61, 195-200.	4.0	13
153	Long-term Seroprotection Rates Following Second Dose of Measles as MMR Vaccine at 15 months in Indian Children. Indian Pediatrics, 2018, 55, 405-407.	0.4	3
154	O27â€ƒProfile of Henoch Schonlein purpura (HSP) nephritis: 23 years of experience at a tertiary care centre in North India. Rheumatology, 2018, 57, .	1.9	0
155	An Infant with Respiratory Distress and Loose Stools. Indian Pediatrics, 2018, 55, 693-697.	0.4	0
156	Autoantibody profile in children with Kawasaki disease on longâ€term followâ€up: A prospective study from North India. International Journal of Rheumatic Diseases, 2018, 21, 2036-2040.	1.9	12
157	Clinical and molecular features of X-linked hyper IgM syndrome â€“ An experience from North India. Clinical Immunology, 2018, 195, 59-66.	3.2	16
158	STAT3-Mediated Transcriptional Regulation of Osteopontin in STAT3 Loss-of-Function Related Hyper IgE Syndrome. Frontiers in Immunology, 2018, 9, 1080.	4.8	16
159	Spectrum of renal disease in HIV-infected children: report of five cases. Paediatrics and International Child Health, 2018, 38, 271-276.	1.0	2
160	Lupus anticoagulant hypoprothrombinemia syndrome associated with systemic lupus erythematosus in children: report of two cases and systematic review of the literature. Rheumatology International, 2018, 38, 1933-1940.	3.0	29
161	O19â€ƒPerforin & granzyme expression: role of natural killer cell cytotoxicity in acute phase of Kawasaki disease. Rheumatology, 2018, 57, .	1.9	0
162	Low Expression of Leucocyte Associated Immunoglobulin Like Receptor-1 (LAIR-1/CD305) in a Cohort of Pediatric Acute Lymphoblastic Leukemia Cases. Asian Pacific Journal of Cancer Prevention, 2018, 19, 3131-3135.	1.2	6

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163	Spectrum of IgG4-related kidney disease at a Tertiary Care Center. Indian Journal of Nephrology, 2018, 28, 209.	0.5	4
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