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List of Publications by Year in descending order

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Version: 2024-02-01

36
papers

1,512
citations

394421

19
h-index

361022

35
g-index

37
all docs

37
docs citations

37
times ranked

2878
citing authors

#	ARTICLE	IF	CITATIONS
1	Revisiting Human IL-12R β 1 Deficiency. <i>Medicine (United States)</i> , 2010, 89, 381-402.	1.0	367
2	B cell subsets in healthy children: Reference values for evaluation of B cell maturation process in peripheral blood. <i>Cytometry Part B - Clinical Cytometry</i> , 2010, 78B, 372-381.	1.5	126
3	Thymus transplantation for complete DiGeorge syndrome: European experience. <i>Journal of Allergy and Clinical Immunology</i> , 2017, 140, 1660-1670.e16.	2.9	108
4	Oxidative stress, mitochondrial abnormalities and antioxidant defense in Ataxia-telangiectasia, Bloom syndrome and Nijmegen breakage syndrome. <i>Redox Biology</i> , 2017, 11, 375-383.	9.0	84
5	Genetic defects in PI3K γ affect B-cell differentiation and maturation leading to hypogammaglobulinemia and recurrent infections. <i>Clinical Immunology</i> , 2017, 176, 77-86.	3.2	80
6	Nijmegen Breakage Syndrome: Clinical and Immunological Features, Long-Term Outcome and Treatment Options – a Retrospective Analysis. <i>Journal of Clinical Immunology</i> , 2015, 35, 538-549.	3.8	73
7	Wiskott–Aldrich Syndrome protein deficiency perturbs the homeostasis of B-cell compartment in humans. <i>Journal of Autoimmunity</i> , 2014, 50, 42-50.	6.5	72
8	Disseminated <i>Bacillus Calmette-Guérin</i> Infection and Immunodeficiency. <i>Emerging Infectious Diseases</i> , 2007, 13, 799-801.	4.3	61
9	Genetic and demographic features of X-linked agammaglobulinemia in Eastern and Central Europe: A cohort study. <i>Molecular Immunology</i> , 2009, 46, 2140-2146.	2.2	50
10	Clinical heterogeneity and diagnostic delay of autoimmune polyendocrinopathy-candidiasis-ectodermal dystrophy syndrome. <i>Clinical Immunology</i> , 2011, 139, 6-11.	3.2	49
11	Efficacy and Safety of Hizentra [®] , a New 20% Immunoglobulin Preparation for Subcutaneous Administration, in Pediatric Patients with Primary Immunodeficiency. <i>Journal of Clinical Immunology</i> , 2011, 31, 752-61.	3.8	47
12	Loss of juxtaposition of RAG-induced immunoglobulin DNA ends is implicated in the precursor B-cell differentiation defect in NBS patients. <i>Blood</i> , 2010, 115, 4770-4777.	1.4	37
13	Genetic characteristics of eighty-seven patients with the Wiskott–Aldrich syndrome. <i>Molecular Immunology</i> , 2011, 48, 788-792.	2.2	35
14	Incomplete penetrance for isolated congenital asplenia in humans with mutations in translated and untranslated <i>RPSA</i> exons. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E8007-E8016.	7.1	31
15	A Novel CDC42 Mutation in an 11-Year Old Child Manifesting as Syndromic Immunodeficiency, Autoinflammation, Hemophagocytic Lymphohistiocytosis, and Malignancy: A Case Report. <i>Frontiers in Immunology</i> , 2020, 11, 318.	4.8	31
16	The defect in humoral immunity in patients with Nijmegen breakage syndrome is explained by defects in peripheral B lymphocyte maturation. <i>Cytometry Part A: the Journal of the International Society for Analytical Cytology</i> , 2012, 81A, 835-842.	1.5	26
17	Nijmegen breakage syndrome: Long-term monitoring of viral and immunological biomarkers in peripheral blood before development of malignancy. <i>Clinical Immunology</i> , 2010, 135, 440-447.	3.2	25
18	Gastrointestinal disorders next to respiratory infections as leading symptoms of X-linked agammaglobulinemia in children – 34-year experience of a single center. <i>Archives of Medical Science</i> , 2017, 2, 412-417.	0.9	23

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19	Antioxidant Defense, Redox Homeostasis, and Oxidative Damage in Children With Ataxia Telangiectasia and Nijmegen Breakage Syndrome. <i>Frontiers in Immunology</i> , 2019, 10, 2322.	4.8	21
20	Common Variable Immune Deficiency in Children—Clinical Characteristics Varies Depending on Defect in Peripheral B Cell Maturation. <i>Journal of Clinical Immunology</i> , 2013, 33, 731-741.	3.8	20
21	EuroFlow Standardized Approach to Diagnostic Immunophenotyping of Severe PID in Newborns and Young Children. <i>Frontiers in Immunology</i> , 2020, 11, 371.	4.8	17
22	Comprehensive activities to increase recognition of primary immunodeficiency and access to immunoglobulin replacement therapy in Poland. <i>European Journal of Pediatrics</i> , 2016, 175, 1099-1105.	2.7	16
23	Comparison of Selected Parameters of Redox Homeostasis in Patients with Ataxia-Telangiectasia and Nijmegen Breakage Syndrome. <i>Oxidative Medicine and Cellular Longevity</i> , 2017, 2017, 1-8.	4.0	16
24	The Clinical and Genetic Spectrum of 82 Patients With RAG Deficiency Including a c.256_257delAA Founder Variant in Slavic Countries. <i>Frontiers in Immunology</i> , 2020, 11, 900.	4.8	16
25	IgG Subclasses and Antibody Response to Pneumococcal Capsular Polysaccharides in Children with Severe Sinopulmonary Infections and Asthma. <i>Immunological Investigations</i> , 1991, 20, 173-185.	2.0	13
26	BCG Moreau Vaccine Safety Profile and NK Cells—Double Protection Against Disseminated BCG Infection in Retrospective Study of BCG Vaccination in 52 Polish Children with Severe Combined Immunodeficiency. <i>Journal of Clinical Immunology</i> , 2020, 40, 138-146.	3.8	13
27	Interstitial Lung Disease in Children With Selected Primary Immunodeficiency Disorders—A Multicenter Observational Study. <i>Frontiers in Immunology</i> , 2020, 11, 1950.	4.8	11
28	Pulmonary Lymphomatoid Granulomatosis in Griscelli Syndrome Type 2. <i>Viral Immunology</i> , 2011, 24, 471-473.	1.3	10
29	Vitamin D deficiency in children with recurrent respiratory infections, with or without immunoglobulin deficiency. <i>Advances in Medical Sciences</i> , 2018, 63, 173-178.	2.1	10
30	COVID-19 Pandemic and Patients with Rare Inherited Metabolic Disorders and Rare Autoinflammatory Diseases—Organizational Challenges from the Point of View of Healthcare Providers. <i>Journal of Clinical Medicine</i> , 2021, 10, 4862.	2.4	9
31	Clinical immunology Disseminated Mycobacterium tuberculosis complex infection in a girl with partial dominant IFN- γ receptor 1 deficiency. <i>Central-European Journal of Immunology</i> , 2012, 4, 378-381.	1.2	4
32	Rapid push: new opportunities in subcutaneous immunoglobulin replacement therapy. <i>Central-European Journal of Immunology</i> , 2013, 3, 388-392.	1.2	3
33	Knowledge Discovery from Medical Data and Development of an Expert System in Immunology. <i>Entropy</i> , 2021, 23, 695.	2.2	3
34	BCG Moreau Polish Substrain Infections in Patients With Inborn Errors of Immunity: 40 Years of Experience in the Department of Immunology, Children's Memorial Health Institute, Warsaw. <i>Frontiers in Pediatrics</i> , 2022, 10, .	1.9	3
35	Clinical and immunological analysis of patients with X-linked agammaglobulinemia — single center experience. <i>Central-European Journal of Immunology</i> , 2013, 3, 367-371.	1.2	1
36	Atypical Hemolytic Uremic Syndrome (aHUS) and Adenosine Deaminase (ADA)-Deficient Severe Combined Immunodeficiency (SCID)—Two Diseases That Exacerbate Each Other: Case Report. <i>International Journal of Molecular Sciences</i> , 2021, 22, 9479.	4.1	1