Ewa Anna Bernatowska

List of Publications by Year in descending order

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55 papers 2,543 citations

304701 22 h-index 50 g-index

59 all docs

59 docs citations

59 times ranked

4020 citing authors

#	Article	IF	CITATIONS
1	Chronic Granulomatous Disease: The European Experience. PLoS ONE, 2009, 4, e5234.	2.5	567
2	Children and adults with primary antibody deficiencies gain quality of life by subcutaneous IgG self-infusions at home. Journal of Allergy and Clinical Immunology, 2004, 114, 936-942.	2.9	237
3	Rationale for pertussis booster vaccination throughout life in Europe. Lancet Infectious Diseases, The, 2011, 11, 557-570.	9.1	222
4	Rapid Subcutaneous IgG Replacement Therapy is Effective and Safe in Children and Adults with Primary Immunodeficiencies—A Prospective, Multi-National Study. Journal of Clinical Immunology, 2006, 26, 177-185.	3.8	176
5	B cell subsets in healthy children: Reference values for evaluation of B cell maturation process in peripheral blood. Cytometry Part B - Clinical Cytometry, 2010, 78B, 372-381.	1.5	126
6	Genetic defects in PI3K $\hat{\Gamma}$ affect B-cell differentiation and maturation leading to hypogammaglobulineamia and recurrent infections. Clinical Immunology, 2017, 176, 77-86.	3.2	80
7	Gross Deletions Involving IGHM, BTK, or Artemis: A Model for Genomic Lesions Mediated by Transposable Elements. American Journal of Human Genetics, 2008, 82, 320-332.	6.2	77
8	Nijmegen Breakage Syndrome: Clinical and Immunological Features, Long-Term Outcome and Treatment Options – a Retrospective Analysis. Journal of Clinical Immunology, 2015, 35, 538-549.	3.8	73
9	Diagnostics of Primary Immunodeficiency Diseases: A Sequencing Capture Approach. PLoS ONE, 2014, 9, e114901.	2.5	73
10	A Global Perspective on Vaccine Safety and Public Health: The Global Advisory Committee on Vaccine Safety. American Journal of Public Health, 2004, 94, 1926-1931.	2.7	72
11	Wiskott–Aldrich Syndrome protein deficiency perturbs the homeostasis of B-cell compartment in humans. Journal of Autoimmunity, 2014, 50, 42-50.	6.5	72
12	The immunophenotypic and immunogenotypic B-cell differentiation arrest in bone marrow of RAG-deficient SCID patients corresponds to residual recombination activities of mutated RAG proteins. Blood, 2002, 100, 2145-52.	1.4	64
13	<i>Cryptosporidium</i> Infection in Patients With Primary Immunodeficiencies. Journal of Pediatric Gastroenterology and Nutrition, 2007, 45, 458-464.	1.8	62
14	Disseminated Bacillus Calmette-Guérin Infection and Immunodeficiency. Emerging Infectious Diseases, 2007, 13, 799-801.	4.3	61
15	Results of a prospective controlled two-dose crossover study with intravenous immunoglobulin and comparison (retrospective) with plasma treatment. Clinical Immunology and Immunopathology, 1987, 43, 153-162.	2.0	54
16	Genetic and demographic features of X-linked agammaglobulinemia in Eastern and Central Europe: A cohort study. Molecular Immunology, 2009, 46, 2140-2146.	2.2	50
17	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. Frontiers in Immunology, 2017, 8, 685.	4.8	50
18	Loss of juxtaposition of RAC-induced immunoglobulin DNA ends is implicated in the precursor B-cell differentiation defect in NBS patients. Blood, 2010, 115, 4770-4777.	1.4	37

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19	Genetic characteristics of eighty-seven patients with the Wiskott–Aldrich syndrome. Molecular Immunology, 2011, 48, 788-792.	2.2	35
20	A Novel CDC42 Mutation in an 11-Year Old Child Manifesting as Syndromic Immunodeficiency, Autoinflammation, Hemophagocytic Lymphohistiocytosis, and Malignancy: A Case Report. Frontiers in Immunology, 2020, 11, 318.	4.8	31
21	Ataxia-Telangiectasia With Hyper-IgM and Wilms Tumor: Fatal Reaction to Irradiation. Journal of Pediatric Hematology/Oncology, 2010, 32, e28-e30.	0.6	29
22	The defect in humoral immunity in patients with Nijmegen breakage syndrome is explained by defects in peripheral B lymphocyte maturation. Cytometry Part A: the Journal of the International Society for Analytical Cytology, 2012, 81A, 835-842.	1.5	26
23	Antioxidant Defense, Redox Homeostasis, and Oxidative Damage in Children With Ataxia Telangiectasia and Nijmegen Breakage Syndrome. Frontiers in Immunology, 2019, 10, 2322.	4.8	21
24	Common Variable Immune Deficiency in Children—Clinical Characteristics Varies Depending on Defect in Peripheral B Cell Maturation. Journal of Clinical Immunology, 2013, 33, 731-741.	3.8	20
25	Meningococcal B Vaccine Immunogenicity in Children With Defects in Complement and Splenic Function. Pediatrics, 2018, 142, .	2.1	17
26	Comprehensive activities to increase recognition of primary immunodeficiency and access to immunoglobulin replacement therapy in Poland. European Journal of Pediatrics, 2016, 175, 1099-1105.	2.7	16
27	Comparison of Selected Parameters of Redox Homeostasis in Patients with Ataxia-Telangiectasia and Nijmegen Breakage Syndrome. Oxidative Medicine and Cellular Longevity, 2017, 2017, 1-8.	4.0	16
28	The Clinical and Genetic Spectrum of 82 Patients With RAG Deficiency Including a c.256_257delAA Founder Variant in Slavic Countries. Frontiers in Immunology, 2020, 11, 900.	4.8	16
29	Successful treatment of refractory autoimmune thrombocytopenia with rituximab and cyclosporin A in a patient with chronic granulomatous disease. Annals of Hematology, 2005, 84, 835-836.	1.8	15
30	Chronic rhinosinusitis in primary antibody immunodeficient patients. International Journal of Pediatric Otorhinolaryngology, 2006, 70, 1587-1592.	1.0	14
31	Vaccination-related <i>Mycobacterium bovis </i> BCG Infection. Emerging Infectious Diseases, 2006, 12, 860-861.	4.3	14
32	Treosulfan-based conditioning regimen in a second matched unrelated peripheral blood stem cell transplantation for a pediatric patient with CGD and invasive aspergillosis, who experienced initial graft failure after RIC. International Journal of Hematology, 2009, 90, 571-575.	1.6	14
33	BCG Moreau Vaccine Safety Profile and NK Cellsâ€"Double Protection Against Disseminated BCG Infection in Retrospective Study of BCG Vaccination in 52 Polish Children with Severe Combined Immunodeficiency. Journal of Clinical Immunology, 2020, 40, 138-146.	3.8	13
34	Genotyping of Cryptosporidium isolates from human clinical cases in Poland. Parasitology Research, 2008, 103, 37-42.	1.6	11
35	Interstitial Lung Disease in Children With Selected Primary Immunodeficiency Disorders—A Multicenter Observational Study. Frontiers in Immunology, 2020, 11, 1950.	4.8	11
36	Vitamin D deficiency in children with recurrent respiratory infections, with or without immunoglobulin deficiency. Advances in Medical Sciences, 2018, 63, 173-178.	2.1	10

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37	A Multicentre Study on the Efficacy, Safety and Pharmacokinetics of IqYmune®, a Highly Purified 10% Liquid Intravenous Immunoglobulin, in Patients with Primary Immune Deficiency. Journal of Clinical Immunology, 2017, 37, 539-547.	3.8	9
38	The use of Staphylococcus aureus Cowan I for evaluation of suppressor-T-cell activity in hypogammaglobulinemia: Evidence for two functionally distinct suppressor T cells. Clinical Immunology and Immunopathology, 1984, 33, 293-300.	2.0	8
39	Zalecenia Polskiej Grupy Roboczej ds. Inwazyjnej Choroby Pneumokowej (IChP) u Dzieci dotyczÄce stosowania siedmiowalentnej skoniugowanej szczepionki przeciwpneumokokowej (PCV7). Pediatria Polska, 2007, 82, 486-491.	0.2	5
40	Neutrophil Phenotypic Characteristics in Children with Congenital Asplenia and Splenectomized for Hereditary Spherocytosis. Immunological Investigations, 2012, 41, 61-74.	2.0	4
41	Peripheral blood T lymphocyte subsets in children with congenital asplenia. Human Immunology, 2012, 73, 1091-1097.	2.4	4
42	Clinical immunology Disseminated Mycobacterium tuberculosis complex infection in a girl with partial dominant IFN- \hat{l}^3 receptor 1 deficiency. Central-European Journal of Immunology, 2012, 4, 378-381.	1.2	4
43	Fabricated or induced illness in the oral cavity in children. A systematic review and personal experience. Central-European Journal of Immunology, 2015, 1, 109-114.	1.2	4
44	A Rose Amongst the Thorns: the Mission of the J Project in a Conflictual World. Journal of Clinical Immunology, 2022, 42, 1151-1155.	3.8	4
45	A novel radiosensitive SCID patient with a pronounced G2/M sensitivity. DNA Repair, 2010, 9, 365-373.	2.8	3
46	Rapid push: new opportunities in subcutaneous immunoglobulin replacement therapy. Central-European Journal of Immunology, 2013, 3, 388-392.	1.2	3
47	Tumor Necrosis Factor Receptor-Associated Periodic Syndrome (TRAPS) with a New Pathogenic Variant in TNFRSF1A Gene in a Family of the Adult Male with Renal AA Amyloidosis—Diagnostic and Therapeutic Challenge for Clinicians. Journal of Clinical Medicine, 2021, 10, 465.	2.4	3
48	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. Frontiers in Pediatrics, 2022, 10, .	1.9	3
49	A Multiâ€'Center, Openâ€'Label, Singleâ€'Arm Trial to Evaluate the Efficacy,ÂPharmacokinetics, and Safety and Tolerability of IGSC 20% in SubjectsAwith Primary Immunodeficiency. Journal of Clinical Immunology, 2022, 42, 500-511.	3.8	2
50	Pierwotne niedobory odpornoÅ;ci. Pediatria Polska, 2008, 83, 697-703.	0.2	1
51	Clinical and immunological analysis of patients with X-linked agammaglobulinemia – single center experience. Central-European Journal of Immunology, 2013, 3, 367-371.	1.2	1
52	Inwazyjne zakaÅ⅓enie Haemophilus influenzae typu b u dziecka z peÅ,nym cyklem szczepieÅ". Pediatria Polska, 2013, 88, 103-106.	0.2	0
53	Clinical immunology Clinical manifestations in the oral cavity in patients with hyper-IgE syndrome Central-European Journal of Immunology, 2013, 1, 92-99.	1.2	0
54	Witamina D a odporność u dzieci. Pediatria Polska, 2016, 91, 251-256.	0.2	0

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55	Program Szczepień Ochronnych w 2017 roku ┠powszechne szczepienia przeciwko pneumokokom u dzieci. Pediatria Polska, 2017, 92, 231-237.	0.2	0