Ewa Anna Bernatowska

List of Publications by Year
in descending order

55 papers	2,543 citations	346980 22 h-index	214428 50 g-index
59 all docs	59 docs citations	59 times ranked	4239 citing authors

#	Article	IF	Citations
1	A Multiâ€'Center, Openâ€'Label, Singleâ€'Arm Trial to Evaluate the Efficacy,ÂPharmacokinetics, and Safety and Tolerability of IGSC 20% in SubjectsÂwith Primary Immunodeficiency. Journal of Clinical Immunology, 2022, 42, 500-511.	2.0	2
2	A Rose Amongst the Thorns: the Mission of the J Project in a Conflictual World. Journal of Clinical Immunology, 2022, 42, 1151-1155.	2.0	4
3	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, .	0.9	3
4	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 Amyloidosisâ€"Diagnostic and Therapeutic Challenge for Clinicians. Journal of Clinical Medicine, 2021, 10, 465.	1.0	3
5	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.	2.0	13
6	Interstitial Lung Disease in Children With Selected Primary Immunodeficiency Disordersâ€"A Multicenter Observational Study. Frontiers in Immunology, 2020, 11, 1950.	2.2	11
7	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.	2.2	16
8	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.	2.2	31
9	Antioxidant Defense, Redox Homeostasis, and Oxidative Damage in Children With Ataxia Telangiectasia and Nijmegen Breakage Syndrome. Frontiers in Immunology, 2019, 10, 2322.	2.2	21
10	Vitamin D deficiency in children with recurrent respiratory infections, with or without immunoglobulin deficiency. Advances in Medical Sciences, 2018, 63, 173-178.	0.9	10
11	Meningococcal B Vaccine Immunogenicity in Children With Defects in Complement and Splenic Function. Pediatrics, 2018, 142, .	1.0	17
12	Genetic defects in PI3KÎ´ affect B-cell differentiation and maturation leading to hypogammaglobulineamia and recurrent infections. Clinical Immunology, 2017, 176, 77-86.	1.4	80
13	Program SzczepieÅ„ Ochronnych w 2017 roku â^' powszechne szczepienia przeciwko pneumokokom u dzieci. Pediatria Polska, 2017, 92, 231-237.	0.1	0
14	A Multicentre Study on the Efficacy, Safety and Pharmacokinetics of IqYmuneÂ®, a Highly Purified 10% Liquid Intravenous Immunoglobulin, in Patients with Primary Immune Deficiency. Journal of Clinical Immunology, 2017, 37, 539-547.	2.0	9
15	Patients with Primary Immunodeficiencies Are a Reservoir of Poliovirus and a Risk to Polio Eradication. Frontiers in Immunology, 2017, 8, 685.	2.2	50
16	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.	1.9	16
17	Comprehensive activities to increase recognition of primary immunodeficiency and access to immunoglobulin replacement therapy in Poland. European Journal of Pediatrics, 2016, 175, 1099-1105.	1.3	16
18	Witamina D a odpornoÅ›Ä‡ u dzieci. Pediatria Polska, 2016, 91, 251-256.	0.1	0

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19	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.	0.4	4
20	Nijmegen Breakage Syndrome: Clinical and Immunological Features, Long-Term Outcome and Treatment Options â€“ a Retrospective Analysis. Journal of Clinical Immunology, 2015, 35, 538-549.	2.0	73
21	Wiskottâ€“Aldrich Syndrome protein deficiency perturbs the homeostasis of B-cell compartment in humans. Journal of Autoimmunity, 2014, 50, 42-50.	3.0	72
22	Diagnostics of Primary Immunodeficiency Diseases: A Sequencing Capture Approach. PLoS ONE, 2014, 9, e114901.	1.1	73
23	InwazyjneÂ zakaÅ¼enie Haemophilus influenzae typu b u dziecka z peÅ‚nym cyklem szczepieÅ„. Pediatria Polska, 2013, 88, 103-106.	0.1	0
24	Common Variable Immune Deficiency in Childrenâ€”Clinical Characteristics Varies Depending on Defect in Peripheral B Cell Maturation. Journal of Clinical Immunology, 2013, 33, 731-741.	2.0	20
25	Rapid push: new opportunities in subcutaneous immunoglobulin replacement therapy. Central-European Journal of Immunology, 2013, 3, 388-392.	0.4	3
26	Clinical immunology Clinical manifestations in the oral cavity in patients with hyper-IgE syndrome.. Central-European Journal of Immunology, 2013, 1, 92-99.	0.4	0
27	Clinical and immunological analysis of patients with X-linked agammaglobulinemia â€“ single center experience. Central-European Journal of Immunology, 2013, 3, 367-371.	0.4	1
28	Neutrophil Phenotypic Characteristics in Children with Congenital Asplenia and Splenectomized for Hereditary Spherocytosis. Immunological Investigations, 2012, 41, 61-74.	1.0	4
29	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.	1.1	26
30	Peripheral blood T lymphocyte subsets in children with congenital asplenia. Human Immunology, 2012, 73, 1091-1097.	1.2	4
31	Clinical immunology Disseminated Mycobacterium tuberculosis complex infection in a girl with partial dominant IFN-Î³ receptor 1 deficiency. Central-European Journal of Immunology, 2012, 4, 378-381.	0.4	4
32	Rationale for pertussis booster vaccination throughout life in Europe. Lancet Infectious Diseases, The, 2011, 11, 557-570.	4.6	222
33	Genetic characteristics of eighty-seven patients with the Wiskottâ€“Aldrich syndrome. Molecular Immunology, 2011, 48, 788-792.	1.0	35
34	Loss of juxtaposition of RAG-induced immunoglobulin DNA ends is implicated in the precursor B-cell differentiation defect in NBS patients. Blood, 2010, 115, 4770-4777.	0.6	37
35	Ataxia-Telangiectasia With Hyper-IgM and Wilms Tumor: Fatal Reaction to Irradiation. Journal of Pediatric Hematology/Oncology, 2010, 32, e28-e30.	0.3	29
36	A novel radiosensitive SCID patient with a pronounced G2/M sensitivity. DNA Repair, 2010, 9, 365-373.	1.3	3

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37	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.	0.7	126
38	Chronic Granulomatous Disease: The European Experience. PLoS ONE, 2009, 4, e5234.	1.1	567
39	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.	0.7	14
40	Genetic and demographic features of X-linked agammaglobulinemia in Eastern and Central Europe: A cohort study. Molecular Immunology, 2009, 46, 2140-2146.	1.0	50
41	Genotyping of Cryptosporidium isolates from human clinical cases in Poland. Parasitology Research, 2008, 103, 37-42.	0.6	11
42	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.	2.6	77
43	Pierwotne niedobory odpornoÅ›ci. Pediatria Polska, 2008, 83, 697-703.	0.1	1
44	Cryptosporidium Infection in Patients With Primary Immunodeficiencies. Journal of Pediatric Gastroenterology and Nutrition, 2007, 45, 458-464.	0.9	62
45	Zalecenia Polskiej Grupy Roboczej ds. Inwazyjnej Choroby Pneumokowej (IChP) u Dzieci dotyczÄ…ce stosowania siedmiowalentnej skoniugowanej szczepionki przeciwpneumokokowej (PCV7). Pediatria Polska, 2007, 82, 486-491.	0.1	5
46	Disseminated Bacillus Calmette-GuÃ©rin Infection and Immunodeficiency. Emerging Infectious Diseases, 2007, 13, 799-801.	2.0	61
47	Chronic rhinosinusitis in primary antibody immunodeficient patients. International Journal of Pediatric Otorhinolaryngology, 2006, 70, 1587-1592.	0.4	14
48	Vaccination-relatedMycobacterium bovisBCG Infection. Emerging Infectious Diseases, 2006, 12, 860-861.	2.0	14
49	Rapid Subcutaneous IgG Replacement Therapy is Effective and Safe in Children and Adults with Primary Immunodeficienciesâ€"A Prospective, Multi-National Study. Journal of Clinical Immunology, 2006, 26, 177-185.	2.0	176
50	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.	0.8	15
51	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.	1.5	237
52	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.	1.5	72
53	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.	0.6	64
54	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.	2.1	54

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55	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.	2.1	8