Wojciech MÅ‚ynarski

List of Publications by Year
in descending order

166	2,738	279701	233338
papers	citations	23 h-index	45 g-index

168	168	168	5093
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Therapy of type 1 diabetes with CD4+CD25highCD127-regulatory T cells prolongs survival of pancreatic islets â€" Results of one year follow-up. Clinical Immunology, 2014, 153, 23-30.	1.4	307
2	ISPAD Clinical Practice Consensus Guidelines 2018: The diagnosis and management of monogenic diabetes in children and adolescents. Pediatric Diabetes, 2018, 19, 47-63.	1.2	227
3	SARS-CoV-2 Mpro inhibitors and activity-based probes for patient-sample imaging. Nature Chemical Biology, 2021, 17, 222-228.	3.9	215
4	GATA4 Mutations Are a Cause of Neonatal and Childhood-Onset Diabetes. Diabetes, 2014, 63, 2888-2894.	0.3	108
5	Sulfonylurea improves CNS function in a case of intermediate DEND syndrome caused by a mutation in KCNJ11. Nature Clinical Practice Neurology, 2007, 3, 640-645.	2.7	102
6	Factors affecting long-term efficacy of T regulatory cell-based therapy in type 1 diabetes. Journal of Translational Medicine, 2016, 14, 332.	1.8	83
7	Defective DNA damage repair leads to frequent catastrophic genomic events in murine and human tumors. Nature Communications, 2018, 9, 4760.	5.8	66
8	High Yield of Pathogenic Germline Mutations Causative or Likely Causative of the Cancer Phenotype in Selected Children with Cancer. Clinical Cancer Research, 2018, 24, 1594-1603.	3.2	52
9	Complications of mammalian target of rapamycin inhibitor anticancer treatment among patients with tuberous sclerosis complex are common and occasionally life-threatening. Anti-Cancer Drugs, 2015, 26, 437-442.	0.7	49
10	Validation of the United Kingdom copy-number alteration classifier in 3239 children with B-cell precursor ALL. Blood Advances, 2019, 3, 148-157.	2.5	48
11	Psychiatric Disorders and Health-Related Quality of Life in Children With Type 1 Diabetes Mellitus. Psychosomatics, 2016, 57, 185-193.	2.5	47
12	Monogenic diabetes syndromes: Locus-specific databases for AlstrÃ¶m, Wolfram, and Thiamine-responsive megaloblastic anemia. Human Mutation, 2017, 38, 764-777.	1.1	47
13	EURO-WABB: an EU rare diseases registry for Wolfram syndrome, AlstrÃ¶m syndrome and Bardet-Biedl syndrome. BMC Pediatrics, 2013, 13, 130.	0.7	43
14	Biallelic loss ofCDKN2Ais associated with poor response to treatment in pediatric acute lymphoblastic leukemia. Leukemia and Lymphoma, 2017, 58, 1162-1171.	0.6	43
15	Natural History and Current Treatment Options for Subependymal Giant Cell Astrocytoma in Tuberous Sclerosis Complex. Seminars in Pediatric Neurology, 2015, 22, 274-281.	1.0	41
16	Clinical course and therapeutic implications for lymphoid malignancies in Nijmegen breakage syndrome. European Journal of Medical Genetics, 2016, 59, 126-132.	0.7	38
17	Dental Anomalies as Late Adverse Effect among Young Children Treated for Cancer. Cancer Research and Treatment, 2016, 48, 658-667.	1.3	33
18	Flash Glucose Measurements in Children with Type 1 Diabetes in Real-Life Settings: To Trust or Not to Trust?. Diabetes Technology and Therapeutics, 2018, 20, 17-24.	2.4	32

#	Article	IF	Citations
19	Novel severe hemophilia A and moyamoya (SHAM) syndrome caused by Xq28 deletions encompassing F8 and BRCC3 genes. Blood, 2014, 123, 4002-4004.	0.6	31
20	Palliative Sedation at Home for Terminally Ill Children With Cancer. Journal of Pain and Symptom Management, 2014, 48, 968-974.	0.6	27
21	Intragenic amplification of PAX5: a novel subgroup in B-cell precursor acute lymphoblastic leukemia?. Blood Advances, 2017, 1, 1473-1477.	2.5	25
22	The Role of Histone Protein Modifications and Mutations in Histone Modifiers in Pediatric B-Cell Progenitor Acute Lymphoblastic Leukemia. Cancers, 2017, 9, 2.	1.7	25
23	Targeting the thioredoxin system as a novel strategy against Bâ€cell acute lymphoblastic leukemia. Molecular Oncology, 2019, 13, 1180-1195.	2.1	24
24	Altered Plateletsâ€™ morphological parameters in children with type 1 diabetes â€“ a case-control study. BMC Endocrine Disorders, 2015, 15, 17.	0.9	23
25	Interleukin 18 as a Marker of Chronic Nephropathy in Children after Anticancer Treatment. Disease Markers, 2013, 35, 811-818.	0.6	22
26	GlyCulator2: an update on a web application for calculation of glycemic variability indices. Acta Diabetologica, 2018, 55, 877-880.	1.2	22
27	Retinal Thinning as a Marker of Disease Progression in Patients With Wolfram Syndrome. Diabetes Care, 2015, 38, e36-e37.	4.3	21
28	Markers influencing the presence of partial clinical remission in patients with newly diagnosed type 1 diabetes. Journal of Pediatric Endocrinology and Metabolism, 2017, 30, 1147-1153.	0.4	21
29	Monogenic diabetes prevalence among Polish children-Summary of 11â€‰years-long nationwide genetic screening program. Pediatric Diabetes, 2018, 19, 53-58.	1.2	21
30	Asparagine synthetase (ASNS) gene polymorphism is associated with the outcome of childhood acute lymphoblastic leukemia by affecting early response to treatment. Leukemia Research, 2014, 38, 180-183.	0.4	20
31	Abnormal serum microRNA profiles in tuberous sclerosis are normalized during treatment with everolimus: possible clinical implications. Orphanet Journal of Rare Diseases, 2016, 11, 129.	1.2	20
32	A family with the Arg103Pro mutation in the NEUROD1 gene detected by next-generation sequencing â€“ Clinical characteristics of mutation carriers. European Journal of Medical Genetics, 2016, 59, 75-79.	0.7	19
33	Retinal thickness as a marker of disease progression in longitudinal observation of patients with Wolfram syndrome. Acta Diabetologica, 2017, 54, 1019-1024.	1.2	19
34	COVID-19 in pediatric cancer patients is associated with treatment interruptions but not with short-term mortality: a Polish national study. Journal of Hematology and Oncology, 2021, 14, 163.	6.9	19
35	Circulating ghrelin level is higher in HNF1Aâ€“MODY and GCKâ€“MODY than in polygenic forms of diabetes mellitus. Endocrine, 2015, 50, 643-649.	1.1	18
36	Differential regulation of serum microRNA expression by HNF1Î² and HNF1Î± transcription factors. Diabetologia, 2016, 59, 1463-1473.	2.9	18

#	Article	IF	Citations
37	Optical coherence tomography and magnetic resonance imaging visual pathway evaluation in Wolfram syndrome. Developmental Medicine and Child Neurology, 2019, 61, 359-365.	1.1	18
38	Neutrophil Elastase Defects in Congenital Neutropenia. Frontiers in Immunology, 2021, 12, 653932.	2.2	18
39	Decreased FOXP3 mRNA expression in children with atopic asthma and IgE-mediated food allergy. Annals of Allergy, Asthma and Immunology, 2015, 115, 415-421.	0.5	17
40	Maintenance therapy with everolimus for subependymal giant cell astrocytoma in patients with tuberous sclerosis (the EMINENTS study). Pediatric Blood and Cancer, 2017, 64, e26347.	0.8	17
41	Measurement of corneal thickness, optic nerve sheath diameter and retinal nerve fiber layer as potential new non-invasive methods in assessing a risk of cerebral edema in type 1 diabetes in children. Acta Diabetologica, 2018, 55, 1295-1301.	1.2	17
42	Temporal dynamics of serum letâ€7g expression mirror the decline of residual betaâ€cell function in longitudinal observation of children with type 1 diabetes. Pediatric Diabetes, 2018, 19, 1407-1415.	1.2	16
43	Combined therapy with <scp>CD4</scp>⁺<scp>CD25highCD127</scp>^{â^'} T regulatory cells and <scp>antiâ€CD20</scp> antibody in recentâ€onset type 1 diabetes is superior to monotherapy: Randomized phase I/<scp>II</scp> trial. Diabetes, Obesity and Metabolism, 2022, 24, 1534-1543.	2.2	15
44	Development of treatment and clinical results in childhood acute myeloid leukemia in Poland. Memo - Magazine of European Medical Oncology, 2013, 6, 54-62.	0.3	14
45	Effect of Insulin Dilution on Lowering Glycemic Variability in Pump-Treated Young Children with Inadequately Controlled Type 1 Diabetes. Diabetes Technology and Therapeutics, 2015, 17, 605-610.	2.4	14
46	Nivolumab for the Treatment of Advanced Pediatric Malignancies. Anticancer Research, 2020, 40, 7095-7100.	0.5	14
47	Proinsulin-specific T regulatory cells may control immune responses in type 1 diabetes: implications for adoptive therapy. BMJ Open Diabetes Research and Care, 2020, 8, e000873.	1.2	14
48	Severe toxicity free survival: physician-derived definitions of unacceptable long-term toxicities following acute lymphocytic leukaemia. Lancet Haematology,the, 2021, 8, e513-e523.	2.2	14
49	Polymorphism inIKZF1gene affects age at onset of childhood acute lymphoblastic leukemia. Leukemia and Lymphoma, 2014, 55, 2174-2178.	0.6	13
50	Continuous Glucose Monitoring in Type 1 Diabetes Pregnancy Shows that Fetal Heart Rate Correlates with Maternal Glycemia. Diabetes Technology and Therapeutics, 2015, 17, 619-624.	2.4	13
51	Surface expression of Cytokine Receptor-Like Factor 2 increases risk of relapse in pediatric acute lymphoblastic leukemia patients harboring IKZF1 deletions. Oncotarget, 2018, 9, 25971-25982.	0.8	13
52	Hematopoietic Stem Cell Transplantation Positively Affects the Natural History of Cancer in Nijmegen Breakage Syndrome. Clinical Cancer Research, 2021, 27, 575-584.	3.2	13
53	The Impact of Pediatric Palliative Care Education on Medical Studentsâ€™ Knowledge and Attitudes. Scientific World Journal, The, 2013, 2013, 1-9.	0.8	12
54	Intima-media thickness and endothelial dysfunction in GCK and HNF1A-MODY patients. European Journal of Endocrinology, 2015, 172, 277-283.	1.9	12

#	Article	IF	Citations
55	HLA-G and MHC Class II Protein Expression in Diffuse Large B-Cell Lymphoma. Archivum Immunologiae Et Therapiae Experimentalis, 2016, 64, 225-240.	1.0	12
56	Unfavorable Outcome of Neuroblastoma in Patients With 2p Gain. Frontiers in Oncology, 2019, 9, 1018.	1.3	12
57	A crossâ€sectional study of patients referred for <scp> HNF1B </scp> â€MODY genetic testing due to cystic kidneys and diabetes. Pediatric Diabetes, 2020, 21, 422-430.	1.2	12
58	Genetic Signature of Acute Lymphoblastic Leukemia and Netherton Syndrome Co-incidenceâ€"First Report in the Literature. Frontiers in Oncology, 2019, 9, 1477.	1.3	12
59	Maintenance Therapy With Everolimus for Subependymal Giant Cell Astrocytoma in Patients With Tuberous Sclerosis â€“ Final Results From the EMINENTS Study. Frontiers in Neurology, 2021, 12, 581102.	1.1	12
60	Breaking the Taboo: Illicit Drug Use among Adolescents with Type 1 Diabetes Mellitus. Journal of Diabetes Research, 2016, 2016, 1-8.	1.0	11
61	Haemophilia A and cardiovascular morbidity in a female SHAM syndrome carrier due to skewed X chromosome inactivation. European Journal of Medical Genetics, 2016, 59, 43-47.	0.7	11
62	Essential oils reduce autonomous response to pain sensation during self-monitoring of blood glucose among children with diabetes. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 47-53.	0.4	11
63	Serum Metabolic Fingerprinting Identified Putatively Annotated Sphinganine Isomer as a Biomarker of Wolfram Syndrome. Journal of Proteome Research, 2017, 16, 4000-4008.	1.8	11
64	NIRCa: An artificial neural network-based insulin resistance calculator. Pediatric Diabetes, 2018, 19, 231-235.	1.2	11
65	Polymorphism of theFTOGene Influences Body Weight in Children with Type 1 Diabetes without Severe Obesity. International Journal of Endocrinology, 2014, 2014, 1-5.	0.6	10
66	The Stricter the Better? The Relationship between Targeted HbA_{1c}Values and Metabolic Control of Pediatric Type 1 Diabetes Mellitus. Journal of Diabetes Research, 2016, 2016, 1-7.	1.0	10
67	Genetic Association Study of IL2RA, IFIH1, and CTLA-4 Polymorphisms With Autoimmune Thyroid Diseases and Type 1 Diabetes. Frontiers in Pediatrics, 2020, 8, 481.	0.9	10
68	Next-Generation Sequencing in the Diagnosis of Patients with Bardetâ€"Biedl Syndromeâ€"New Variants and Relationship with Hyperglycemia and Insulin Resistance. Genes, 2020, 11, 1283.	1.0	10
69	Advances in the First Line Treatment of Pediatric Acute Myeloid Leukemia in the Polish Pediatric Leukemia and Lymphoma Study Group from 1983 to 2019. Cancers, 2021, 13, 4536.	1.7	10
70	Surface Expression of CRLF2 Protein Is Associated with Lower Minimal Residual Disease (MRD) Among Children with IKZF1-deleted Acute Lymphoblastic Leukemia (ALL). Blood, 2014, 124, 2400-2400.	0.6	10
71	Central Nervous System PET-CT Imaging Reveals Regional Impairments in Pediatric Patients with Wolfram Syndrome. PLoS ONE, 2014, 9, e115605.	1.1	9
72	Dental caries among childhood cancer survivors. Medicine (United States), 2019, 98, e14279.	0.4	9

#	Article	IF	Citations
73	GATA3 germline variant is associated with CRLF2 expression and predicts outcome in pediatric Bâ€cell precursor acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2019, 58, 619-626.	1.5	9
74	Results of two consecutive treatment protocols in Polish children with acute lymphoblastic leukemia. Scientific Reports, 2020, 10, 20168.	1.6	9
75	Genetic predisposition to lymphomas: Overview of rare syndromes and inherited familial variants. Mutation Research - Reviews in Mutation Research, 2021, 788, 108386.	2.4	9
76	Potent, p53-independent induction of NOXA sensitizes MLL-rearranged B-cell acute lymphoblastic leukemia cells to venetoclax. Oncogene, 2022, 41, 1600-1609.	2.6	9
77	Longâ€term treatment results of Polish pediatric and adolescent patients enrolled in the ALL ICâ€BFM 2002 trial. American Journal of Hematology, 2019, 94, E307-E310.	2.0	8
78	The HD-OCT Study May Be Useful in Searching for Markers of Preclinical Stage of Diabetic Retinopathy in Patients with Type 1 Diabetes. Diagnostics, 2019, 9, 105.	1.3	8
79	Six molecular patterns leading to hemophilia A phenotype in 18 females from Poland. Thrombosis Research, 2020, 193, 9-14.	0.8	8
80	Ovarian carcinoma in children with constitutional mutation of SMARCA4: single-family report and literature review. Familial Cancer, 2021, 20, 355-362.	0.9	8
81	The first investigation of Wilms' tumour atomic structure-nitrogen and carbon isotopic composition as a novel biomarker for the most individual approach in cancer disease. Oncotarget, 2016, 7, 76726-76734.	0.8	8
82	Metabolic bone markers can be related to preserved insulin secretion in children with newly diagnosed type 1 diabetes. Pediatric Endocrinology, Diabetes and Metabolism, 2020, 26, 10-16.	0.3	8
83	Ocena czÄ™stoÅ›ci wystÄ™powania zespoÅ‚u Wolframa w populacji dzieci z cukrzycÄ…. Endokrynologia Polska, 2014, 65, 295-297.	0.3	8
84	Prognostic significance of IKZF1 deletions and IKZF1^{plus} profile in children with Bâ€cell precursor acute lymphoblastic leukemia treated according to the ALLâ€IC BFM 2009 protocol. Hematological Oncology, 2022, 40, 430-441.	0.8	8
85	Chromosome 18q deletion syndrome with autoimmune diabetes mellitus: putative genomic loci for autoimmunity and immunodeficiency. Pediatric Diabetes, 2016, 17, 153-159.	1.2	7
86	Microarray testing as an efficient tool to redefine hyperdiploid paediatric B-cell precursor acute lymphoblastic leukaemia patients. Leukemia Research, 2019, 83, 106163.	0.4	7
87	Corneal Abnormalities Are Novel Clinical Feature in Wolfram Syndrome. American Journal of Ophthalmology, 2020, 217, 140-151.	1.7	7
88	Accessibility of the reference center as a protective factor against ketoacidosis at the onset of diabetes in children. Journal of Pediatric Endocrinology and Metabolism, 2014, 27, 1137-43.	0.4	6
89	Polymorphism in IKZF1 gene affects clinical outcome in diffuse large B-cell lymphoma. International Journal of Hematology, 2017, 106, 794-800.	0.7	6
90	Resistance to Data Loss of Glycemic Variability Measurements in Long-Term Continuous Glucose Monitoring. Diabetes Technology and Therapeutics, 2018, 20, 833-842.	2.4	6

#	Article	IF	Citations
91	Clinical Outcome in Pediatric Patients with Philadelphia Chromosome Positive ALL Treated with Tyrosine Kinase Inhibitors Plus Chemotherapyâ€"The Experience of a Polish Pediatric Leukemia and Lymphoma Study Group. Cancers, 2020, 12, 3751.	1.7	6
92	Above 40% of Polish children and young adults with type 1 diabetes achieve international <scp>HbA1c</scp> target â€•results of a nationwide crossâ€sectional evaluation of glycemic control: The <scp>PolPeDiab HbA1c</scp> study. Pediatric Diabetes, 2021, 22, 1003-1013.	1.2	6
93	Immune Dysregulation in Patients With Chromosome 18q Deletionsâ€"Searching for Putative Loci for Autoimmunity and Immunodeficiency. Frontiers in Immunology, 2021, 12, 742834.	2.2	6
94	Multiomic analysis on human cell model of wolfram syndrome reveals changes in mitochondrial morphology and function. Cell Communication and Signaling, 2021, 19, 116.	2.7	6
95	One-step nucleic acid amplification testing in medullary thyroid cancer lymph nodes: a case series. Archives of Medical Science, 2015, 1, 137-141.	0.4	5
96	Heterozygous carriers of germline c.657_661del5 founder mutation in NBN gene are at risk of central nervous system relapse of B-cell precursor acute lymphoblastic leukemia. Haematologica, 2018, 103, e200-e203.	1.7	5
97	Multiple Retinal Anomalies in Wfs1-Deficient Mice. Diagnostics, 2020, 10, 607.	1.3	5
98	Retrospective Analysis of the Treatment Outcome in Myeloid Leukemia of Down Syndrome in Polish Pediatric Leukemia and Lymphoma Study Group From 2005 to 2019. Frontiers in Pediatrics, 2020, 8, 277.	0.9	5
99	Can we effectively predict the occurrence of cerebral edema in children with ketoacidosis in the course of type 1 diabetes? â€“ case report and literature review. Journal of Pediatric Endocrinology and Metabolism, 2020, 33, 319-322.	0.4	5
100	Determination and interpretation of MTHFR gene mutations in gynecology and internal medicine. Polish Archives of Internal Medicine, 2019, 129, 728-732.	0.3	5
101	Consensus Recommendations for the Clinical Management of Hematological Malignancies in Patients with DNA Double Stranded Break Disorders. Cancers, 2022, 14, 2000.	1.7	5
102	Tackling the Turmoil of Transformation: Radiation Oncology in Poland. International Journal of Radiation Oncology Biology Physics, 2014, 90, 480-486.	0.4	4
103	Impact of mTOR expression on clinical outcome in paediatric patients with B-cell acute lymphoblastic leukaemia â€“ preliminary report. Wspolczesna Onkologia, 2016, 4, 291-296.	0.7	4
104	Perforin gene variation influences survival in childhood acute lymphoblastic leukemia. Leukemia Research, 2018, 65, 29-33.	0.4	4
105	Achieving target levels for vascular risk parameters in Polish school-age children with type 1 diabetes â€“ a single center study. Journal of Pediatric Endocrinology and Metabolism, 2018, 31, 1073-1079.	0.4	4
106	MLPA as a complementary tool for diagnosis of chromosome 21 aberrations in childhood BCP-ALL. Journal of Applied Genetics, 2019, 60, 347-355.	1.0	4
107	Dynamic changes in specific anti-L-asparaginase antibodies generation during acute lymphoblastic leukemia treatment. Pharmacological Reports, 2019, 71, 311-318.	1.5	4
108	Advantages and Limitations of SNP Array in the Molecular Characterization of Pediatric T-Cell Acute Lymphoblastic Leukemia. Frontiers in Oncology, 2020, 10, 1184.	1.3	4

#	Article	IF	Citations
109	High Frequency of Fusion Gene Transcript Resulting From t(10;11)(p12;q23) Translocation in Pediatric Acute Myeloid Leukemia in Poland. Frontiers in Pediatrics, 2020, 8, 278.	0.9	4
110	A multistep approach to the genotype-phenotype analysis of Polish patients with tuberous sclerosis complex. European Journal of Medical Genetics, 2021, 64, 104309.	0.7	4
111	Alkaptonuria in a boy with type 1 diabetes mellitus, vitiligo, autoimmune thyroiditis and immunoglobulin A deficiency â€“ a case report. Pediatric Endocrinology, Diabetes and Metabolism, 2016, 22, 71-75.	0.3	4
112	Reduced Corneal Sensitivity With Neuronal Degeneration is a Novel Clinical Feature in Wolfram Syndrome. American Journal of Ophthalmology, 2022, 236, 63-68.	1.7	4
113	Molecular Genetics Diversity of Primary Hemophagocytic Lymphohistiocytosis among Polish Pediatric Patients. Archivum Immunologiae Et Therapiae Experimentalis, 2021, 69, 31.	1.0	4
114	Case report: Neonatal diabetes mellitus with congenital hypothyroidism as a result of biallelic heterozygous mutations in <scp> GLIS3 </scp> â€‰gene. Pediatric Diabetes, 2022, 23, 668-674.	1.2	4
115	Single patient in GCK-MODY family successfully re-diagnosed into GCK-PNDM through targeted next-generation sequencing technology. Acta Diabetologica, 2016, 53, 337-338.	1.2	3
116	Puzzling outcome of the nationwide genetic survey of severe/moderate female haemophilia B in Poland. Haemophilia, 2019, 25, e373-e376.	1.0	3
117	The Broad Variability in Dental Age Observed among Childhood Survivors Is Cancer Specific. Cancer Research and Treatment, 2021, 53, 252-260.	1.3	3
118	Severe and fatal toxicity after hematopoietic stem cell transplantation in GNE defect-associated thrombocytopenia. Bone Marrow Transplantation, 2021, 56, 1714-1716.	1.3	3
119	Prognostic Value of Rare IKZF1 deletions in Childhood B-Cell Precursor Acute Lymphoblastic Leukemia: An International Collaborative Study. Blood, 2014, 124, 368-368.	0.6	3
120	Evaluation of Changes to the Oral Microbiome Based on 16S rRNA Sequencing among Children Treated for Cancer. Cancers, 2022, 14, 7.	1.7	3
121	mTOR Inhibitor Treatment in Patients with Tuberous Sclerosis Complex Is Associated with Specific Changes in microRNA Serum Profile. Journal of Clinical Medicine, 2022, 11, 3395.	1.0	3
122	Availability and outcomes of radiotherapy in Central Poland during the 2005-2012 period - an observational study. BMC Cancer, 2015, 15, 214.	1.1	2
123	Everolimus treatment among patients with tuberous sclerosis affects serum lipid profile. Pharmacological Reports, 2016, 68, 1002-1007.	1.5	2
124	Uncommon reasons of the digestive tract-related paraneoplastic syndromes in children with neuroblastic tumors: three case reports. Wspolczesna Onkologia, 2018, 22, 42-46.	0.7	2
125	Recurrent and novel diseaseâ€causing F8 variants in boys with severe haemophilia A in Poland. Haemophilia, 2019, 25, e311-e314.	1.0	2
126	Salivary immunoglobulin A level during steroids and chemotherapy treatment administered in remission induction phase among pediatric patients with acute lymphoblastic leukemia. Medicine (United States), 2020, 99, e22802.	0.4	2

#	Article	IF	Citations
127	A Retrospective Evaluation of the Effects of Severe Aplastic Anemia Treatment in Children with Horse and Rabbit ATG.Polish Pediatric Hematology Group. Blood, 2011, 118, 4375-4375.	0.6	2
128	A randomized pharmacokinetic and pharmacodynamic trial of two regular human insulins demonstrates bioequivalence in type 1 diabetes and availability of biosimilar insulin may improve access to this medication. Diabetes, Obesity and Metabolism, 2022, , .	2.2	2
129	Editorial Room for the candidate gene approach in the genome-wide association era: an example from GSTP1 gene polymorphism. Archives of Medical Science, 2012, 4, 606-607.	0.4	1
130	<scp>UVR</scp> protection influences fructosamine level after sun exposure of healthy adults. Photodermatology Photoimmunology and Photomedicine, 2016, 32, 296-303.	0.7	1
131	Neonatal outcome and diabetes course in children with GCK-MODY born from women with GCK-MODY. Pediatric Endocrinology, Diabetes and Metabolism, 2018, 24, 167-173.	0.3	1
132	Central Corneal Thickness can be Related to Diabetic Peripheral Neuropathy in Children with Type 1 Diabetes. Experimental and Clinical Endocrinology and Diabetes, 2019, 127, 672-676.	0.6	1
133	Serum microRNA profiles in patients with autosomal dominant polycystic kidney disease show systematic dysregulation partially reversible by hemodialysis. Archives of Medical Science, 2019, 17, 1730-1741.	0.4	1
134	Pulmonary Exacerbation of Undiagnosed Toxocariasis in Intensively-Treated High-Risk Neuroblastoma Patients. Children, 2020, 7, 169.	0.6	1
135	HLA-A gene variation modulates residual function of the pancreatic Î²-cells in children with type 1 diabetes. Pediatric Endocrinology, Diabetes and Metabolism, 2020, 26, 73-78.	0.3	1
136	Evaluation of Three Lancing Devices: What Do Blood Volume and Lancing Pain Depend On?. Journal of Diabetes Science and Technology, 2021, 15, 1076-1083.	1.3	1
137	Serum microRNA as indicators of Wolfram syndromeâ€™s progression in neuroimaging studies. BMJ Open Diabetes Research and Care, 2020, 8, e001379.	1.2	1
138	Treatment Outcome and the Genetic Characteristics of Acute Promyelocytic Leukemia in Children in Poland From 2005 to 2018. Frontiers in Pediatrics, 2020, 8, 86.	0.9	1
139	Broad phenotypic spectrum of germ line 7p12.1 microdeletions encompassing the IKZF1 gene includes predisposition to acute lymphoblastic leukemia. Genes Chromosomes and Cancer, 2021, 60, 79-87.	1.5	1
140	Successful Salvage Haploidentical Bone Marrow Transplantation in a Child With Hemophagocytic Lymphohistiocytosis, When the Previously Matched Unrelated Donor Tested Positive for SARS-CoV-2 on the Day of Stem Cells Collection. Transplantation Proceedings, 2021, 53, 2498-2501.	0.3	1
141	New Insights into Red Blood Cell Microcytosis upon mTOR Inhibitor Administration. International Journal of Molecular Sciences, 2021, 22, 6802.	1.8	1
142	Genetic counseling in monogenic diabetes GCK MODY. Pediatric Endocrinology, Diabetes and Metabolism, 2016, 22, 54-59.	0.3	1
143	Gene expression of ASNS, LGMN and CTSB is elevated in a subgroup of childhood BCPâ€‘ALL with PAX5 deletion. Oncology Letters, 2019, 18, 6926-6932.	0.8	1
144	Polymorphisms Of Human Leukocyte Antigen-G Gene and Clinical Outcome Of Patients With Chronic Lymphocytic Leukemia. Blood, 2013, 122, 4151-4151.	0.6	1

#	Article	IF	Citations
145	Coexisting psoriasis affects the clinical course of type 1 diabetes in children. Pediatric Endocrinology, Diabetes and Metabolism, 2017, 23, 139-145.	0.3	1
146	Elevated level of lysophosphatidic acid among patients with HNF1B mutations and its role in RCAD syndrome: a multiomic study. Metabolomics, 2022, 18, 15.	1.4	1
147	Comparative Study of Malocclusions between Cancer Patients and Healthy Peers. International Journal of Environmental Research and Public Health, 2022, 19, 4045.	1.2	1
148	Selected risk factors of fractures in children â€" own observation. Open Medicine (Poland), 2012, 7, 635-641.	0.6	0
149	Successful combination treatment of a bifocal secretory germinoma with brain stem compression in a 17-year-old girl. Archives of Medical Science, 2016, 3, 678-680.	0.4	0
150	Successful Salvage Haploidentical Alpha-Beta T Cellâ€“Depleted Stem Cell Transplantation After Busulfan-Based Myeloablation in a Patient With IPEX Syndrome: A Case Report. Transplantation Proceedings, 2019, 51, 3150-3154.	0.3	0
151	Response to the Letter to the Editor: â€œCorneal thickness, optic nerve sheath diameter and retinal nerve fiber layer evaluation to assess the risk of cerebral edema in type 1 diabetes in childrenâ€• Acta Diabetologica, 2019, 56, 487-488.	1.2	0
152	Sex hormones and insulin sensitivity in adolescent girls with type 1 diabetes. Diabetes and Metabolism, 2020, 46, 75-77.	1.4	0
153	Novel FANCA mutation in the first fully-diagnosed patient with Fanconi anemia in Polish population â€" case report. Molecular Cytogenetics, 2020, 13, 33.	0.4	0
154	Successful Allogeneic Stem Cell Transplantation in Nuclear Factor-Kappa B Essential Modulator Deficiency Syndrome After Treosulfan-Based Conditioning: A Case Report. Transplantation Proceedings, 2020, 52, 647-652.	0.3	0
155	Evaluation of skin autofluorescence as a surrogate of advanced glycation end products accumulation in children and adolescents with normal haemoglobin A1c values. Pediatric Endocrinology, Diabetes and Metabolism, 2020, 26, 1-9.	0.3	0
156	Clinical heterogeneity among pediatric patients with autoimmune type 1 diabetes stratified by immunoglobulin deficiency. Pediatric Diabetes, 2021, 22, 707-716.	1.2	0
157	HL-A*11:01, -B*51:01, -DQB1*02:02 and -DRB1*07:01 are associated with inhibitor development in boys with severe haemophilia A receiving rFVIII prophylaxis in Poland. Thrombosis Research, 2021, 202, 170-172.	0.8	0
158	Results of Immunosupresive Therapy in Children with Aquired Severe Aplastic Anaemia (SAA). Report Polish Pediatric Hematology Group. Blood, 2008, 112, 4123-4123.	0.6	0
159	Antithymocyte Rabbit Globulin Is Safe and Effective in Children with Severe Aplastic Anemia Report Polish Pediatric Hematology Group (Warszawa, Krakow, Wroclaw, Bydgoszcz, Poznan, Lodz, Bialystok,) Tj ETQq1 1 0.784314 rgBT /Ov		
160	The Pro12Ala PPARg2 gene polymorphism involves residual C-peptide secretionand BMI in type 1 diabetes. Pediatric Endocrinology, Diabetes and Metabolism, 2014, 20, 88-94.	0.3	0
161	Human Leukocyte Antigen-G Polymorphisms Influence Clinical Outcome in Diffuse Large B-Cell Lymphoma. Blood, 2014, 124, 1643-1643.	0.6	0
162	HLA-G and MHC Class II Protein Expression in Diffuse Large B-Cell Lymphoma. Blood, 2014, 124, 1642-1642.	0.6	0

#	Article	IF	Citations
163	Infectious Complications in Children with ALL Treated with ALL-IC-2009 Protocol: Multicenter National Study of Polish Society of Pediatric Hematology and Oncology. Blood, 2014, 124, 5247-5247.	0.6	0
164	Vaccination in children with chronic severe neutropenia â€“ review of recommendations and a practical approach. Central-European Journal of Immunology, 2020, 45, 202-205.	0.4	0
165	Germline 3p22.1 microdeletion encompassing RPSA gene is an ultra-rare cause of isolated asplenia. Molecular Cytogenetics, 2021, 14, 51.	0.4	0
166	Psychiatric comorbidities in pediatric monogenic diabetes due to GCK mutation and their impact on the diabetes-related quality of life compared with type 1 diabetes.. Journal of the Academy of Consultation-Liaison Psychiatry, 2022, , .	0.2	0