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	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
2	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
3	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
4	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
5	Comparative Study of Malocclusions between Cancer Patients and Healthy Peers. International Journal of Environmental Research and Public Health, 2022, 19, 4045.	1.2	1
6	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
7	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
8	Evaluation of Changes to the Oral Microbiome Based on 16S rRNA Sequencing among Children Treated for Cancer. Cancers, 2022, 14, 7.	1.7	3
9	Consensus Recommendations for the Clinical Management of Hematological Malignancies in Patients with DNA Double Stranded Break Disorders. Cancers, 2022, 14, 2000.	1.7	5
10	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
11	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
12	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
13	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
14	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
15	SARS-CoV-2 Mpro inhibitors and activity-based probes for patient-sample imaging. Nature Chemical Biology, 2021, 17, 222-228.	3.9	215
16	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
17	The Broad Variability in Dental Age Observed among Childhood Survivors Is Cancer Specific. Cancer Research and Treatment, 2021, 53, 252-260.	1.3	3
18	Severe and fatal toxicity after hematopoietic stem cell transplantation in GNE defect-associated thrombocytopenia. Bone Marrow Transplantation, 2021, 56, 1714-1716.	1.3	3

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19	Clinical heterogeneity among pediatric patients with autoimmune type 1 diabetes stratified by immunoglobulin deficiency. Pediatric Diabetes, 2021, 22, 707-716.	1.2	0
20	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
21	Ovarian carcinoma in children with constitutional mutation of SMARCA4: single-family report and literature review. Familial Cancer, 2021, 20, 355-362.	0.9	8
22	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
23	Neutrophil Elastase Defects in Congenital Neutropenia. Frontiers in Immunology, 2021, 12, 653932.	2.2	18
24	New Insights into Red Blood Cell Microcytosis upon mTOR Inhibitor Administration. International Journal of Molecular Sciences, 2021, 22, 6802.	1.8	1
25	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
26	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
27	Genetic predisposition to lymphomas: Overview of rare syndromes and inherited familial variants. Mutation Research - Reviews in Mutation Research, 2021, 788, 108386.	2.4	9
28	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
29	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
30	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
31	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
32	Molecular Genetics Diversity of Primary Hemophagocytic Lymphohistiocytosis among Polish Pediatric Patients. Archivum Immunologiae Et Therapiae Experimentalis, 2021, 69, 31.	1.0	4
33	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
34	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
35	Germline 3p22.1 microdeletion encompassing RPSA gene is an ultra-rare cause of isolated asplenia. Molecular Cytogenetics, 2021, 14, 51.	0.4	0
36	Sex hormones and insulin sensitivity in adolescent girls with type 1 diabetes. Diabetes and Metabolism, 2020, 46, 75-77.	1.4	0

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37	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
38	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
39	Pulmonary Exacerbation of Undiagnosed Toxocariasis in Intensively-Treated High-Risk Neuroblastoma Patients. Children, 2020, 7, 169.	0.6	1
40	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
41	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
42	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
43	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
44	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
45	Multiple Retinal Anomalies in Wfs1-Deficient Mice. Diagnostics, 2020, 10, 607.	1.3	5
46	Results of two consecutive treatment protocols in Polish children with acute lymphoblastic leukemia. Scientific Reports, 2020, 10, 20168.	1.6	9
47	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
48	Nivolumab for the Treatment of Advanced Pediatric Malignancies. Anticancer Research, 2020, 40, 7095-7100.	0.5	14
49	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
50	Serum microRNA as indicators of Wolfram syndromeâ€™s progression in neuroimaging studies. BMJ Open Diabetes Research and Care, 2020, 8, e001379.	1.2	1
51	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
52	Six molecular patterns leading to hemophilia A phenotype in 18 females from Poland. Thrombosis Research, 2020, 193, 9-14.	0.8	8
53	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
54	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

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55	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
56	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
57	Corneal Abnormalities Are Novel Clinical Feature in Wolfram Syndrome. American Journal of Ophthalmology, 2020, 217, 140-151.	1.7	7
58	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
59	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
60	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
61	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
62	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
63	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
64	Puzzling outcome of the nationwide genetic survey of severe/moderate female haemophilia B in Poland. Haemophilia, 2019, 25, e373-e376.	1.0	3
65	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
66	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
67	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
68	Dental caries among childhood cancer survivors. Medicine (United States), 2019, 98, e14279.	0.4	9
69	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
70	Recurrent and novel diseaseâ€causing F8 variants in boys with severe haemophilia A in Poland. Haemophilia, 2019, 25, e311-e314.	1.0	2
71	Targeting the thioredoxin system as a novel strategy against Bâ€cell acute lymphoblastic leukemia. Molecular Oncology, 2019, 13, 1180-1195.	2.1	24
72	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

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73	Dynamic changes in specific anti-L-asparaginase antibodies generation during acute lymphoblastic leukemia treatment. Pharmacological Reports, 2019, 71, 311-318.	1.5	4
74	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
75	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
76	Unfavorable Outcome of Neuroblastoma in Patients With 2p Gain. Frontiers in Oncology, 2019, 9, 1018.	1.3	12
77	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
78	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
79	Determination and interpretation of MTHFR gene mutations in gynecology and internal medicine. Polish Archives of Internal Medicine, 2019, 129, 728-732.	0.3	5
80	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
81	GlyCulator2: an update on a web application for calculation of glycemic variability indices. Acta Diabetologica, 2018, 55, 877-880.	1.2	22
82	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
83	Perforin gene variation influences survival in childhood acute lymphoblastic leukemia. Leukemia Research, 2018, 65, 29-33.	0.4	4
84	Monogenic diabetes prevalence among Polish children-Summary of 11â€‰years-long nationwide genetic screening program. Pediatric Diabetes, 2018, 19, 53-58.	1.2	21
85	NIRCa: An artificial neural network-based insulin resistance calculator. Pediatric Diabetes, 2018, 19, 231-235.	1.2	11
86	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
87	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
88	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
89	Defective DNA damage repair leads to frequent catastrophic genomic events in murine and human tumors. Nature Communications, 2018, 9, 4760.	5.8	66
90	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

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91	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
92	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
93	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
94	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
95	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
96	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
97	Monogenic diabetes syndromes: Locus-specific databases for AlstrÃ¶m, Wolfram, and Thiamine-responsive megaloblastic anemia. Human Mutation, 2017, 38, 764-777.	1.1	47
98	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
99	Polymorphism in IKZF1 gene affects clinical outcome in diffuse large B-cell lymphoma. International Journal of Hematology, 2017, 106, 794-800.	0.7	6
100	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
101	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
102	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
103	Biallelic loss ofCDKN2Ais associated with poor response to treatment in pediatric acute lymphoblastic leukemia. Leukemia and Lymphoma, 2017, 58, 1162-1171.	0.6	43
104	Intragenic amplification of PAX5: a novel subgroup in B-cell precursor acute lymphoblastic leukemia?. Blood Advances, 2017, 1, 1473-1477.	2.5	25
105	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
106	Coexisting psoriasis affects the clinical course of type 1 diabetes in children. Pediatric Endocrinology, Diabetes and Metabolism, 2017, 23, 139-145.	0.3	1
107	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
108	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

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109	Dental Anomalies as Late Adverse Effect among Young Children Treated for Cancer. Cancer Research and Treatment, 2016, 48, 658-667.	1.3	33
110	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
111	Breaking the Taboo: Illicit Drug Use among Adolescents with Type 1 Diabetes Mellitus. Journal of Diabetes Research, 2016, 2016, 1-8.	1.0	11
112	Chromosome 18q deletion syndrome with autoimmune diabetes mellitus: putative genomic loci for autoimmunity and immunodeficiency. Pediatric Diabetes, 2016, 17, 153-159.	1.2	7
113	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
114	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
115	Differential regulation of serum microRNA expression by HNF1Î² and HNF1Î± transcription factors. Diabetologia, 2016, 59, 1463-1473.	2.9	18
116	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
117	Everolimus treatment among patients with tuberous sclerosis affects serum lipid profile. Pharmacological Reports, 2016, 68, 1002-1007.	1.5	2
118	<scp>UVR</scp> protection influences fructosamine level after sun exposure of healthy adults. Photodermatology Photoimmunology and Photomedicine, 2016, 32, 296-303.	0.7	1
119	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
120	Psychiatric Disorders and Health-Related Quality of Life in Children With Type 1 Diabetes Mellitus. Psychosomatics, 2016, 57, 185-193.	2.5	47
121	Clinical course and therapeutic implications for lymphoid malignancies in Nijmegen breakage syndrome. European Journal of Medical Genetics, 2016, 59, 126-132.	0.7	38
122	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
123	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
124	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
125	Genetic counseling in monogenic diabetes GCK MODY. Pediatric Endocrinology, Diabetes and Metabolism, 2016, 22, 54-59.	0.3	1
126	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

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127	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
128	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
129	Availability and outcomes of radiotherapy in Central Poland during the 2005-2012 period - an observational study. BMC Cancer, 2015, 15, 214.	1.1	2
130	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
131	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
132	Intima-media thickness and endothelial dysfunction in GCK and HNF1A-MODY patients. European Journal of Endocrinology, 2015, 172, 277-283.	1.9	12
133	Retinal Thinning as a Marker of Disease Progression in Patients With Wolfram Syndrome. Diabetes Care, 2015, 38, e36-e37.	4.3	21
134	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
135	Altered Plateletsâ€™ morphological parameters in children with type 1 diabetes â€“ a case-control study. BMC Endocrine Disorders, 2015, 15, 17.	0.9	23
136	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
137	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
138	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
139	Central Nervous System PET-CT Imaging Reveals Regional Impairments in Pediatric Patients with Wolfram Syndrome. PLoS ONE, 2014, 9, e115605.	1.1	9
140	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
141	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
142	GATA4 Mutations Are a Cause of Neonatal and Childhood-Onset Diabetes. Diabetes, 2014, 63, 2888-2894.	0.3	108
143	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
144	Polymorphism inIKZF1gene affects age at onset of childhood acute lymphoblastic leukemia. Leukemia and Lymphoma, 2014, 55, 2174-2178.	0.6	13

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145	Tackling the Turmoil of Transformation: Radiation Oncology in Poland. International Journal of Radiation Oncology Biology Physics, 2014, 90, 480-486.	0.4	4
146	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
147	Palliative Sedation at Home for Terminally Ill Children With Cancer. Journal of Pain and Symptom Management, 2014, 48, 968-974.	0.6	27
148	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
149	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
150	Ocena czÄ™stoÅ›ci wystÄ™powania zespoÅ‚u Wolframa w populacji dzieci z cukrzycÄ…. Endokrynologia Polska, 2014, 65, 295-297.	0.3	8
151	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
152	Human Leukocyte Antigen-G Polymorphisms Influence Clinical Outcome in Diffuse Large B-Cell Lymphoma. Blood, 2014, 124, 1643-1643.	0.6	0
153	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
154	HLA-G and MHC Class II Protein Expression in Diffuse Large B-Cell Lymphoma. Blood, 2014, 124, 1642-1642.	0.6	0
155	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
156	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
157	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
158	Interleukin 18 as a Marker of Chronic Nephropathy in Children after Anticancer Treatment. Disease Markers, 2013, 35, 811-818.	0.6	22
159	The Impact of Pediatric Palliative Care Education on Medical Studentsâ€™ Knowledge and Attitudes. Scientific World Journal, The, 2013, 2013, 1-9.	0.8	12
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