

Kamil Hozyasz

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

90
papers

607
citations

14
h-index

22
g-index

102
ext. papers

699
ext. citations

2
avg, IF

3.85
L-index

#	Paper	IF	Citations
90	Maternal MTR genotype contributes to the risk of non-syndromic cleft lip and palate in the Polish population. <i>Clinical Genetics</i> , 2006 , 69, 512-7	4	69
89	Associations of folate and choline metabolism gene polymorphisms with orofacial clefts. <i>Journal of Medical Genetics</i> , 2010 , 47, 809-15	5.8	60
88	Association between genetic variants of reported candidate genes or regions and risk of cleft lip with or without cleft palate in the Polish population. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2010 , 88, 538-45		47
87	Immune Abnormalities in Autism Spectrum Disorder-Could They Hold Promise for Causative Treatment?. <i>Molecular Neurobiology</i> , 2018 , 55, 6387-6435	6.2	45
86	Genotype and haplotype analysis of WNT genes in non-syndromic cleft lip with or without cleft palate. <i>European Journal of Oral Sciences</i> , 2012 , 120, 1-8	2.3	34
85	Polymorphisms located in the region containing BHMT and BHMT2 genes as maternal protective factors for orofacial clefts. <i>European Journal of Oral Sciences</i> , 2010 , 118, 325-32	2.3	24
84	Relation between the concentration of zinc in maternal whole blood and the risk of an infant being born with an orofacial cleft. <i>British Journal of Oral and Maxillofacial Surgery</i> , 2009 , 47, 466-9	1.4	24
83	Stem cell regenerative therapy in alveolar cleft reconstruction. <i>Archives of Oral Biology</i> , 2015 , 60, 1517-328		21
82	Polymorphic variants at 10q25.3 and 17q22 loci and the risk of non-syndromic cleft lip and palate in the Polish population. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2012 , 94, 42-6		21
81	Common variants in DLG1 locus are associated with non-syndromic cleft lip with or without cleft palate. <i>Clinical Genetics</i> , 2018 , 93, 784-793	4	20
80	Nasolabial symmetry and aesthetics in children with complete unilateral cleft lip and palate. <i>British Journal of Oral and Maxillofacial Surgery</i> , 2012 , 50, 621-5	1.4	20
79	Polymorphic variants of genes involved in homocysteine metabolism in celiac disease. <i>Molecular Biology Reports</i> , 2012 , 39, 3123-30	2.8	19
78	Nucleotide variants of the cancer predisposing gene CDH1 and the risk of non-syndromic cleft lip with or without cleft palate. <i>Familial Cancer</i> , 2014 , 13, 415-21	3	18
77	Association of DVL2 and AXIN2 gene polymorphisms with cleft lip with or without cleft palate in a Polish population. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2012 , 94, 943-50		17
76	Association between polymorphisms at the GREM1 locus and the risk of nonsyndromic cleft lip with or without cleft palate in the Polish population. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2015 , 103, 847-56		9
75	Identification of novel cystinuria mutations in pediatric patients. <i>Journal of Pediatric Urology</i> , 2006 , 2, 575-8	1.5	9
74	Polymorphisms of stress-related genes and the risk of nonsyndromic cleft lip with or without cleft palate. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2011 , 91, 948-55		8

73	PAX7 nucleotide variants and the risk of non-syndromic orofacial clefts in the Polish population. <i>Oral Diseases</i> , 2019 , 25, 1608-1618	3.5	7
72	A mutation in mouse Pak1ip1 causes orofacial clefting while human PAK1IP1 maps to 6p24 translocation breaking points associated with orofacial clefting. <i>PLoS ONE</i> , 2013 , 8, e69333	3.7	7
71	High manganese levels in milk-based infant formulas. <i>NeuroToxicology</i> , 2004 , 25, 733	4.4	7
70	Coeliac disease and birth defects in offspring. <i>Gut</i> , 2001 , 49, 738	19.2	7
69	Genetic variants in BRIP1 (BACH1) contribute to risk of nonsyndromic cleft lip with or without cleft palate. <i>Birth Defects Research Part A: Clinical and Molecular Teratology</i> , 2014 , 100, 670-8		6
68	Malonylcarnitine in newborns with non-syndromic cleft lip with or without cleft palate. <i>International Journal of Oral Science</i> , 2010 , 2, 136-41	27.9	6
67	Polymorphic variants of genes encoding main antioxidant enzymes and the risk of CL/P-affected pregnancies. <i>Clinical Biochemistry</i> , 2007 , 40, 416-9	3.5	5
66	Whole blood citrulline concentrations in newborns with non-syndromic oral clefts--a preliminary report. <i>Asia Pacific Journal of Clinical Nutrition</i> , 2010 , 19, 217-22	1	5
65	Further Evidence of the Association of the Diacylglycerol Kinase Kappa (DGKK) Gene With Hypospadias. <i>Urology Journal</i> , 2018 , 15, 272-276	0.9	5
64	WartoŹdrowotna produktŹ kokosowych. <i>Pediatrics Polska</i> , 2015 , 90, 415-423	0.1	4
63	Association of CDKAL1 nucleotide variants with the risk of non-syndromic cleft lip with or without cleft palate. <i>Journal of Human Genetics</i> , 2018 , 63, 397-406	4.3	4
62	Re: High serum levels of 8-hydroxy-2'-deoxyguanosine (8-OHdG) in mothers of children with cleft lip. <i>British Journal of Oral and Maxillofacial Surgery</i> , 2003 , 41, 205-6	1.4	4
61	Analysis of the concentration of vitamin E in erythrocytes of patients with celiac disease. <i>Przegląd Gastroenterologiczny</i> , 2016 , 11, 282-285	6	3
60	Dental arch relationship in 5-year-olds with complete unilateral cleft lip and palate after early alveolar bone grafting. <i>Orthodontics and Craniofacial Research</i> , 2012 , 15, 117-23	3	3
59	Whole blood propionylcarnitine in newborns with orofacial cleft. <i>Maternal and Child Nutrition</i> , 2011 , 7, 100-3	3.4	3
58	The search for risk factors that contribute to the etiology of non-syndromic cleft lip with or without cleft palate (CL/P) in the Polish population. <i>Pediatrics Polska</i> , 2010 , 85, 609-623	0.1	3
57	Polymorphic variants of genes related to arginine metabolism and the risk of orofacial clefts. <i>Archives of Oral Biology</i> , 2010 , 55, 861-6	2.8	3
56	Re: Pregnancy outcomes in celiac women. <i>American Journal of Gastroenterology</i> , 2000 , 95, 1373-4	0.7	3

55	Searching for new genes and loci involved in cleft lip and palate in the Polish population – genome-wide association study. <i>Journal of Medical Science</i> , 2014 , 83, 265-268	1.6	3
54	Propionylcarnitine and methionine concentrations in newborns with hypospadias. <i>Central European Journal of Urology</i> , 2013 , 66, 377-80	0.9	3
53	80-lecie fenylketonurii. Czł: historia nazwy i nietuzinkowi pionierzy badań chorób. <i>Pediatrica Polska</i> , 2016 , 91, 386-391	0.1	2
52	Ekspozycja na napoje w sklepach a żywienie niemowląt: brak społecznie odpowiedzialnego marketingu w Polsce?. <i>Pediatrica Polska</i> , 2013 , 88, 164-169	0.1	2
51	Alternative n-3 PUFAs Sources in Central European Diet before Westernization – Case Report from Poland. <i>Journal of Food Research</i> , 2013 , 2, 29	1.3	2
50	The Mediterranean diet for Polish infants: a losing struggle or a battle still worth fighting?. <i>Mediterranean Journal of Nutrition and Metabolism</i> , 2010 , 3, 227-232	1.3	2
49	Sex ratio variation in offspring of women with celiac disease. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2002 , 105, 195	2.4	2
48	Lack of association of polymorphic variants of genes encoding zinc transporters with the risk of orofacial cleft-affected pregnancies. <i>Folia Histochemica Et Cytobiologica</i> , 2010 , 48, 618-23	1.4	2
47	Non-celiac gluten sensitivity (NCGS) – an old diagnosis recently rediscovered. <i>Family Medicine and Primary Care Review</i> , 2016 , 1, 79-83	0.6	2
46	Napiętnowanie społeczne w chorobach przewlekłych. <i>Pediatrica Polska</i> , 2017 , 92, 316-320	0.1	1
45	Pieczyno z odroczonego wypieku. Badanie dostępności i składu. <i>Pediatrica Polska</i> , 2017 , 92, 156-163	0.1	1
44	Nucleotide Variants of the BH4 Biosynthesis Pathway Gene GCH1 and the Risk of Orofacial Clefts. <i>Molecular Neurobiology</i> , 2016 , 53, 769-776	6.2	1
43	Kompendium wiedzy o jajach. <i>Pediatrica Polska</i> , 2014 , 89, 313-322	0.1	1
42	Ortoreksja – patologiczna kontrola nad odżywianiem. Zagrożenie dla dzieci i młodzieży. <i>Pediatrica Polska</i> , 2014 , 89, 119-124	0.1	1
41	Aronia czarnoowocowa – kliniczne perspektywy. <i>Pediatrica Polska</i> , 2013 , 88, 452-458	0.1	1
40	Nieprawidłowa relacja pomiędzy matką a dzieckiem jako przyczyna zaburzeń karmienia u niemowląt – opis przypadku. <i>Pediatrica Polska</i> , 2013 , 88, 472-476	0.1	1
39	Ocena spożycia wód smakowych przez dzieci kierowane do oddziału pediatrycznego. <i>Pediatrica Polska</i> , 2017 , 92, 401-405	0.1	1
38	Słabikowe dania z mlekiem dla niemowląt – przykład zdrowej diety. <i>Pediatrica Polska</i> , 2014 , 89, 240-244	0.1	1

37	Association of common variants in PAH and LAT1 with non-syndromic cleft lip with or without cleft palate (NSCL/P) in the Polish population. <i>Archives of Oral Biology</i> , 2014 , 59, 363-9	2.8	1
36	Paleodieta [czy dieta przodków] może zapobiegać chorobom cywilizacyjnym?. <i>Pediatrics Polska</i> , 2014 , 89, 261-268	0.1	1
35	Gluten-free diet in the treatment of autism spectrum disorders – short overview. <i>Przegląd Gastroenterologiczny</i> , 2010 , 4, 195-201	6	1
34	Stężenia glutarylkarnityny (C5DC) u noworodków z rozszczepem twarzoczaszki. <i>Pediatrics Polska</i> , 2010 , 85, 239-242	0.1	1
33	Fever and high C-reactive protein level as the sole manifestation of allergy to cow milk. <i>Explore: the Journal of Science and Healing</i> , 2006 , 2, 520-1	1.4	1
32	Low increase in phenylalanine tolerance during pregnancies in PKU woman with high prepregnancy BMI and postconceptional initiation of diet: A case report. <i>International Journal of Reproductive BioMedicine</i> , 2019 , 17, 763-770	1.3	1
31	Celiac disease presenting during puerperium. <i>Journal of Clinical Gastroenterology</i> , 2003 , 36, 81-2	3	1
30	From rediscovered "biche" to "hainstream" – <i>Glyceria fluitans</i> as a candidate grain for manufacturing premium food products. <i>Plants People Planet</i> , 2020 , 2, 104-106	4.1	1
29	80-lecie fenyloketonurii. Cz. IV: Stanisław F. Bieszko i Ada J. Susi a historia opracowania testu Guthriego. <i>Pediatrics Polska</i> , 2017 , 92, 658-663	0.1	
28	80-lecie fenyloketonurii. Cz. III: Charles E. Dent – biochemik-naukowiec i klinicysta, który dostrzegł choroby matki u potomstwa i zrutynizował badanie profilu aminokwasów. <i>Pediatrics Polska</i> , 2017 , 92, 352-359	0.1	
27	Wpływ otyłości u matki w czasie ciąży na ryzyko rozwoju autyzmu u dziecka. <i>Pediatrics Polska</i> , 2015 , 90, 229-235	0.1	
26	RDW w praktyce klinicznej. <i>Pediatrics Polska</i> , 2015 , 90, 130-134	0.1	
25	Produkty imitujące mięsian – badanie dostępności w sklepach sieciowych oraz przyczynek do dyskusji o wartościach odżywczych i roli mięsian w żywieniu. <i>Pediatrics Polska</i> , 2016 , 91, 227-232	0.1	
24	Ocena sposobu żywienia dzieci kierowanych do oddziału pediatrycznego oraz wiedzy żywieniowej ich opiekunów – wyniki badania pilotażowego. <i>Pediatrics Polska</i> , 2016 , 91, 208-213	0.1	
23	Celiakia w polskojęzycznych publikacjach 1990-2012 – analiza bibliometryczna. <i>Pediatrics Polska</i> , 2014 , 89, 106-111	0.1	
22	Współwystępowanie dwóch rzadkich chorób genetycznych: fenyloketonurii oraz zespołu Pradera i Williego. Opis przypadku. <i>Pediatrics Polska</i> , 2014 , 89, 297-301	0.1	
21	Whole-blood 3-hydroxyisovalerylcarnitine as a risk factor for orofacial clefts. <i>Archives of Oral Biology</i> , 2013 , 58, 459-61	2.8	
20	Rola szczepu <i>Lactobacillus plantarum</i> 299v w zapobieganiu i leczeniu zaburzeń układu pokarmowego. <i>Pediatrics Polska</i> , 2013 , 88, 347-352	0.1	

- 19 Niemowlę opóźnionym rozwojem psychoruchowym i pomarańczowymi kryształami na pieluszcze
Opis przypadku zespołu Lescha i Nyhana. *Pediatrics Polska*, **2013**, 88, 286-289 0.1
- 18 Właściwości psychometryczne polskiej adaptacji kwestionariusza oceny zadowolenia rodziców z
ambulatoryjnej opieki pediatrycznej. *Pediatrics Polska*, **2017**, 92, 271-280 0.1
- 17 80-lecie fenylketonurii. Część I: pierwszy literacki oraz inne niemedyczne opisy choroby. *Pediatrics
Polska*, **2017**, 92, 218-226 0.1
- 16 Chrzan tarty dostępny na rynku a tradycja kulinarna w Polsce. *Pediatrics Polska*, **2015**, 90, 308-311 0.1
- 15 Zażółta w galaktozemii. *Pediatrics Polska*, **2014**, 89, 192-197 0.1
- 14 Trehalozę dwucukier o unikatowych właściwościach. *Pediatrics Polska*, **2012**, 87, 569-573 0.1
- 13 C14 and C16 acylcarnitines in newborns with orofacial clefts. *Przegląd Gastroenterologiczny*, **2012**,
5, 276-280 6
- 12 Alternatives to standard cow milk: pros and cons. *Przegląd Gastroenterologiczny*, **2013**, 2, 98-107 6
- 11 Letter to the editor re: Comino, I., et al. *Nutrients* 2013, 5, 4250-4268. *Nutrients*, **2013**, 5, 4964-5 6.7
- 10 Całkowita aktywność przeciwutleniająca osocza u chorych z kamieniami nerkowymi. *Pediatrics Polska*,
2007, 82, 857-859 0.1
- 9 A variable latent interval after exposure to gluten in persons developing celiac disease. *American
Journal of Medical Genetics, Part A*, **2008**, 146A, 539 2.5
- 8 Therapeutic modalities for celiac disease. *Explore: the Journal of Science and Healing*, **2006**, 2, 291;
author reply 291 1.4
- 7 European "gluten-free" solid foods for infants may be a risky food for celiacs. *Journal of the
American Dietetic Association*, **2002**, 102, 637
- 6 Is low male sex ratio in offspring of celiacs an advantage?. *American Journal of Gastroenterology*,
2002, 97, 1574 0.7
- 5 Addison's disease mimicking anorexia nervosa. *Clinical Pediatrics*, **1999**, 38, 561-2 1.2
- 4 Neurological manifestations in celiacs and vitamin E status. *Arquivos De Neuro-Psiquiatria*, **2005**, 63,
371; author reply 371-2 1.6
- 3 EVC gene polymorphisms and risks of isolated hypospadias - a preliminary study. *Central European
Journal of Urology*, **2015**, 68, 257-62 0.9
- 2 Comparison of phenylalanine tolerance in singleton and twin pregnancies in patients with
phenylketonuria. *Journal of International Medical Research*, **2020**, 48, 300060520934623 1.4

- 1 Niedobór witaminy B12 jako przyczyna nieustępujących trudności we wprowadzeniu posiłków uzupełniających u niemowlęcia. Opis przypadku. *Pediatrics Polska*, **2016**, 91, 484-491

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