Irena Jankowska

List of Publications by Year in Descending Order

69 papers	1,453 citations	14 h-index	37 g-index
71 ext. papers	1,689 ext. citations	3 avg, IF	3.33 L-index

#	Paper	IF	Citations
69	Autosomal Recessive Polycystic Kidney Disease-The Clinical Aspects and Diagnostic Challenges. *Journal of Pediatric Genetics*, **2021**, 10, 1-8	0.7	0
68	Cholestasis Due to USP53 Deficiency. *Journal of Pediatric Gastroenterology and Nutrition*, **2021**, 72, 667-67	2.8	8
67	A Report of 2 Infant Siblings with Progressive Intrahepatic Familial Cholestasis Type 1 and a Novel Homozygous Mutation in the ATP8B1 Gene Treated with Partial External Biliary Diversion and Liver Transplant. *American Journal of Case Reports*, **2021**, 22, e932374	1.3	2
66	Acute liver failure due to DGUOK deficiency-is liver transplantation justified?. *Clinics and Research in Hepatology and Gastroenterology*, **2021**, 45, 101408	2.4	2
65	Coronavirus Disease 2019-Liver Injury-Literature Review and Guidelines Based on the Recommendations of Hepatological Societies. *Pediatric Gastroenterology, Hepatology and Nutrition*, **2021**, 24, 119-126	2.3	1
64	Impact of Genotype, Serum Bile Acids, and Surgical Biliary Diversion on Native Liver Survival in FIC1 Deficiency. *Hepatology*, **2021**, 74, 892-906	11.2	6
63	NBAS deficiency due to biallelic c.2809C > G variant presenting with recurrent acute liver failure with severe hyperammonemia, acquired microcephaly and progressive brain atrophy. *Metabolic Brain Disease*, **2021**, 36, 2169-2172	3.9	0
62	Successful Liver Transplantation in Two Polish Brothers with Transaldolase Deficiency. *Children*, **2021**, 8,	2.8	0
61	Progressive familial intrahepatic cholestasis type 3: Report of four clinical cases, novel ABCB4 variants and long-term follow-up. *Annals of Hepatology*, **2021**, 25, 100342	3.1	1
60	Genotype correlates with the natural history of severe bile salt export pump deficiency. *Journal of Hepatology*, **2020**, 73, 84-93	13.4	22
59	Non-Hodgkin lymphoma after liver and kidney transplantation in children. Experience from one center. *Advances in Clinical and Experimental Medicine*, **2020**, 29, 197-202	1.8	1
58	Occurrence of Portal Hypertension and Its Clinical Course in Patients With Molecularly Confirmed Autosomal Recessive Polycystic Kidney Disease (ARPKD). *Frontiers in Pediatrics*, **2020**, 8, 591379	3.4	1
57	Transient Elastography for Detection of Liver Fibrosis in Children With Autosomal Recessive Polycystic Kidney Disease. *Frontiers in Pediatrics*, **2018**, 6, 422	3.4	8
56	Outcomes of surgical management of familial intrahepatic cholestasis 1 and bile salt export protein deficiencies. *Hepatology Communications*, **2018**, 2, 515-528	6	32
55	Long-Term Systematic Monitoring of Four Polish Transaldolase Deficient Patients. *JIMD Reports*, **2018**, 42, 79-87	1.9	8
54	Newborn presentation of Niemann-Pick disease type C - Difficulties and limitations of diagnostic methods. *Pediatrics and Neonatology*, **2018**, 59, 317-318	1.8	2
53	Endoscopic treatment in biliary strictures after pediatric liver transplantation. *Pediatric Transplantation*, **2018**, 22, e13271	1.8	2

52	Acute liver failure in children-Is living donor liver transplantation justified?. *PLoS ONE*, **2018**, 13, e0193327	3.7	7
51	Percutaneous Treatment of Biliary Strictures After Pediatric Liver Transplantation. *Annals of Transplantation*, **2018**, 23, 845-851	1.4	6
50	Prevalence of Cryptosporidium, Blastocystis, and other opportunistic infections in patients with primary and acquired immunodeficiency. *Parasitology Research*, **2018**, 117, 2869-2879	2.4	28
49	Congenital hepatic fibrosis associated with Gilbert's syndrome and cholelithiasis in a pediatric patient – The crucial role of liver biopsy in diagnostic process. *Pediatria Polska*, **2017**, 92, 446-449	0.1	
48	Co nowego w cholestazie – cz. 2. Cholestaza z podwyższoną aktywnością gamma-glutamylotranspeptydazy. *Pediatria Polska*, **2017**, 92, 412-416	0.1	
47	Co nowego w cholestazie – cz. 1. Cholestaza z prawidłową aktywnością gamma-glutamylotranspeptydazy. *Pediatria Polska*, **2017**, 92, 366-372	0.1	
46	Acute-on-chronic hepatitis. A case report of autoimmune hepatitis/primary sclerosing cholangitis/ulcerative colitis overlap syndrome in a 15-year-old patient. *Clinical and Experimental Hepatology*, **2017**, 3, 28-32	2.2	1
45	Łagodna nawracająca cholestaza wewnątrzwątrobowa. *Pediatria Polska*, **2017**, 92, 575-578	0.1	
44	Świąd skóry w przebiegu cholestatycznych chorób wątroby – aktualny stan wiedzy. *Pediatria Polska*, **2017**, 92, 669-674	0.1	
43	Co nowego w cholestazie – cz. 3. Wrodzone zaburzenia syntezy kwasów żółciowych. *Pediatria Polska*, **2017**, 92, 567-574	0.1	
42	Witamina K w cholestazie w świetle aktualnych wytycznych – opis przypadku. *Pediatria Polska*, **2017**, 92, 789-792	0.1	
41	Metabolizm bilirubiny i jego znaczenie w patogenezie zespołów Rotora oraz Dubina i Johnsona – aktualny stan wiedzy. *Pediatria Polska*, **2017**, 92, 745-747	0.1	
40	Choroby wątroby i nerek w przebiegu ciliopatii. *Pediatria Polska*, **2017**, 92, 121-128	0.1	
39	Obesity, lipid profiles and oxidative stress in children after liver transplantation. *Acta Biochimica Polonica*, **2017**, 64, 661-665	2	5
38	Congenital hepatic fibrosis in a 9-year-old female patient - a case report. *Clinical and Experimental Hepatology*, **2017**, 3, 176-179	2.2	5
37	Diagnostic Approach in Biliary Strictures After Pediatric Liver Transplantation. *Annals of Transplantation*, **2017**, 22, 257-264	1.4	4
36	Ogniskowy rozrost guzkowy wątroby – opis przypadków oraz przegląd literatury. *Pediatria Polska*, **2016**, 91, 359-365	0.1	
35	Choroba czy zespół Caroliego? Od zaburzeń embriogenezy do przeszczepienia wątroby. *Pediatria Polska*, **2016**, 91, 453-457	0.1	

34	Influence of Partial External Biliary Diversion on the Lipid Profile in Children With Progressive Familial Intrahepatic Cholestasis. *Journal of Pediatric Gastroenterology and Nutrition*, **2016**, 63, 598-602	2.8	6
33	Exocrine pancreatic function in children with Alagille syndrome. *Scientific Reports*, **2016**, 6, 35229	4.9	4
32	Successful pregnancy after ileal exclusion in progressive familial intrahepatic cholestasis type 2. *Annals of Hepatology*, **2015**, 14, 550-552	3.1	4
31	Immunological factors and liver fibrosis in pediatric liver transplant recipients. *Annals of Transplantation*, **2015**, 20, 279-84	1.4	18
30	Ileal exclusion in children with progressive familial intrahepatic cholestasis. *Journal of Pediatric Gastroenterology and Nutrition*, **2014**, 58, 92-5	2.8	19
29	Treatment of pruritus with Prometheus dialysis and absorption system in a patient with benign recurrent intrahepatic cholestasis. *Hepatology Research*, **2014**, 44, E304-E308	5.1	9
28	Antibody-mediated rejection in pediatric liver transplant recipients. *Annals of Transplantation*, **2014** , 19, 119-23	1.4	9
27	Cardiovascular risk factors after conversion from cyclosporine to tacrolimus in children after liver transplantation. *Annals of Transplantation*, **2014**, 19, 604-8	1.4	
26	Small intestinal bacterial overgrowth in patients with progressive familial intrahepatic cholestasis. *Acta Biochimica Polonica*, **2014**, 61, 103-7	2	3
25	The first case of Enterocytozoon bieneusi infection in Poland. *Annals of Agricultural and Environmental Medicine*, **2013**, 20, 287-8	1.4	10
24	Successful sirolimus rescue in tacrolimus-induced thrombotic microangiopathy after living-related liver transplantation. *Pediatric Transplantation*, **2012**, 16, E261-4	1.8	6
23	Lipid metabolism and oxidative stress in children after liver transplantation treated with sirolimus. *Pediatric Transplantation*, **2012**, 16, 901-6	1.8	4
22	The impact of cytokine gene polymorphisms on Epstein-Barr virus infection outcome in pediatric liver transplant recipients. *Journal of Clinical Virology*, **2012**, 55, 226-32	14.5	9
21	Progressive familial intrahepatic cholestasis and inborn errors of bile acid synthesis. *Clinics and Research in Hepatology and Gastroenterology*, **2012**, 36, 271-4	2.4	12
20	Cholestatic Syndromes in Childhood and Catch-Up Growth **2012**, 863-879		
19	Epstein-Barr virus gene expression and latent membrane protein 1 gene polymorphism in pediatric liver transplant recipients. *Journal of Medical Virology*, **2011**, 83, 2182-90	19.7	5
18	Wrodzone wnienie wroby **2011**, 11, 97-102		2
17	Long-term monitoring of Epstein-Barr virus DNA load and humoral parameter abnormalities in pediatric liver transplant recipients before development of malignancy. *Pediatric Transplantation*, **2010**, 14, 629-35	1.8	4

16	Differences in presentation and progression between severe FIC1 and BSEP deficiencies. *Journal of Hepatology*, **2010**, 53, 170-8	13.4	145
15	beta-Carotene deficiency in cholestatic liver disease of childhood is caused by beta-carotene malabsorption. *Journal of Pediatric Gastroenterology and Nutrition*, **2010**, 51, 106-9	2.8	3
14	Normal levels of serum pancreatic enzymes in patients with progressive familial intrahepatic cholestasis type 2. *Acta Biochimica Polonica*, **2010**, 57, 573-5	2	1
13	Hemodynamic failure as an indication to urgent liver transplantation in infants with giant hepatic hemangiomas or vascular malformations--report of four cases. *Pediatric Transplantation*, **2009**, 13, 906-12	1.8	11
12	Severe bile salt export pump deficiency: 82 different ABCB11 mutations in 109 families. *Gastroenterology*, **2008**, 134, 1203-14	13.3	278
11	Reversal of tacrolimus-related hypertrophic cardiomyopathy after conversion to rapamycin in a pediatric liver transplant recipient. *Pediatric Transplantation*, **2007**, 11, 319-23	1.8	21
10	Recurrence of non-alcoholic steatohepatitis after liver transplantation in a 13-yr-old boy. *Pediatric Transplantation*, **2007**, 11, 796-8	1.8	20
9	Short-term effects of parenteral nutrition of cholestatic infants with lipid emulsions based on medium-chain and long-chain triacylglycerols. *Nutrition*, **2007**, 23, 121-6	4.8	20
8	Hepatocellular carcinoma in ten children under five years of age with bile salt export pump deficiency. *Hepatology*, **2006**, 44, 478-86	11.2	303
7	Normal pancreatic secretion in children with progressive familial intrahepatic cholestasis type 1. *Scandinavian Journal of Gastroenterology*, **2006**, 41, 1480-3	2.4	7
6	Exocrine pancreatic function in children with progressive familial intrahepatic cholestasis type 2. *Journal of Pediatric Gastroenterology and Nutrition*, **2006**, 42, 416-8	2.8	14
5	Characterization of mutations in ATP8B1 associated with hereditary cholestasis. *Hepatology*, **2004**, 40, 27-38	11.2	221
4	Long-chain PUFA supplementation improves PUFA profile in infants with cholestasis. *Lipids*, **2002**, 37, 953-7	1.6	5
3	Deficiency of the expression of CD45RA isoform of CD45 common leukocyte antigen in CD4+ T lymphocytes in children with infantile cholestasis. *Immunology Letters*, **2001**, 75, 179-84	4.1	13
2	Treatment of progressive familial intrahepatic cholestasis: liver transplantation or partial external biliary diversion. *Pediatric Transplantation*, **1999**, 3, 219-24	1.8	84
1	Fine-resolution mapping by haplotype evaluation: the examples of PFIC1 and BRIC. *Human Genetics*, **1999**, 104, 241-8	6.3	29