Irena Jankowska

List of Publications by Year
in descending order

65	1,887	430442	253896
papers	citations	18 h-index	43 g-index

71	71	71	1807
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Hepatocellular carcinoma in ten children under five years of age with bile salt export pump deficiency. Hepatology, 2006, 44, 478-486.	3.6	345
2	Severe Bile Salt Export Pump Deficiency: 82 Different ABCB11 Mutations in 109 Families. Gastroenterology, 2008, 134, 1203-1214.e8.	0.6	331
3	Characterization of mutations inATP8B1associated with hereditary cholestasis. Hepatology, 2004, 40, 27-38.	3.6	263
4	Differences in presentation and progression between severe FIC1 and BSEP deficiencies. Journal of Hepatology, 2010, 53, 170-178.	1.8	182
5	Treatment of progressive familial intrahepatic cholestasis: Liver transplantation or partial external biliary diversion. Pediatric Transplantation, 1999, 3, 219-224.	0.5	90
6	Genotype correlates with the natural history of severe bile salt export pump deficiency. Journal of Hepatology, 2020, 73, 84-93.	1.8	61
7	Outcomes of surgical management of familial intrahepatic cholestasis 1 and bile salt export protein deficiencies. Hepatology Communications, 2018, 2, 515-528.	2.0	54
8	Prevalence of Cryptosporidium, Blastocystis, and other opportunistic infections in patients with primary and acquired immunodeficiency. Parasitology Research, 2018, 117, 2869-2879.	0.6	37
9	Fine-resolution mapping by haplotype evaluation: the examples of PFIC1 and BRIC. Human Genetics, 1999, 104, 241-248.	1.8	33
10	Ileal Exclusion in Children With Progressive Familial Intrahepatic Cholestasis. Journal of Pediatric Gastroenterology and Nutrition, 2014, 58, 92-95.	0.9	27
11	Maralixibat for the treatment of PFIC: Longâ€term, IBAT inhibition in an openâ€label, Phase 2 study. Hepatology Communications, 2022, 6, 2379-2390.	2.0	26
12	Impact of Genotype, Serum Bile Acids, and Surgical Biliary Diversion on Native Liver Survival in FIC1 Deficiency. Hepatology, 2021, 74, 892-906.	3.6	25
13	Reversal of tacrolimus-related hypertrophic cardiomyopathy after conversion to rapamycin in a pediatric liver transplant recipient. Pediatric Transplantation, 2007, 11, 319-323.	0.5	24
14	Recurrence of nonâ€alcoholic steatohepatitis after liver transplantation in a 13â€yrâ€old boy. Pediatric Transplantation, 2007, 11, 796-798.	0.5	24
15	Immunological Factors and Liver Fibrosis in Pediatric Liver Transplant Recipients. Annals of Transplantation, 2015, 20, 279-284.	0.5	23
16	Short-term effects of parenteral nutrition of cholestatic infants with lipid emulsions based on medium-chain and long-chain triacylglycerols. Nutrition, 2007, 23, 121-126.	1.1	21
17	Exocrine Pancreatic Function in Children with Progressive Familial Intrahepatic Cholestasis Type 2. Journal of Pediatric Gastroenterology and Nutrition, 2006, 42, 416-418.	0.9	20
18	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.	0.9	19

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19	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, 179-184.	1.1	18
20	Cholestasis Due to USP53 Deficiency. Journal of Pediatric Gastroenterology and Nutrition, 2021, 72, 667-673.	0.9	18
21	Hemodynamic failure as an indication to urgent liver transplantation in infants with giant hepatic hemangiomas or vascular malformations â€“ Report of four cases. Pediatric Transplantation, 2009, 13, 906-912.	0.5	14
22	Progressive familial intrahepatic cholestasis and inborn errors of bile acid synthesis. Clinics and Research in Hepatology and Gastroenterology, 2012, 36, 271-274.	0.7	14
23	Acute liver failure in childrenâ€”Is living donor liver transplantation justified?. PLoS ONE, 2018, 13, e0193327.	1.1	14
24	Acute liver failure due to DGUOK deficiencyâ€“is liver transplantation justified?. Clinics and Research in Hepatology and Gastroenterology, 2021, 45, 101408.	0.7	13
25	Antibody-mediated rejection in pediatric liver transplant recipients. Annals of Transplantation, 2014, 19, 119-123.	0.5	13
26	Long-Term Systematic Monitoring of Four Polish Transaldolase Deficient Patients. JIMD Reports, 2018, 42, 79-87.	0.7	11
27	Normal pancreatic secretion in children with progressive familial intrahepatic cholestasis type 1. Scandinavian Journal of Gastroenterology, 2006, 41, 1480-1483.	0.6	10
28	The impact of cytokine gene polymorphisms on Epsteinâ€“Barr virus infection outcome in pediatric liver transplant recipients. Journal of Clinical Virology, 2012, 55, 226-232.	1.6	10
29	Successful sirolimus rescue in tacrolimusâ€induced thrombotic microangiopathy after livingâ€related liver transplantation. Pediatric Transplantation, 2012, 16, E261-4.	0.5	10
30	Treatment of pruritus with <scp>P</scp>rometheus dialysis and absorption system in a patient with benign recurrent intrahepatic cholestasis. Hepatology Research, 2014, 44, E304-E308.	1.8	10
31	The first case of Enterocytozoon bieneusi infection in Poland. Annals of Agricultural and Environmental Medicine, 2013, 20, 287-8.	0.5	10
32	Long-chain PUFA supplementation improves PUFA profile in infants with cholestasis. Lipids, 2002, 37, 953-957.	0.7	8
33	Transient Elastography for Detection of Liver Fibrosis in Children With Autosomal Recessive Polycystic Kidney Disease. Frontiers in Pediatrics, 2018, 6, 422.	0.9	8
34	Percutaneous Treatment of Biliary Strictures After Pediatric Liver Transplantation. Annals of Transplantation, 2018, 23, 845-851.	0.5	8
35	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.	0.3	7
36	Autosomal Recessive Polycystic Kidney Diseaseâ€”The Clinical Aspects and Diagnostic Challenges. Journal of Pediatric Genetics, 2021, 10, 001-008.	0.3	7

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37	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, 629-635.	0.5	6
38	Epsteinâ€“Barr virus gene expression and latent membrane protein 1 gene polymorphism in pediatric liver transplant recipients. Journal of Medical Virology, 2011, 83, 2182-2190.	2.5	6
39	Obesity, lipid profiles and oxidative stress in children after liver transplantation. Acta Biochimica Polonica, 2017, 64, 661-665.	0.3	6
40	Congenital hepatic fibrosis in a 9-year-old female patient â€“ a case report. Clinical and Experimental Hepatology, 2017, 3, 176-179.	0.6	6
41	NBAS deficiency due to biallelic c.2809Câ€‰>â€‰G variant presenting with recurrent acute liver failure with severe hyperammonemia, acquired microcephaly and progressive brain atrophy. Metabolic Brain Disease, 2021, 36, 2169-2172.	1.4	5
42	Diagnostic Approach in Biliary Strictures After Pediatric Liver Transplantation. Annals of Transplantation, 2017, 22, 257-264.	0.5	5
43	Lipid metabolism and oxidative stress in children after liver transplantation treated with sirolimus. Pediatric Transplantation, 2012, 16, 901-906.	0.5	4
44	Successful pregnancy after ileal exclusion in progressive familial intrahepatic cholestasis type 2. Annals of Hepatology, 2015, 14, 550-552.	0.6	4
45	Exocrine pancreatic function in children with Alagille syndrome. Scientific Reports, 2016, 6, 35229.	1.6	4
46	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.	0.6	4
47	Non-Hodgkin lymphoma after liver and kidney transplantation in children. Experience from one center. Advances in Clinical and Experimental Medicine, 2020, 29, 197-202.	0.6	4
48	Î²â€Carotene Deficiency in Cholestatic Liver Disease of Childhood Is Caused by Î²â€Carotene Malabsorption. Journal of Pediatric Gastroenterology and Nutrition, 2010, 51, 106-109.	0.9	3
49	Endoscopic treatment in biliary strictures after pediatric liver transplantation. Pediatric Transplantation, 2018, 22, e13271.	0.5	3
50	Small intestinal bacterial overgrowth in patients with progressive familial intrahepatic cholestasis. Acta Biochimica Polonica, 2014, 61, 103-7.	0.3	3
51	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, 1, 28-32.	0.6	2
52	Newborn presentation of Niemannâ€“Pick disease type C â€“ Difficulties and limitations of diagnostic methods. Pediatrics and Neonatology, 2018, 59, 317-318.	0.3	2
53	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.	0.9	2
54	Coronavirus Disease 2019-Liver Injury-Literature Review and Guidelines Based on the Recommendations of Hepatological Societies. Pediatric Gastroenterology, Hepatology and Nutrition, 2021, 24, 119.	0.4	2

#	Article	IF	Citations
55	Liver Histopathology in Late Protocol Biopsies after Pediatric Liver Transplantation. Children, 2021, 8, 671.	0.6	2
56	Successful Liver Transplantation in Two Polish Brothers with Transaldolase Deficiency. Children, 2021, 8, 746.	0.6	2
57	Wrodzone wÅ‚Ã³knienie wÄ…troby. Medical Science Review - Hepatologia, 2011, 11, 97-102.	0.0	2
58	Normal levels of serum pancreatic enzymes in patients with progressive familial intrahepatic cholestasis type 2. Acta Biochimica Polonica, 2010, 57, 573-5.	0.3	1
59	Ogniskowy rozrost guzkowy wÄ…troby â€“ opis przypadkÃ³w oraz przeglÄ…d literatury. Pediatria Polska, 2016, 91, 359-365.	0.1	0
60	Choroba czy zespÃ³Å‚ Caroliego? Od zaburzeÅ„ embriogenezy do przeszczepienia wÄ…troby. Pediatria Polska, 2016, 91, 453-457.	0.1	0
61	Congenital hepatic fibrosis associated with Gilbert's syndrome and cholelithiasis in a pediatric patient â€“ The crucial role of liver biopsy in diagnostic process. Pediatria Polska, 2017, 92, 446-449.	0.1	0
62	Co nowego w cholestazie â€“ czÄ™Å›Ä‡ 2. Cholestaza z podwyÅ¼szonÄ… aktywnoÅ›ciÄ… gamma-glutamylotranspeptydazy. Pediatria Polska, 2017, 92, 412-416.	0.1	0
63	Co nowego w cholestazie â€“ czÄ™Å›Ä‡ 3. Wrodzone zaburzenia syntezy kwasÃ³w Å¼Ã³Å‚ciowych. Pediatria Polska, 2017, 92, 567-574.	0.1	0
64	Cholestatic Syndromes in Childhood and Catch-Up Growth. , 2012, , 863-879.		0
65	Cardiovascular Risk Factors after Conversion from Cyclosporine to Tacrolimus in Children after Liver Transplantation. Annals of Transplantation, 2014, 19, 604-608.	0.5	0