

Joanna Pawlowska

List of Publications by Year in Descending Order

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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.

65
papers

2,485
citations

18
h-index

49
g-index

67
ext. papers

2,768
ext. citations

3.7
avg, IF

3.85
L-index

#	Paper	IF	Citations
65	Coronavirus Disease 2019-Liver Injury-Literature Review and Guidelines Based on the Recommendations of Hepatological Societies. <i>Pediatric Gastroenterology, Hepatology and Nutrition</i> , 2021 , 24, 119-126	2.3	1
64	Perspectives of Hospital Pharmacists Towards Biosimilar Medicines: A Survey of Polish Pharmacy Practice in General Hospitals. <i>BioDrugs</i> , 2019 , 33, 183-191	7.9	6
63	Immune Status in Children Before Liver Transplantation-A Cross-Sectional Analysis Within the ChilsSFree Multicentre Cohort Study. <i>Frontiers in Immunology</i> , 2019 , 10, 52	8.4	2
62	Pharmaceutical care in the neonatal intensive care unit: Perspectives of Polish medical and pharmacy students. <i>Currents in Pharmacy Teaching and Learning</i> , 2019 , 11, 361-372	1.5	0
61	Similarities and Differences in Allocation Policies for Pediatric Liver Transplantation Across the World. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2019 , 68, 700-705	2.8	6
60	Phase I/II Trial of Liver-derived Mesenchymal Stem Cells in Pediatric Liver-based Metabolic Disorders: A Prospective, Open Label, Multicenter, Partially Randomized, Safety Study of One Cycle of Heterologous Human Adult Liver-derived Progenitor Cells (HepaStem) in Urea Cycle Disorders and Crigler-Najjar Syndrome Patients. <i>Transplantation</i> , 2019 , 103, 1903-1915	1.8	33
59	Health-related quality of life in pre-adolescent liver transplant recipients with biliary atresia: A cross-sectional study. <i>Clinics and Research in Hepatology and Gastroenterology</i> , 2019 , 43, 427-435	2.4	7
58	Long-Term Systematic Monitoring of Four Polish Transaldolase Deficient Patients. <i>JIMD Reports</i> , 2018 , 42, 79-87	1.9	8
57	Oral mucosa lesions and gingival bleeding can indicate the progression of liver disease in children and adolescents aged two to 18 years. <i>Acta Paediatrica, International Journal of Paediatrics</i> , 2018 , 107, 886-892	3.1	1
56	Endoscopic treatment in biliary strictures after pediatric liver transplantation. <i>Pediatric Transplantation</i> , 2018 , 22, e13271	1.8	2
55	Immune monitoring after pediatric liver transplantation - the prospective ChilsSFree cohort study. <i>BMC Gastroenterology</i> , 2018 , 18, 63	3	7
54	New Insights in Genetic Cholestasis: From Molecular Mechanisms to Clinical Implications. <i>Canadian Journal of Gastroenterology and Hepatology</i> , 2018 , 2018, 2313675	2.8	41
53	Percutaneous Treatment of Biliary Strictures After Pediatric Liver Transplantation. <i>Annals of Transplantation</i> , 2018 , 23, 845-851	1.4	6
52	Pharmacist perspectives towards pharmaceutical care services in neonatal intensive care units in Australia and Poland. <i>Drugs and Therapy Perspectives</i> , 2018 , 34, 573-582	1.5	1
51	Congenital hepatic fibrosis associated with Gilbert's syndrome and cholelithiasis in a pediatric patient - The crucial role of liver biopsy in diagnostic process. <i>Pediatrica Polska</i> , 2017 , 92, 446-449	0.1	
50	Acute-on-chronic hepatitis. A case report of autoimmune hepatitis/primary sclerosing cholangitis/ulcerative colitis overlap syndrome in a 15-year-old patient. <i>Clinical and Experimental Hepatology</i> , 2017 , 3, 28-32	2.2	1
49	Āgodna nawracajāda cholestaza wewnĀrzwĀrobowa. <i>Pediatrica Polska</i> , 2017 , 92, 575-578	0.1	

48	Obesity, lipid profiles and oxidative stress in children after liver transplantation. <i>Acta Biochimica Polonica</i> , 2017 , 64, 661-665	2	5
47	Congenital hepatic fibrosis in a 9-year-old female patient - a case report. <i>Clinical and Experimental Hepatology</i> , 2017 , 3, 176-179	2.2	5
46	Diagnostic Approach in Biliary Strictures After Pediatric Liver Transplantation. <i>Annals of Transplantation</i> , 2017 , 22, 257-264	1.4	4
45	Polish Experience with Liver Transplantation and Post-Transplant Outcomes in Children with Urea Cycle Disorders. <i>Annals of Transplantation</i> , 2017 , 22, 555-562	1.4	4
44	Follow-Up of Pediatric Liver Transplant Patients After Reaching Adulthood. <i>Annals of Transplantation</i> , 2016 , 21, 644-648	1.4	4
43	The importance of nutrition for pediatric liver transplant patients. <i>Clinical and Experimental Hepatology</i> , 2016 , 2, 105-108	2.2	9
42	Influence of Partial External Biliary Diversion on the Lipid Profile in Children With Progressive Familial Intrahepatic Cholestasis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016 , 63, 598-602	2.8	6
41	Clinical and conventional pharmacy services in Polish hospitals: a national survey. <i>International Journal of Clinical Pharmacy</i> , 2016 , 38, 271-9	2.3	19
40	The importance of anti-transglutaminase IgA antibody detection in the diagnosis of celiac disease - case report of an inappropriate diagnostic approach. <i>Przegląd Gastroenterologiczny</i> , 2015 , 10, 250-3	6	0
39	Liver involvement in children with collagen vascular diseases. <i>Clinical and Experimental Hepatology</i> , 2015 , 1, 117-119	2.2	1
38	The limited prognostic value of liver histology in children with biliary atresia. <i>Annals of Hepatology</i> , 2015 , 14, 902-9	3.1	13
37	Post-Transplant Lymphoproliferative Disorder (PTLD) Manifesting in the Oral Cavity of a 13-Year-Old Liver Transplant Recipient (LTX). <i>Annals of Transplantation</i> , 2015 , 20, 478-82	1.4	7
36	Ileal exclusion in children with progressive familial intrahepatic cholestasis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2014 , 58, 92-5	2.8	19
35	Oral health and liver function in children and adolescents with cirrhosis of the liver. <i>Przegląd Gastroenterologiczny</i> , 2014 , 9, 24-31	6	4
34	Biliary atresia in children with aberrations involving chromosome 11q. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2014 , 58, e26-9	2.8	5
33	Treatment of pruritus with Prometheus dialysis and absorption system in a patient with benign recurrent intrahepatic cholestasis. <i>Hepatology Research</i> , 2014 , 44, E304-E308	5.1	9
32	Cardiovascular risk factors after conversion from cyclosporine to tacrolimus in children after liver transplantation. <i>Annals of Transplantation</i> , 2014 , 19, 604-8	1.4	
31	Successful sirolimus rescue in tacrolimus-induced thrombotic microangiopathy after living-related liver transplantation. <i>Pediatric Transplantation</i> , 2012 , 16, E261-4	1.8	6

30	Bone metabolism in cholestatic children before and after living-related liver transplantation--a long-term prospective study. <i>Journal of Clinical Densitometry</i> , 2012 , 15, 233-40	3.5	12
29	The impact of cytokine gene polymorphisms on Epstein-Barr virus infection outcome in pediatric liver transplant recipients. <i>Journal of Clinical Virology</i> , 2012 , 55, 226-32	14.5	9
28	The status of dental and jaw bones in children and adolescents after kidney and liver transplantation. <i>Annals of Transplantation</i> , 2012 , 17, 72-81	1.4	9
27	Blastomyces in pathological lesions on oral mucous membrane in children and adolescents after transplant and with kidney or liver diseases. <i>Journal of Stomatology</i> , 2012 , 65, 676-692	0.5	2
26	Cholestatic Syndromes in Childhood and Catch-Up Growth 2012 , 863-879		
25	Epstein-Barr virus gene expression and latent membrane protein 1 gene polymorphism in pediatric liver transplant recipients. <i>Journal of Medical Virology</i> , 2011 , 83, 2182-90	19.7	5
24	Current status of oxidative stress in pediatric liver transplantation. <i>Pediatric Transplantation</i> , 2010 , 14, 169-77	1.8	9
23	Long-term monitoring of Epstein-Barr virus DNA load and humoral parameter abnormalities in pediatric liver transplant recipients before development of malignancy. <i>Pediatric Transplantation</i> , 2010 , 14, 629-35	1.8	4
22	Differences in presentation and progression between severe FIC1 and BSEP deficiencies. <i>Journal of Hepatology</i> , 2010 , 53, 170-8	13.4	145
21	beta-Carotene deficiency in cholestatic liver disease of childhood is caused by beta-carotene malabsorption. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2010 , 51, 106-9	2.8	3
20	Genomic alterations in biliary atresia suggest region of potential disease susceptibility in 2q37.3. <i>American Journal of Medical Genetics, Part A</i> , 2010 , 152A, 886-95	2.5	48
19	Transaldolase deficiency in two new patients with a relative mild phenotype. <i>Molecular Genetics and Metabolism</i> , 2009 , 97, 15-7	3.7	26
18	Relation between genotype of hepatitis C virus with the course of hepatitis in children and teenagers and treatment efficiency. <i>Pediatrics Polska</i> , 2009 , 84, 402-411	0.1	
17	Severe bile salt export pump deficiency: 82 different ABCB11 mutations in 109 families. <i>Gastroenterology</i> , 2008 , 134, 1203-14	13.3	278
16	Reversal of tacrolimus-related hypertrophic cardiomyopathy after conversion to rapamycin in a pediatric liver transplant recipient. <i>Pediatric Transplantation</i> , 2007 , 11, 319-23	1.8	21
15	Aspergillosis in children after liver transplantation: Single center experience. <i>Pediatric Transplantation</i> , 2007 , 11, 868-75	1.8	12
14	Recurrence of non-alcoholic steatohepatitis after liver transplantation in a 13-yr-old boy. <i>Pediatric Transplantation</i> , 2007 , 11, 796-8	1.8	20
13	Hepatocellular carcinoma in ten children under five years of age with bile salt export pump deficiency. <i>Hepatology</i> , 2006 , 44, 478-86	11.2	303

12	Normal pancreatic secretion in children with progressive familial intrahepatic cholestasis type 1. <i>Scandinavian Journal of Gastroenterology</i> , 2006 , 41, 1480-3	2.4	7
11	Exocrine pancreatic function in children with progressive familial intrahepatic cholestasis type 2. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2006 , 42, 416-8	2.8	14
10	Successful treatment of a child with fulminant liver failure and coma caused by <i>Amanita phalloides</i> intoxication with albumin dialysis without liver transplantation. <i>Pediatric Transplantation</i> , 2004 , 8, 295-300	1.8	21
9	Characterization of mutations in ATP8B1 associated with hereditary cholestasis. <i>Hepatology</i> , 2004 , 40, 27-38	11.2	221
8	Long-chain PUFA supplementation improves PUFA profile in infants with cholestasis. <i>Lipids</i> , 2002 , 37, 953-7	1.6	5
7	Deficiency of the expression of CD45RA isoform of CD45 common leukocyte antigen in CD4+ T lymphocytes in children with infantile cholestasis. <i>Immunology Letters</i> , 2001 , 75, 179-84	4.1	13
6	Treatment of progressive familial intrahepatic cholestasis: liver transplantation or partial external biliary diversion. <i>Pediatric Transplantation</i> , 1999 , 3, 219-24	1.8	84
5	Fine-resolution mapping by haplotype evaluation: the examples of PFIC1 and BRIC. <i>Human Genetics</i> , 1999 , 104, 241-8	6.3	29
4	A gene encoding a liver-specific ABC transporter is mutated in progressive familial intrahepatic cholestasis. <i>Nature Genetics</i> , 1998 , 20, 233-8	36.3	869
3	Essential fatty acid status in children with cholestasis, in relation to serum bilirubin concentration. <i>Journal of Pediatrics</i> , 1997 , 131, 700-6	3.6	29
2	Treatment of cholestatic children with water-soluble vitamin E (alpha-tocopheryl polyethylene glycol succinate): effects on serum vitamin E, lipid peroxides, and polyunsaturated fatty acids. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 1997 , 24, 189-93	2.8	28
1	The value of quantitative measurement of HBeAg and HBsAg before interferon-alpha treatment of chronic hepatitis B in children. <i>Journal of Hepatology</i> , 1994 , 21, 1097-102	13.4	16