

Joanna Pawlowska

List of Publications by Year in descending order

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Version: 2024-02-01

66
papers

2,992
citations

361296

20
h-index

161767

54
g-index

67
all docs

67
docs citations

67
times ranked

2618
citing authors

#	ARTICLE	IF	CITATIONS
1	A gene encoding a liver-specific ABC transporter is mutated in progressive familial intrahepatic cholestasis. <i>Nature Genetics</i> , 1998, 20, 233-238.	9.4	968
2	Hepatocellular carcinoma in ten children under five years of age with bile salt export pump deficiency. <i>Hepatology</i> , 2006, 44, 478-486.	3.6	345
3	Severe Bile Salt Export Pump Deficiency: 82 Different ABCB11 Mutations in 109 Families. <i>Gastroenterology</i> , 2008, 134, 1203-1214.e8.	0.6	331
4	Characterization of mutations in ATP8B1 associated with hereditary cholestasis. <i>Hepatology</i> , 2004, 40, 27-38.	3.6	263
5	Differences in presentation and progression between severe FIC1 and BSEP deficiencies. <i>Journal of Hepatology</i> , 2010, 53, 170-178.	1.8	182
6	Treatment of progressive familial intrahepatic cholestasis: Liver transplantation or partial external biliary diversion. <i>Pediatric Transplantation</i> , 1999, 3, 219-224.	0.5	90
7	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, 886-895.	0.7	64
8	New Insights in Genetic Cholestasis: From Molecular Mechanisms to Clinical Implications. <i>Canadian Journal of Gastroenterology and Hepatology</i> , 2018, 2018, 1-12.	0.8	59
9	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.	0.5	47
10	Treatment of Cholestatic Children with Water-soluble Vitamin E (Î±-Tocopheryl Polyethylene Glycol) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 5 <i>Pediatric Gastroenterology and Nutrition</i> , 1997, 24, 189-193.	0.9	44
11	Essential fatty acid status in children with cholestasis, in relation to serum bilirubin concentration. <i>Journal of Pediatrics</i> , 1997, 131, 700-706.	0.9	38
12	Fine-resolution mapping by haplotype evaluation: the examples of PFIC1 and BRIC. <i>Human Genetics</i> , 1999, 104, 241-248.	1.8	33
13	Transaldolase deficiency in two new patients with a relative mild phenotype. <i>Molecular Genetics and Metabolism</i> , 2009, 97, 15-17.	0.5	30
14	Ileal Exclusion in Children With Progressive Familial Intrahepatic Cholestasis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2014, 58, 92-95.	0.9	27
15	Successful treatment of a child with fulminant liver failure and coma caused by Amanita phalloides intoxication with albumin dialysis without liver transplantation. <i>Pediatric Transplantation</i> , 2004, 8, 295-300.	0.5	26
16	Clinical and conventional pharmacy services in Polish hospitals: a national survey. <i>International Journal of Clinical Pharmacy</i> , 2016, 38, 271-279.	1.0	25
17	Reversal of tacrolimus-related hypertrophic cardiomyopathy after conversion to rapamycin in a pediatric liver transplant recipient. <i>Pediatric Transplantation</i> , 2007, 11, 319-323.	0.5	24
18	Recurrence of non-alcoholic steatohepatitis after liver transplantation in a 13-year-old boy. <i>Pediatric Transplantation</i> , 2007, 11, 796-798.	0.5	24

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19	The value of quantitative measurement of HBeAg and HBsAg before interferon- α treatment of chronic hepatitis B in children. <i>Journal of Hepatology</i> , 1994, 21, 1097-1102.	1.8	22
20	Exocrine Pancreatic Function in Children with Progressive Familial Intrahepatic Cholestasis Type 2. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2006, 42, 416-418.	0.9	20
21	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.	0.9	19
22	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, 179-184.	1.1	18
23	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, 233-240.	0.5	18
24	The limited prognostic value of liver histology in children with biliary atresia. <i>Annals of Hepatology</i> , 2015, 14, 902-909.	0.6	17
25	The importance of nutrition for pediatric liver transplant patients. <i>Clinical and Experimental Hepatology</i> , 2016, 3, 105-108.	0.6	15
26	Perspectives of Hospital Pharmacists Towards Biosimilar Medicines: A Survey of Polish Pharmacy Practice in General Hospitals. <i>BioDrugs</i> , 2019, 33, 183-191.	2.2	15
27	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.	0.9	15
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.	0.5	14
29	Aspergillosis in children after liver transplantation: Single center experience. <i>Pediatric Transplantation</i> , 2007, 11, 868-875.	0.5	13
30	Immune monitoring after pediatric liver transplantation — the prospective ChilSFree cohort study. <i>BMC Gastroenterology</i> , 2018, 18, 63.	0.8	12
31	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.	0.7	12
32	Current status of oxidative stress in pediatric liver transplantation. <i>Pediatric Transplantation</i> , 2010, 14, 169-177.	0.5	11
33	Long-Term Systematic Monitoring of Four Polish Transaldolase Deficient Patients. <i>JIMD Reports</i> , 2018, 42, 79-87.	0.7	11
34	Normal pancreatic secretion in children with progressive familial intrahepatic cholestasis type 1. <i>Scandinavian Journal of Gastroenterology</i> , 2006, 41, 1480-1483.	0.6	10
35	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, 226-232.	1.6	10
36	Successful sirolimus rescue in tacrolimus-induced thrombotic microangiopathy after living-related liver transplantation. <i>Pediatric Transplantation</i> , 2012, 16, E261-4.	0.5	10

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37	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.	1.8	10
38	Long-chain PUFA supplementation improves PUFA profile in infants with cholestasis. <i>Lipids</i> , 2002, 37, 953-957.	0.7	8
39	Percutaneous Treatment of Biliary Strictures After Pediatric Liver Transplantation. <i>Annals of Transplantation</i> , 2018, 23, 845-851.	0.5	8
40	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, 478-482.	0.5	7
41	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, 629-635.	0.5	6
42	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, 2182-2190.	2.5	6
43	Obesity, lipid profiles and oxidative stress in children after liver transplantation. <i>Acta Biochimica Polonica</i> , 2017, 64, 661-665.	0.3	6
44	Congenital hepatic fibrosis in a 9-year-old female patient – a case report. <i>Clinical and Experimental Hepatology</i> , 2017, 3, 176-179.	0.6	6
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.	0.5	6
46	Oral health and liver function in children and adolescents with cirrhosis of the liver. <i>Przegląd Gastroenterologiczny</i> , 2014, 1, 24-31.	0.3	5
47	Biliary Atresia in Children With Aberrations Involving Chromosome 11q. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2014, 58, e26-9.	0.9	5
48	Immune Status in Children Before Liver Transplantation – A Cross-Sectional Analysis Within the ChildSAFE Multicentre Cohort Study. <i>Frontiers in Immunology</i> , 2019, 10, 52.	2.2	5
49	Diagnostic Approach in Biliary Strictures After Pediatric Liver Transplantation. <i>Annals of Transplantation</i> , 2017, 22, 257-264.	0.5	5
50	Follow-Up of Pediatric Liver Transplant Patients After Reaching Adulthood. <i>Annals of Transplantation</i> , 2016, 21, 644-648.	0.5	5
51	β-Carotene Deficiency in Cholestatic Liver Disease of Childhood Is Caused by β-Carotene Malabsorption. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2010, 51, 106-109.	0.9	3
52	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.	0.7	3
53	Endoscopic treatment in biliary strictures after pediatric liver transplantation. <i>Pediatric Transplantation</i> , 2018, 22, e13271.	0.5	3
54	Liver involvement in children with collagen vascular diseases. <i>Clinical and Experimental Hepatology</i> , 2015, 3, 117-119.	0.6	2

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55	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, 1, 28-32.	0.6	2
56	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.	0.4	2
57	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.	0.4	2
58	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.	0.1	2
59	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, 4, 250-253.	0.3	1
60	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.	0.3	1
61	Blastocystis infection in a 5-year-old boy – a case report. <i>Pediatrics I Medycyna Rodzinna</i> , 2018, 14, 324-326.	2.3	1
62	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	0
63	Congenital hepatic fibrosis associated with Gilbert's syndrome and cholelithiasis in a pediatric patient – The crucial role of liver biopsy in diagnostic process. <i>Pediatrics Polska</i> , 2017, 92, 446-449.	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. <i>Annals of Transplantation</i> , 2014, 19, 604-608.	0.5	0
66	7 ESSENTIAL FATTY ACID PROFILE IN THE EARLY CHOLESTASIS OF INFANCY. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 1996, 22, 411.	0.9	0