Elizabeth B Rand

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2956738/publications.pdf

Version: 2024-02-01

39 papers 3,065 citations

394421 19 h-index 395702 33 g-index

40 all docs

40 docs citations

40 times ranked

3398 citing authors

#	Article	IF	CITATIONS
1	Alagille syndrome is caused by mutations in human Jagged1, which encodes a ligand for Notch1. Nature Genetics, 1997, 16, 243-251.	21.4	1,184
2	The Precarious State of the Liver After a Fontan Operation: Summary of a Multidisciplinary Symposium. Pediatric Cardiology, 2012, 33, 1001-1012.	1.3	262
3	Hepatic Fibrosis Is Universal Following Fontan Operation, and Severity is Associated With Time From Surgery: A Liver Biopsy and Hemodynamic Study. Journal of the American Heart Association, 2017, 6, .	3.7	195
4	Treatment of Neonatal Hemochromatosis with Exchange Transfusion and Intravenous Immunoglobulin. Journal of Pediatrics, 2009, 155, 566-571.e1.	1.8	153
5	Liver transplantation for erythropoietic protoporphyria liver disease. Liver Transplantation, 2005, 11, 1590-1596.	2.4	136
6	Evidence of Chronic Allograft Injury in Liver Biopsies From Long-term Pediatric Recipients of Liver Transplants. Gastroenterology, 2018, 155, 1838-1851.e7.	1.3	125
7	Total Serum Bilirubin within 3ÂMonths of Hepatoportoenterostomy Predicts Short-Term Outcomes in Biliary Atresia. Journal of Pediatrics, 2016, 170, 211-217.e2.	1.8	100
8	Measles vaccination after orthotopic liver transplantation. Journal of Pediatrics, 1993, 123, 87-89.	1.8	98
9	Prevalence and characterization of fibrosis in surveillance liver biopsies of patients with Fontan circulation. Human Pathology, 2016, 57, 106-115.	2.0	86
10	Alagille syndrome mutation update: Comprehensive overview of <i>JAG1</i> nd <i>NOTCH2</i> mutation frequencies and insight into missense variant classification. Human Mutation, 2019, 40, 2197-2220.	2.5	84
11	Hepatic jagged1 expression studies. Hepatology, 1999, 30, 1269-1275.	7. 3	79
12	End-organ consequences of the Fontan operation: liver fibrosis, protein-losing enteropathy and plastic bronchitis. Cardiology in the Young, 2013, 23, 831-840.	0.8	79
13	Sequential Liver and Bone Marrow Transplantation for Treatment of Erythropoietic Protoporphyria. Pediatrics, 2006, 118, e1896-e1899.	2.1	76
14	Deletions of 20p12 in Alagille syndrome: Frequency and molecular characterization., 1997, 70, 80-86.		71
15	Cost-effectiveness of Universal Hepatitis C Virus Screening of Pregnant Women in the United States. Clinical Infectious Diseases, 2019, 69, 1888-1895.	5.8	61
16	Efficacy and Safety of Immunosuppression Withdrawal in Pediatric Liver Transplant Recipients: Moving Toward Personalized Management. Hepatology, 2021, 73, 1985-2004.	7.3	57
17	Central venulitis in pediatric liver allografts. Hepatology, 2001, 33, 1141-1147.	7.3	51
18	Hepatic Abnormalities Are Present Before and Early After the Fontan Operation. Annals of Thoracic Surgery, 2015, 100, 2298-2304.	1.3	36

#	Article	IF	CITATIONS
19	Overview of pediatric liver transplantation. Gastroenterology Clinics of North America, 2003, 32, 913-929.	2.2	29
20	Protein Losing Enteropathy After Fontan Operation: Glimpses of Clarity Through the Lifting Fog. World Journal for Pediatric & Congenital Heart Surgery, 2020, 11, 92-96.	0.8	26
21	A Path FORWARD: Development of a Comprehensive Multidisciplinary Clinic to Create Health and Wellness for the Child and Adolescent with a Fontan Circulation. Pediatric Cardiology, 2022, 43, 1175-1192.	1.3	9
22	Steroid Free Treatment of Autoimmune Hepatitis in Selected Children. Journal of Pediatrics, 2019, 207, 244-247.	1.8	8
23	Live virus vaccination following pediatric liver transplantation: Outcomes from two academic children's hospitals. American Journal of Transplantation, 2022, 22, 1201-1212.	4.7	8
24	Growth in Children with a Fontan Circulation. Journal of Pediatrics, 2021, 235, 149-155.e2.	1.8	7
25	Hepatic Involvement in Aicardi-Goutià res Syndrome. Neuropediatrics, 2021, 52, 441-447.	0.6	6
26	Multiple organ involvement and ICU considerations for the care of acute liver failure (ALF) and acute on chronic liver failure (ACLF) in children. Translational Pediatrics, 2021, 10, 2749-2762.	1.2	6
27	Early detection of SARSâ€CoVâ€2 and other infections in solid organ transplant recipients and household members using wearable devices. Transplant International, 2021, 34, 1019-1031.	1.6	6
28	Defining the role of liver biopsy in the assessment of liver fibrosis in patients with Fontan circulationâ€"reply. Human Pathology, 2017, 69, 141.	2.0	4
29	Splenic Rupture in Children With Portal Hypertension. Journal of Pediatric Gastroenterology and Nutrition, 2018, 66, 447-450.	1.8	4
30	Biliary Excretion Noted on Hepatobiliary Iminodiacetic Acid Scan Does Not Exclude Diagnosis of Biliary Atresia. Journal of Pediatrics, 2020, 220, 245-248.	1.8	4
31	Center Variability in Acute Rejection and Biliary Complications After Pediatric Liver Transplantation. Liver Transplantation, 2022, 28, 454-465.	2.4	4
32	Status 1B designation does not adequately prioritize children with acuteâ€onâ€chronic liver failure for liver transplantation. Liver Transplantation, 2022, 28, 1288-1298.	2.4	4
33	Evaluation of the Candidate. , 0, , 207-212.		3
34	Inborn error of metabolism patients after liver transplantation: Outcomes of 35 patients over 27 years in one pediatric quaternary hospital. American Journal of Medical Genetics, Part A, 2022, 188, 1443-1447.	1.2	2
35	Liver disease in the survivor with congenital heart disease. Clinical Liver Disease, 2014, 4, 77-79.	2.1	1
36	Cirrhotic Cardiomyopathy in Children With Biliary Atresia: A New Objective Parameter to Predict Morbidity and Mortality on the Wait List—and Beyond!. Hepatology, 2019, 69, 940-942.	7.3	0

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37	Reply. Journal of Pediatrics, 2019, 210, 243-244.	1.8	O
38	Back Cover, Volume 40, Issue 12. Human Mutation, 2019, 40, iii.	2.5	0
39	1396. Live Virus Vaccination Following Pediatric Liver Transplantation: Results from Two Academic Children's Hospitals. Open Forum Infectious Diseases, 2020, 7, S707-S708.	0.9	O