Nedim Hadzic

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.

130
papers5,026
citations41
h-index69
g-index150
ext. papers5,923
ext. citations4.6
avg, IF5.22
L-index

#	Paper	IF	Citations
130	Liver Disease in Primary Immunodeficiencies 2022 , 851-854		
129	Autoimmune Liver Disease. <i>In Clinical Practice</i> , 2022 , 129-141	O	
128	Alpha-1 Antitrypsin Deficiency in the Transition Period. <i>In Clinical Practice</i> , 2022 , 99-106	О	
127	Biliary Atresia: Clinical Phenotypes and Aetiological Heterogeneity. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	3
126	Liver Disease in Immunodeficiencies in Children 2021 , 383-400		
125	Tumors of the Liver in Children 2021 , 769-800		
124	Human Leukocyte Antigen Profile Predicts Severity of Autoimmune Liver Disease in Children of European Ancestry. <i>Hepatology</i> , 2021 , 74, 2032-2046	11.2	5
123	Liver histology in children with glycogen storage disorders type VI and IX. <i>Digestive and Liver Disease</i> , 2021 , 53, 86-93	3.3	3
122	Clinical management of sickle cell liver disease in children and young adults. <i>Archives of Disease in Childhood</i> , 2021 , 106, 315-320	2.2	4
121	Fatal metabolic stroke in a child with propionic acidemia 11 years post liver transplant. <i>American Journal of Transplantation</i> , 2021 , 21, 1637-1640	8.7	1
120	Successful Auxiliary Liver Transplant Followed by Hematopoietic Stem Cell Transplantation in X-Linked Lymphoproliferative Disease Type 1. <i>Liver Transplantation</i> , 2021 , 27, 450-455	4.5	1
119	Biliary Atresia and Neonatal Disorders of the Bile Ducts 2021 , 757-768.e5		
118	Sclerosing Cholangitis in Pediatric Inflammatory Bowel Disease: Early Diagnosis and Management Affect Clinical Outcome. <i>Journal of Pediatrics</i> , 2021 , 238, 50-56.e3	3.6	2
117	Reply. Journal of Pediatrics, 2021,	3.6	
116	Liver and Hematopoietic Stem Cell Transplantation 2021 , 80-86		
115	Seamless Management of Juvenile Autoimmune Liver Disease: Long-Term Medical and Social Outcome. <i>Journal of Pediatrics</i> , 2020 , 218, 121-129.e3	3.6	16
114	Hepatoblastoma surveillance in infants born with very low birth weight: Has the time come?. <i>Journal of Pediatrics</i> , 2020 , 216, 248-249	3.6	4

(2018-2020)

113	Liver Transplantation in Children With Propionic Acidemia: Medium-Term Outcomes. <i>Liver Transplantation</i> , 2020 , 26, 419-430	4.5	9
112	Defining clinical subgroups and genotype-phenotype correlations in NBAS-associated disease across 110 patients. <i>Genetics in Medicine</i> , 2020 , 22, 610-621	8.1	22
111	Endoscopic Retrograde Cholangiopancreatography in Infants: Availability Under Threat: A Survey on Availability, Need, and Clinical Practice in Europe and Israel. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2020 , 71, e54-e58	2.8	3
110	Adjuvant therapy of cytomegalovirus IgM + ve associated biliary atresia: Prima facie evidence of effect. <i>Journal of Pediatric Surgery</i> , 2019 , 54, 1941-1945	2.6	16
109	Inherited Metabolic Disorders 2019 , 271-278		
108	Chronic Cholangiopathy Associated with Primary Immune Deficiencies Can Be Resolved by Effective Hematopoietic Stem Cell Transplantation. <i>Journal of Pediatrics</i> , 2019 , 209, 97-106.e2	3.6	8
107	Prognostic markers at adolescence in patients requiring liver transplantation for biliary atresia in adulthood. <i>Journal of Hepatology</i> , 2019 , 71, 71-77	13.4	26
106	Letter to the Editor: Is Nodular Regenerative Hyperplasia an Immune-Mediated Phenomenon?. <i>Hepatology</i> , 2019 , 69, 2717-2718	11.2	3
105	Combined liver and hematopoietic stem cell transplantation in patients with X-linked hyper-IgM syndrome. <i>Journal of Allergy and Clinical Immunology</i> , 2019 , 143, 1952-1956.e6	11.5	9
104	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
103	Young People With Biliary Atresia Requiring Liver Transplantation: A Distinct Population Requiring Specialist Care. <i>Transplantation</i> , 2019 , 103, e99-e107	1.8	14
102	Autoimmune pancreatitis in children: A single centre experience in diagnosis, management and long term follow up. <i>Pancreatology</i> , 2019 , 19, 169-176	3.8	6
101	Treatment of Chronic Hepatitis C Virus Infection in Children: A Position Paper by the Hepatology Committee of European Society of Paediatric Gastroenterology, Hepatology and Nutrition. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2018 , 66, 505-515	2.8	66
100	Chapter 6. ESPGHAN B Contributions to Paediatric Hepatology. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2018 , 66 Suppl 1, S116	2.8	
99	The Health Care Transition of Youth With Liver Disease Into the Adult Health System: Position Paper From ESPGHAN and EASL. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2018 , 66, 976-990	2.8	21
98	Wilsonß Disease in Children: A Position Paper by the Hepatology Committee of the European Society for Paediatric Gastroenterology, Hepatology and Nutrition. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2018 , 66, 334-344	2.8	98
97	Liver transplantation for neonatal-onset citrullinemia. <i>Pediatric Transplantation</i> , 2018 , 22, e13191	1.8	6
96	Fibropolycystic Liver Diseases and Congenital Biliary Abnormalities 2018 , 308-327		1

95	Survival of children after liver transplantation for hepatocellular carcinoma. <i>Liver Transplantation</i> , 2018 , 24, 246-255	4.5	28
94	Diagnosis and Management of Pediatric Autoimmune Liver Disease: ESPGHAN Hepatology Committee Position Statement. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2018 , 66, 345-360	2.8	136
93	Reduced Hepatocellular Expression of Canalicular Transport Proteins in Infants with Neonatal Cholestasis and Congenital Hypopituitarism. <i>Journal of Pediatrics</i> , 2018 , 200, 181-187	3.6	8
92	Early and Late Factors Impacting Patient and Graft Outcome in Pediatric Liver Transplantation: Summary of an ESPGHAN Monothematic Conference. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2017 , 65, e53-e59	2.8	16
91	Attempt to Determine the Prevalence of Two Inborn Errors of Primary Bile Acid Synthesis: Results of a European Survey. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2017 , 64, 864-868	2.8	14
90	Managing pre-symptomatic Wilson disease in genetic era - More questions than answers. <i>Clinics and Research in Hepatology and Gastroenterology</i> , 2017 , 41, 626-628	2.4	2
89	Guideline for the Evaluation of Cholestatic Jaundice in Infants: Joint Recommendations of the North American Society for Pediatric Gastroenterology, Hepatology, and Nutrition and the European Society for Pediatric Gastroenterology, Hepatology, and Nutrition. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2017 , 64, 154-168	2.8	195
88	Autoimmune Liver Disease in Children with Sickle Cell Disease. <i>Journal of Pediatrics</i> , 2017 , 189, 79-85.e	23.6	16
87	Long-term challenges and perspectives of pre-adolescent liver disease. <i>The Lancet Gastroenterology and Hepatology</i> , 2017 , 2, 435-445	18.8	10
86	Therapeutic Options in Alpha-1 Antitrypsin Deficiency: Liver Transplantation. <i>Methods in Molecular Biology</i> , 2017 , 1639, 263-265	1.4	3
85	Ruxolitinib treatment in an infant with JAK2+ polycythaemia vera-associated Budd-Chiari syndrome. <i>BMJ Case Reports</i> , 2017 , 2017,	0.9	7
84	Liver Disease in Primary Immunodeficiencies 2016 , 701-704		
83	Vitamin D Levels in Infants With Biliary Atresia: Pre- and Post-Kasai Portoenterostomy. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016 , 62, 746-50	2.8	16
82	The Changing Spectrum of Neonatal Hepatitis. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016 , 63, 316-9	2.8	5
81	Congenital Vascular Malformations of the Liver: An Association With Trisomy 21. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016 , 63, e141-e146	2.8	9
80	Hepatitis E in Children: A Position Paper by the ESPGHAN Hepatology Committee. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2016 , 63, 288-94	2.8	18
79	Th-17 cells infiltrate the liver in human biliary atresia and are related to surgical outcome. <i>Journal of Pediatric Surgery</i> , 2015 , 50, 1297-303	2.6	44
78	Cytomegalovirus-associated biliary atresia: An aetiological and prognostic subgroup. <i>Journal of Pediatric Surgery</i> , 2015 , 50, 1739-45	2.6	95

(2012-2015)

77	Inherited metabolic disorders presenting as acute liver failure in newborns and young children: King & College Hospital experience. <i>European Journal of Pediatrics</i> , 2015 , 174, 1387-92	4.1	38
76	Successful use of long-acting octreotide for intractable chronic gastrointestinal bleeding in children. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2015 , 60, 48-53	2.8	7
75	Indications and limitations of bariatric intervention in severely obese children and adolescents with and without nonalcoholic steatohepatitis: ESPGHAN Hepatology Committee Position Statement. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2015 , 60, 550-61	2.8	69
74	Liver biopsy in children: position paper of the ESPGHAN Hepatology Committee. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2015 , 60, 408-20	2.8	119
73	Autoimmune liver disease: novelties in management. <i>Clinics and Research in Hepatology and Gastroenterology</i> , 2014 , 38, 273-6	2.4	7
72	Dedicator of cytokinesis 8 deficiency: a predisposition to sclerosing cholangitis. <i>Clinical Immunology</i> , 2014 , 155, 71-73	9	12
71	Hepatic neoplasms in children: a focus on differential diagnosis. <i>Clinics and Research in Hepatology and Gastroenterology</i> , 2014 , 38, 399-402	2.4	7
70	Clinical, biochemical, cellular and molecular characterization of mitochondrial DNA depletion syndrome due to novel mutations in the MPV17 gene. <i>European Journal of Human Genetics</i> , 2014 , 22, 184-91	5.3	43
69	Childhood hepatocellular carcinoma: a clinicopathological study of 12 cases with special reference to EpCAM. <i>Histopathology</i> , 2014 , 64, 671-82	7.3	26
68	Steroids in acute liver failure?. Journal of Pediatrics, 2014, 165, 211-2	3.6	
68 67	Steroids in acute liver failure?. <i>Journal of Pediatrics</i> , 2014 , 165, 211-2 Liver transplantation for pediatric metabolic disease. <i>Molecular Genetics and Metabolism</i> , 2014 , 111, 418		82
			82
67	Liver transplantation for pediatric metabolic disease. <i>Molecular Genetics and Metabolism</i> , 2014 , 111, 418	8 <i>-92-7</i>	
67 66	Liver transplantation for pediatric metabolic disease. <i>Molecular Genetics and Metabolism</i> , 2014 , 111, 418 A genetic study of Wilson® disease in the United Kingdom. <i>Brain</i> , 2013 , 136, 1476-87 Steroids in biliary atresia: single surgeon, single centre, prospective study. <i>Journal of Hepatology</i> ,	8 <i>3</i> 27 11.2	219
67 66 65	Liver transplantation for pediatric metabolic disease. <i>Molecular Genetics and Metabolism</i> , 2014 , 111, 418 A genetic study of Wilsonß disease in the United Kingdom. <i>Brain</i> , 2013 , 136, 1476-87 Steroids in biliary atresia: single surgeon, single centre, prospective study. <i>Journal of Hepatology</i> , 2013 , 59, 1054-8 BCSH/BSBMT guideline: diagnosis and management of veno-occlusive disease (sinusoidal obstruction syndrome) following haematopoietic stem cell transplantation. <i>British Journal of</i>	11.2	219
67 66 65	Liver transplantation for pediatric metabolic disease. <i>Molecular Genetics and Metabolism</i> , 2014 , 111, 418 A genetic study of Wilsonß disease in the United Kingdom. <i>Brain</i> , 2013 , 136, 1476-87 Steroids in biliary atresia: single surgeon, single centre, prospective study. <i>Journal of Hepatology</i> , 2013 , 59, 1054-8 BCSH/BSBMT guideline: diagnosis and management of veno-occlusive disease (sinusoidal obstruction syndrome) following haematopoietic stem cell transplantation. <i>British Journal of Haematology</i> , 2013 , 163, 444-57 Diagnosis and management of acute graft-versus-host disease. <i>British Journal of Haematology</i> ,	11.2 13.4 4.5	21953200
67 66 65 64 63	Liver transplantation for pediatric metabolic disease. <i>Molecular Genetics and Metabolism</i> , 2014 , 111, 418 A genetic study of Wilson® disease in the United Kingdom. <i>Brain</i> , 2013 , 136, 1476-87 Steroids in biliary atresia: single surgeon, single centre, prospective study. <i>Journal of Hepatology</i> , 2013 , 59, 1054-8 BCSH/BSBMT guideline: diagnosis and management of veno-occlusive disease (sinusoidal obstruction syndrome) following haematopoietic stem cell transplantation. <i>British Journal of Haematology</i> , 2013 , 163, 444-57 Diagnosis and management of acute graft-versus-host disease. <i>British Journal of Haematology</i> , 2012 , 158, 30-45 Organ-specific management and supportive care in chronic graft-versus-host disease. <i>British</i>	11.2 13.4 4.5	21953200232

59	Immune Deficiency-Related Liver Disease. Pediatric and Adolescent Medicine, 2012, 64-75	0.4	
58	Medical management of the RailingRKasai portoenterostomy. South African Medical Journal, 2012 , 102, 868-71	1.5	4
57	Long-term outcome for immune suppression and immune related lymphoproliferative disorder: prospective data from the United Kingdom Children Leukaemia and Cancer Group registry 1994-2004. Leukemia and Lymphoma, 2012, 53, 842-8	1.9	6
56	Diagnosis in bile acid-CoA: amino acid N-acyltransferase deficiency. <i>World Journal of Gastroenterology</i> , 2012 , 18, 3322-6	5.6	18
55	Liver neoplasia in children. <i>Clinics in Liver Disease</i> , 2011 , 15, 443-62, vii-x	4.6	49
54	Biliary atresia in England and Wales: results of centralization and new benchmark. <i>Journal of Pediatric Surgery</i> , 2011 , 46, 1689-94	2.6	169
53	Poststreptococcal acute glomerulonephritis and dense deposit disease after pediatric liver transplantation. <i>Transplantation</i> , 2011 , 91, e44-6	1.8	
52	Evolving changes in aetiology of gallstones in children. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2011 , 52, 499	2.8	
51	Cysts and Congenital Biliary Abnormalities 2011 , 312-328		
50	Hepatocellular carcinoma in biliary atresia: Kingß College Hospital experience. <i>Journal of Pediatrics</i> , 2011 , 159, 617-22.e1	3.6	56
49	Liver transplantation for propionic acidemia in children. Liver Transplantation, 2011, 17, 661-7	4.5	63
48	Selective rather than routine approach to endosopic retrograde cholangio-pancreatography in diagnosis of biliary atresia. <i>Journal of Hepatology</i> , 2010 , 52, 777	13.4	3
47	Variable clinical spectrum of the most common inborn error of bile acid metabolism3beta-hydroxy-Delta 5-C27-steroid dehydrogenase deficiency. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2010 , 50, 61-6	2.8	44
46	Alagille syndrome and liver transplantation. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2010 , 50, 11-5	2.8	73
45	A novel monoclonal antibody to characterize pathogenic polymers in liver disease associated with alpha1-antitrypsin deficiency. <i>Hepatology</i> , 2010 , 52, 1078-88	11.2	111
44	Polymorphism in the endoplasmic reticulum mannosidase I (MAN1B1) gene is not associated with liver disease in individuals homozygous for the Z variant of the alpha1-antitrypsin protease inhibitor (PiZZ individuals). <i>Hepatology</i> , 2009 , 50, 1315; author reply 1315-6	11.2	13
43	Selective use of endoscopic retrograde cholangiopancreatography in the diagnosis of biliary atresia in infants younger than 100 days. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2009 , 49, 435-41	2.8	59
42	Evolution in the management of acute liver failure-associated aplastic anaemia in children: a single centre experience. <i>Journal of Hepatology</i> , 2008 , 48, 68-73	13.4	115

(2004-2008)

41	Drug-related hepatotoxicity and acute liver failure. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2008 , 47, 395-405	2.8	90	
40	Surgical outcome in biliary atresia: etiology affects the influence of age at surgery. <i>Annals of Surgery</i> , 2008 , 247, 694-8	7.8	83	
39	A polymorphism of the alpha1-antitrypsin gene represents a risk factor for liver disease. <i>Hepatology</i> , 2008 , 47, 127-32	11.2	45	
38	Efficacy and tolerability of high-dose methotrexate in central nervous system positive or relapsed lymphoproliferative disease following liver transplant in children. <i>British Journal of Haematology</i> , 2008 , 140, 191-6	4.5	30	
37	Randomized, double-blind, placebo-controlled trial of corticosteroids after Kasai portoenterostomy for biliary atresia. <i>Hepatology</i> , 2007 , 46, 1821-7	11.2	104	
36	Hepatobiliary scintigraphy after Kasai procedure for biliary atresia: clinical correlation and prognostic value. <i>Journal of Pediatric Surgery</i> , 2007 , 42, 1107-13	2.6	7	
35	Liver mtDNA content increases during development: a comparison of methods and the importance of age- and tissue-specific controls for the diagnosis of mtDNA depletion. <i>Mitochondrion</i> , 2007 , 7, 386-9	95 ^{4.9}	47	
34	Hepatocellular carcinoma in a 12-year-old child with PiZZ alpha1-antitrypsin deficiency. <i>Hepatology</i> , 2006 , 43, 194	11.2	66	
33	Single agent efficacy of rituximab in childhood immunosuppression related lymphoproliferative disease: a United Kingdom Childrenß Cancer Study Group (UKCCSG) retrospective review. <i>Leukemia and Lymphoma</i> , 2006 , 47, 2584-9	1.9	30	
32	The biliary atresia splenic malformation syndrome: a 28-year single-center retrospective study. Journal of Pediatrics, 2006 , 149, 393-400	3.6	181	
31	Neonatal herpes simplex virus infection presenting as acute liver failure: prevalent role of herpes simplex virus type I. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2006 , 42, 282-6	2.8	46	
30	Heterogeneity of the epsilon gamma delta beta-thalassaemias: characterization of three novel English deletions. <i>British Journal of Haematology</i> , 2005 , 128, 722-9	4.5	33	
29	Outcome of PiSS and PiSZ alpha-1-antitrypsin deficiency presenting with liver involvement. <i>European Journal of Pediatrics</i> , 2005 , 164, 250-2	4.1	12	
28	Disseminated neonatal herpes simplex virus (HSV) type 2 infection diagnosed by HSV DNA detection in blood and successfully managed by liver transplantation. <i>European Journal of Pediatrics</i> , 2004 , 163, 166-9	4.1	22	
27	Hepatic veno-occlusive disease and portal vein thrombosis; closer than we think?. <i>European Journal of Cancer</i> , 2004 , 40, 2643-4	7.5		
26	Antenatal presentation of biliary atresia. <i>Journal of Pediatrics</i> , 2004 , 144, 43-6	3.6	66	
25	Steatohepatitis and unsuspected micronodular cirrhosis in Dorfman-Chanarin syndrome with documented ABHD5 mutation. <i>Journal of Pediatrics</i> , 2004 , 144, 662-5	3.6	37	
24	Liver disease in children with primary immunodeficiencies. <i>Journal of Pediatrics</i> , 2004 , 145, 333-9	3.6	67	

23	Long-term survival following Kasai portoenterostomy: is chronic liver disease inevitable?. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2003 , 37, 430-3	2.8	70
22	. Pediatric Infectious Disease Journal, 2003 , 22, 329-334	3.4	4
21	Polymerase chain reaction-based diagnosis of infection with Cryptosporidium in children with primary immunodeficiencies. <i>Pediatric Infectious Disease Journal</i> , 2003 , 22, 329-35	3.4	55
20	Humoral immunity in children with biliary atresia splenic malformation syndrome. <i>European Journal of Pediatrics</i> , 2003 , 162, 539-540	4.1	9
19	HLA and cytokine gene polymorphisms in biliary atresia. <i>Liver</i> , 2002 , 22, 213-9		41
18	Hepatic hemangioendothelioma associated with production of humoral thyrotropin-like factor. <i>Journal of Pediatrics</i> , 2001 , 138, 932-5	3.6	39
17	Mycobacterium tuberculosis infection in pediatric liver transplant recipients. <i>Pediatric Infectious Disease Journal</i> , 2000 , 19, 625-30	3.4	30
16	Response to varicella immunization in pediatric liver transplant recipients. <i>Transplantation</i> , 2000 , 70, 1401-4	1.8	39
15	Recurrence of Langerhans cell histiocytosis in the graft after pediatric liver transplantation. <i>Transplantation</i> , 2000 , 70, 815-9	1.8	44
14	Bone marrow failure in children with acute liver failure. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2000 , 31, 557-61	2.8	39
13	Nonmyeloablative stem cell transplantation for congenital immunodeficiencies. <i>Blood</i> , 2000 , 96, 1239-1	1246	128
12	Correction of the hyper-IgM syndrome after liver and bone marrow transplantation. <i>New England Journal of Medicine</i> , 2000 , 342, 320-4	59.2	90
11	Superoxide dismutase activity in children with chronic liver diseases. <i>Journal of Hepatology</i> , 2000 , 32, 188-92	13.4	21
10	Prognosis of alpha-1-antitrypsin deficiency-related liver disease in the era of paediatric liver transplantation. <i>Journal of Hepatology</i> , 2000 , 32, 986-92	13.4	78
9	Liver disease in primary immunodeficiencies. <i>Journal of Hepatology</i> , 2000 , 32, 9-10	13.4	7
8	Nonmyeloablative stem cell transplantation for congenital immunodeficiencies. <i>Blood</i> , 2000 , 96, 1239-1	1246	5
7	Symptomatic neonatal hepatitis A disease from a virus variant acquired in utero. <i>Pediatric Infectious Disease Journal</i> , 1999 , 18, 389-91	3.4	6
6	De-novo autoimmune hepatitis after liver transplantation. <i>Lancet, The</i> , 1998 , 351, 409-13	40	350

LIST OF PUBLICATIONS

5	Role of HLA compatibility in pediatric liver transplantation. <i>Transplantation</i> , 1998 , 66, 53-8	1.8	25	
4	Gender matching and outcome after pediatric liver transplantation. <i>Transplantation</i> , 1998 , 66, 602-5	1.8	26	
3	Successful orthotopic liver transplantation for fulminant liver failure in a child with autosomal recessive chronic granulomatous disease. <i>Transplantation</i> , 1995 , 60, 1185-6	1.8	2	
2	Liver disease in immunodeficiencies370-386			
1	Liver Disease in Immunodeficiencies513-528		1	