David A Piccoli

List of Publications by Year
in descending order

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67 papers	7,857 citations	81900 39 h-index	144013 57 g-index
68 all docs	68 docs citations	68 times ranked	4958 citing authors

#	Article	IF	Citations
1	Genome sequencing increases diagnostic yield in clinically diagnosed Alagille syndrome patients with previously negative test results. Genetics in Medicine, 2021, 23, 323-330.	2.4	17
2	Alagille Syndrome. , 2021, , 222-241.		2
3	Alagille syndrome and risk for hepatocellular carcinoma: Need for increased surveillance in adults with mild liver phenotypes. American Journal of Medical Genetics, Part A, 2021, 185, 719-731.	1.2	12
4	Bone geometry and microarchitecture deficits in children with Alagille syndrome. Bone, 2020, 141, 115576.	2.9	9
5	Proteinâ€elongating mutations inMYH11are implicated in a dominantly inherited smooth muscle dysmotility syndrome with severe esophageal, gastric, and intestinal disease. Human Mutation, 2020, 41, 973-982.	2.5	18
6	Outcomes of Childhood Cholestasis in Alagille Syndrome: Results of a Multicenter Observational Study. Hepatology Communications, 2020, 4, 387-398.	4.3	52
7	Alagille syndrome mutation update: Comprehensive overview ofJAG1andNOTCH2mutation frequencies and insight into missense variant classification. Human Mutation, 2019, 40, 2197-2220.	2.5	84
8	Intrahepatic dynamic contrast MR lymphangiography: initial experience with a new technique for the assessment of liver lymphatics. European Radiology, 2019, 29, 5190-5196.	4.5	51
9	Back Cover, Volume 40, Issue 12. Human Mutation, 2019, 40, iii.	2.5	0
10	Histological features of ileitis differentiating pediatric Crohn disease from ulcerative colitis with backwash ileitis. Digestive and Liver Disease, 2018, 50, 147-153.	0.9	11
11	Alagille Syndrome: Overview and Introduction. , 2018, , 1-9.		0
12	Recommendations for Diagnosis and Management of Autoimmune Pancreatitis in Childhood. Journal of Pediatric Gastroenterology and Nutrition, 2018, 67, 232-236.	1.8	35
13	Protein-Losing Enteropathy in Patients With Congenital Heart Disease. Journal of the American College of Cardiology, 2017, 69, 2929-2937.	2.8	136
14	Autoimmune Pancreatitis in Children: Characteristic Features, Diagnosis, and Management. American Journal of Gastroenterology, 2017, 112, 1604-1611.	0.4	70
15	Early life predictive markers of liver disease outcome in an International, Multicentre Cohort of children with Alagille syndrome. Liver International, 2016, 36, 755-760.	3.9	37
16	Compound heterozygous mutations in NEK8 in siblings with endâ€stage renal disease with hepatic and cardiac anomalies. American Journal of Medical Genetics, Part A, 2016, 170, 750-753.	1.2	22
17	Chromosome 10q23 Deletion Syndrome: An Overlap of Bannayanâ€“Rileyâ€“Ruvalcaba Syndrome and Juvenile Polyposis Syndrome. Journal of Paediatrics and Child Health, 2016, 52, 852-852.	0.8	3
18	Anterior Chamber Pathology in Alagille Syndrome. Ocular Oncology and Pathology, 2016, 2, 270-275.	1.0	6

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19	Severe Eosinophilic Gastroenteritis in a Crohnâ€™s Disease Patient Treated With Infliximab and Adalimumab. American Journal of Gastroenterology, 2016, 111, 437-438.	0.4	44
20	A de novo whole gene deletion of XIAP detected by exome sequencing analysis in very early onset inflammatory bowel disease: a case report. BMC Gastroenterology, 2015, 15, 160.	2.0	38
21	Moyamoya Syndrome Associated with Alagille Syndrome: Outcome after Surgical Revascularization. Journal of Pediatrics, 2015, 166, 470-473.	1.8	29
22	Alagille syndrome. , 2014, , 216-233.		10
23	NOTCH2mutations in Alagille syndrome. Journal of Medical Genetics, 2012, 49, 138-144.	3.2	197
24	Pancreatic Insufficiency Is Not a Prevalent Problem in Alagille Syndrome. Journal of Pediatric Gastroenterology and Nutrition, 2012, 55, 612-614.	1.8	20
25	Renal anomalies in Alagille syndrome: A diseaseâ€defining feature. American Journal of Medical Genetics, Part A, 2012, 158A, 85-89.	1.2	102
26	Pathologic Lower Extremity Fractures in Children With Alagille Syndrome. Journal of Pediatric Gastroenterology and Nutrition, 2010, 51, 66-70.	1.8	69
27	Medical Management of Alagille Syndrome. Journal of Pediatric Gastroenterology and Nutrition, 2010, 50, 580-586.	1.8	61
28	A Longitudinal Study to Identify Laboratory Predictors of Liver Disease Outcome in Alagille Syndrome. Journal of Pediatric Gastroenterology and Nutrition, 2010, 50, 526-530.	1.8	47
29	Alagille Syndrome and JAGGED1/NOTCH Sequence. , 2010, , 159-178.		1
30	SNP array mapping of chromosome 20p deletions: Genotypes, phenotypes, and copy number variation. Human Mutation, 2009, 30, 371-378.	2.5	61
31	Frequency and Clinical Correlations of Granulomas in Children With Crohn Disease. Journal of Pediatric Gastroenterology and Nutrition, 2008, 46, 392-398.	1.8	91
32	Alagille Syndrome. , 2008, , 227-232.		1
33	Alagille Syndrome. , 2007, , 326-345.		15
34	NOTCH2 Mutations Cause Alagille Syndrome, a Heterogeneous Disorder of the Notch Signaling Pathway. American Journal of Human Genetics, 2006, 79, 169-173.	6.2	663
35	Intracranial Vascular Abnormalities in Patients with Alagille Syndrome. Journal of Pediatric Gastroenterology and Nutrition, 2005, 41, 99-107.	1.8	101
36	Deficits in Size-Adjusted Bone Mass in Children with Alagille Syndrome. Journal of Pediatric Gastroenterology and Nutrition, 2005, 40, 76-82.	1.8	41

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37	Vascular Anomalies in Alagille Syndrome. Circulation, 2004, 109, 1354-1358.	1.6	333
38	Alagille Syndrome. , 2004, , 20-24.		0
39	Heritable disorders of the bile ducts. Gastroenterology Clinics of North America, 2003, 32, 857-875.	2.2	63
40	Inflammatory bowel disease in children 5 years of age and younger. American Journal of Gastroenterology, 2002, 97, 2005-2010.	0.4	189
41	Rethinking Growth Failure in Alagille Syndrome: The Role of Dietary Intake and Steatorrhea. Journal of Pediatric Gastroenterology and Nutrition, 2002, 35, 495-502.	1.8	36
42	Analysis of Cardiovascular Phenotype and Genotype-Phenotype Correlation in Individuals With a JAG1 Mutation and/or Alagille Syndrome. Circulation, 2002, 106, 2567-2574.	1.6	273
43	Vitamin D status in children, adolescents, and young adults with Crohn disease,,. American Journal of Clinical Nutrition, 2002, 76, 1077-1081.	4.7	172
44	Facial features in Alagille syndrome: Specific or cholestasis facies?. American Journal of Medical Genetics Part A, 2002, 112, 163-170.	2.4	101
45	Characterization of Notch receptor expression in the developing mammalian heart and liver. American Journal of Medical Genetics Part A, 2002, 112, 181-189.	2.4	113
46	Craniosynostosis in Alagille syndrome. American Journal of Medical Genetics Part A, 2002, 112, 176-180.	2.4	46
47	Monozygotic twins with a severe form of Alagille syndrome and phenotypic discordance. American Journal of Medical Genetics Part A, 2002, 112, 194-197.	2.4	49
48	Alagille syndrome inherited from a phenotypically normal mother with a mosaic 20p microdeletion. American Journal of Medical Genetics Part A, 2002, 112, 190-193.	2.4	26
49	Mutation analysis of Jagged1 (JAG1) in Alagille syndrome patients. Human Mutation, 2001, 17, 151-152.	2.5	76
50	Alagille Syndrome and the Jagged1 Gene. Seminars in Liver Disease, 2001, 21, 525-534.	3.6	127
51	Growth, Body Composition, and Nutritional Status in Children and Adolescents With Crohn's Disease. Journal of Pediatric Gastroenterology and Nutrition, 2000, 31, 33-40.	1.8	130
52	Features of alagille syndrome in 92 patients: Frequency and relation to prognosis. Hepatology, 1999, 29, 822-829.	7.3	591
53	Jagged1 mutations in patients ascertained with isolated congenital heart defects. , 1999, 84, 56-60.		137
54	INFLAMMATORY BOWEL DISEASE IN PEDIATRIC AND ADOLESCENT PATIENTS. Gastroenterology Clinics of North America, 1999, 28, 445-458.	2.2	183

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55	Risk factors for low bone mineral density in children and young adults with Crohnâ€™s disease. Journal of Pediatrics, 1999, 135, 593-600.	1.8	147
56	Growth, nutritional status, body composition, and energy expenditure in prepubertal children with Alagille syndrome. Journal of Pediatrics, 1999, 134, 172-177.	1.8	39
57	Clinical and molecular genetics of Alagille syndrome. Current Opinion in Pediatrics, 1999, 11, 558-564.	2.0	96
58	Bone Mineral Density in Children and Young Adults with CrohnÊ¼s Disease. Inflammatory Bowel Diseases, 1999, 5, 161-166.	1.9	60
59	Experiences with 6-Mercaptopurine and Azathioprine Therapy in Pediatric Patients with Severe Ulcerative Colitis. Journal of Pediatric Gastroenterology and Nutrition, 1999, 28, 54-58.	1.8	60
60	Single toxin detection is inadequate to diagnose Clostridium difficile diarrhea in pediatric patients. Gastroenterology, 1998, 115, 1329-1334.	1.3	49
61	Spectrum and Frequency of Jagged1 (JAG1) Mutations in Alagille Syndrome Patients and Their Families. American Journal of Human Genetics, 1998, 62, 1361-1369.	6.2	218
62	Mutations in the human Jagged1 gene are responsible for Alagille syndrome. Nature Genetics, 1997, 16, 235-242.	21.4	1,072
63	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
64	Deletions of 20p12 in Alagille syndrome: Frequency and molecular characterization. , 1997, 70, 80-86.		71
65	Inflammatory bowel disease in children. Medical Clinics of North America, 1994, 78, 1281-1302.	2.5	24
66	Bile canalicular morphometry in arteriohepatic dysplasia. Hepatology, 1987, 7, 1262-1266.	7.3	11
67	Bone mineral density in children and young adults with Crohn's disease. Inflammatory Bowel Diseases, 0, 5, 161-166.	1.9	21