Keith D Lindor

List of Publications by Year in descending order

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259 papers 29,464 citations

91 h-index 166 g-index

300 all docs 300 does citations

300 times ranked

11483 citing authors

#	Article	IF	CITATIONS
1	Potential Association of Doxycycline With the Onset of Primary Sclerosing Cholangitis: A Case Series. American Journal of Therapeutics, 2022, 29, e437-e443.	0.5	5
2	Primary biliary cholangitis: 2021 practice guidance update from the American Association for the Study of Liver Diseases. Hepatology, 2022, 75, 1012-1013.	3.6	34
3	Dynamic Risk Prediction of Response to Ursodeoxycholic Acid Among Patients with Primary Biliary Cholangitis in the USA. Digestive Diseases and Sciences, 2022, 67, 4170-4180.	1.1	3
4	Alan Hofmann (1931â€2021): A career well spent understanding bile acids. Hepatology, 2022, 75, 238-239.	3.6	0
5	Machine learning in primary biliary cholangitis: A novel approach for risk stratification. Liver International, 2022, 42, 615-627.	1.9	7
6	A pilot study of vidofludimus calcium for treatment of primary sclerosing cholangitis. Hepatology Communications, 2022, 6, 1589-1597.	2.0	7
7	Measurement of Gamma Glutamyl Transferase to Determine Risk of Liver Transplantation or Death in Patients With Primary Biliary Cholangitis. Clinical Gastroenterology and Hepatology, 2021, 19, 1688-1697.e14.	2.4	30
8	Successful response of primary sclerosing cholangitis and associated ulcerative colitis to oral vancomycin may depend on brand and personalized dose: report in an adolescent. Clinical Journal of Gastroenterology, 2021, 14, 684-689.	0.4	8
9	Early Cholangiocarcinoma Detection With Magnetic Resonance Imaging Versus Ultrasound in Primary Sclerosing Cholangitis. Hepatology, 2021, 73, 1868-1881.	3.6	25
10	Assessing and managing symptom burden and quality of life in primary sclerosing cholangitis patients. Expert Opinion on Orphan Drugs, 2021, 9, 53-66.	0.5	1
11	Safety of fibrates in cholestatic liver diseases. Liver International, 2021, 41, 1335-1343.	1.9	25
12	A Comparison of Prognostic Scores (Mayo, UK-PBC, and GLOBE) in Primary Biliary Cholangitis. American Journal of Gastroenterology, 2021, 116, 1514-1522.	0.2	14
13	Global incidence, prevalence and features of primary sclerosing cholangitis: A systematic review and metaâ€analysis. Liver International, 2021, 41, 2418-2426.	1.9	21
14	Liver Stiffness Measured by Either Magnetic Resonance or Transient Elastography Is Associated With Liver Fibrosis and Is an Independent Predictor of Outcomes Among Patients With Primary Biliary Cholangitis. Journal of Clinical Gastroenterology, 2021, 55, 449-457.	1.1	34
15	Factors Associated With Progression and Outcomes of Early Stage Primary Biliary Cholangitis. Clinical Gastroenterology and Hepatology, 2020, 18, 684-692.e6.	2.4	17
16	Consensus guidelines: best practices for detection, assessment and management of suspected acute drugâ€induced liver injury occurring during clinical trials in adults with chronic cholestatic liver disease. Alimentary Pharmacology and Therapeutics, 2020, 51, 90-109.	1.9	21
17	Open-label prospective therapeutic clinical trials: oral vancomycin in children and adults with primary sclerosing cholangitis. Scandinavian Journal of Gastroenterology, 2020, 55, 941-950.	0.6	31
18	The long-term outcomes of patients with immunoglobulin G4-related sclerosing cholangitis: the Mayo Clinic experience. Journal of Gastroenterology, 2020, 55, 1087-1097.	2.3	10

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19	Goals of Treatment for Improved Survival in Primary Biliary Cholangitis: Treatment Target Should Be Bilirubin Within the Normal Range and Normalization of Alkaline Phosphatase. American Journal of Gastroenterology, 2020, 115, 1066-1074.	0.2	74
20	Ursodeoxycholic Acid Treatment Preferentially Improves Overall Survival Among African Americans With Primary Biliary Cholangitis. American Journal of Gastroenterology, 2020, 115, 262-270.	0.2	14
21	Primary Biliary Cholangitis: 2018 Practice Guidance From the American Association for the Study of Liver Diseases. Clinical Liver Disease, 2020, 15, 1-2.	1.0	13
22	An update on primary sclerosing cholangitis epidemiology, outcomes and quantification of alkaline phosphatase variability in a population-based cohort. Journal of Gastroenterology, 2020, 55, 523-532.	2.3	22
23	Number needed to treat with ursodeoxycholic acid therapy to prevent liver transplantation or death in primary biliary cholangitis. Gut, 2020, 69, 1502-1509.	6.1	28
24	Simplified care-pathway selection for nonspecialist practice. European Journal of Gastroenterology and Hepatology, 2020, Publish Ahead of Print, .	0.8	2
25	Primary Biliary Cholangitis: 2018 Practice Guidance from the American Association for the Study of Liver Diseases. Hepatology, 2019, 69, 394-419.	3.6	507
26	Fibrosis stage is an independent predictor of outcome in primary biliary cholangitis despite biochemical treatment response. Alimentary Pharmacology and Therapeutics, 2019, 50, 1127-1136.	1.9	66
27	AGA Clinical Practice Update on Surveillance for Hepatobiliary Cancers in Patients With Primary Sclerosing Cholangitis: Expert Review. Clinical Gastroenterology and Hepatology, 2019, 17, 2416-2422.	2.4	60
28	Efficacy and safety of curcumin in primary sclerosing cholangitis: an open label pilot study. Scandinavian Journal of Gastroenterology, 2019, 54, 633-639.	0.6	23
29	Effects of Age and Sex of Response to Ursodeoxycholic Acid and Transplant-free Survival in Patients With Primary Biliary Cholangitis. Clinical Gastroenterology and Hepatology, 2019, 17, 2076-2084.e2.	2.4	54
30	Ursodeoxycholic acid therapy and liver transplant-free survival in patients with primary biliary cholangitis. Journal of Hepatology, 2019, 71, 357-365.	1.8	148
31	Current and promising therapy for primary biliary cholangitis. Expert Opinion on Pharmacotherapy, 2019, 20, 1161-1167.	0.9	10
32	NGM282, an FGF19 analogue, in primary sclerosing cholangitis: A nebulous matter. Journal of Hepatology, 2019, 70, 348-350.	1.8	5
33	Cancer risk, screening and surveillance in primary sclerosing cholangitis. Minerva Gastroenterologica E Dietologica, 2019, 65, 214-228.	2.2	7
34	Cancer risk in primary sclerosing cholangitis: Epidemiology, prevention, and surveillance strategies. World Journal of Gastroenterology, 2019, 25, 659-671.	1.4	75
35	Editorial: is proton pump inhibitor use associated with worse outcomes in patients with liver abscesses?. Alimentary Pharmacology and Therapeutics, 2018, 47, 1226-1227.	1.9	1
36	Design and Endpoints for Clinical Trials in Primary Sclerosing Cholangitis. Hepatology, 2018, 68, 1174-1188.	3.6	42

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37	Review article: the evidence that vancomycin is a therapeutic option for primary sclerosing cholangitis. Alimentary Pharmacology and Therapeutics, 2018, 47, 886-895.	1.9	57
38	Increasing Prevalence of Primary Biliary Cholangitis and Reduced Mortality With Treatment. Clinical Gastroenterology and Hepatology, 2018, 16, 1342-1350.e1.	2.4	73
39	Factors Associated With Prevalence and Treatment of Primary Biliary Cholangitis in United States Health Systems. Clinical Gastroenterology and Hepatology, 2018, 16, 1333-1341.e6.	2.4	42
40	Complications, symptoms, quality of life and pregnancy in cholestatic liver disease. Liver International, 2018, 38, 399-411.	1.9	30
41	Major Hepatic Complications in Ursodeoxycholic Acid-Treated Patients With Primary Biliary Cholangitis: Risk Factors and Time Trends in Incidence and Outcome. American Journal of Gastroenterology, 2018, 113, 254-264.	0.2	64
42	Surveillance for hepatobiliary cancers in patients with primary sclerosing cholangitis. Hepatology, 2018, 67, 2338-2351.	3.6	92
43	Managing PBC: Expanding the Provider Comfort Zone. Digestive Diseases and Sciences, 2018, 63, 2487-2488.	1.1	0
44	Primary sclerosing cholangitis in children versus adults: lessons for the clinic. Expert Review of Gastroenterology and Hepatology, 2018, 12, 1025-1032.	1.4	11
45	Emerging therapeutic targets for primary sclerosing cholangitis. Expert Opinion on Orphan Drugs, 2018, 6, 393-401.	0.5	О
46	Antimitochondrial Antibody–Negative Primary Biliary Cholangitis. Clinics in Liver Disease, 2018, 22, 589-601.	1.0	11
47	Dominant strictures in primary sclerosing cholangitis: A multicenter survey of clinical definitions and practices. Hepatology Communications, 2018, 2, 836-844.	2.0	28
48	Primary Sclerosing Cholangitis, Part 1: Epidemiology, Etiopathogenesis, Clinical Features, and Treatment. Gastroenterology and Hepatology, 2018, 14, 293-304.	0.2	9
49	Primary Sclerosing Cholangitis, Part 2: Cancer Risk, Prevention, and Surveillance. Gastroenterology and Hepatology, 2018, 14, 427-432.	0.2	4
50	Prospective Clinical Trial of Rifaximin Therapy for Patients With Primary Sclerosing Cholangitis. American Journal of Therapeutics, 2017, 24, e56-e63.	0.5	55
51	A Randomized, Placebo-Controlled Clinical Trial of Efficacy and Safety: Modafinil in the Treatment of Fatigue in Patients With Primary Biliary Cirrhosis. American Journal of Therapeutics, 2017, 24, e167-e176.	0.5	46
52	Update on pharmacotherapies for cholestatic liver disease. Hepatology Communications, 2017, 1, 7-17.	2.0	16
53	Patient Age, Sex, and Inflammatory Bowel Disease Phenotype Associate With Course of Primary Sclerosing Cholangitis. Gastroenterology, 2017, 152, 1975-1984.e8.	0.6	355
54	Emerging treatments for primary sclerosing cholangitis. Expert Review of Gastroenterology and Hepatology, 2017, 11, 451-459.	1.4	12

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55	Combination Therapy of All-Trans Retinoic Acid With Ursodeoxycholic Acid in Patients With Primary Sclerosing Cholangitis. Journal of Clinical Gastroenterology, 2017, 51, e11-e16.	1.1	38
56	Old and new treatments for primary biliary cholangitis. Liver International, 2017, 37, 490-499.	1.9	37
57	Clinical implications of serial versus isolated biliary fluorescence <i>in situ</i> hybridization (FISH) polysomy in primary sclerosing cholangitis. Scandinavian Journal of Gastroenterology, 2017, 52, 377-381.	0.6	26
58	An update on cancer risk and surveillance in primary sclerosing cholangitis. Liver International, 2017, 37, 1103-1109.	1.9	40
59	Heterogeneity of Outcomes Following Liver Transplantation for Primary Sclerosing Cholangitis: Age Matters. Digestive Diseases and Sciences, 2017, 62, 3210-3211.	1.1	2
60	Investigational drugs in phase II clinical trials for primary biliary cholangitis. Expert Opinion on Investigational Drugs, 2017, 26, 1115-1121.	1.9	6
61	Curcumin in Hepatobiliary Disease: Pharmacotherapeutic Properties and Emerging Potential Clinical Applications. Annals of Hepatology, 2017, 16, 835-841.	0.6	39
62	Ursodeoxycholic Acid Treatment in Primary Sclerosing Cholangitis., 2017,, 145-152.		7
63	Oral Vancomycin Therapy in a Child with Primary Sclerosing Cholangitis and Severe Ulcerative Colitis. Pediatric Gastroenterology, Hepatology and Nutrition, 2016, 19, 210.	0.4	18
64	Ustekinumab for patients with primary biliary cholangitis who have an inadequate response to ursodeoxycholic acid: A proofâ \in ofâ \in oncept study. Hepatology, 2016, 64, 189-199.	3.6	101
65	Novel treatments in primary sclerosing cholangitis. Clinical Liver Disease, 2016, 8, 132-135.	1.0	1
66	The Microbiome and Primary Sclerosing Cholangitis. Seminars in Liver Disease, 2016, 36, 340-348.	1.8	15
67	Targets and investigative treatments for primary biliary cholangitis. Expert Opinion on Orphan Drugs, 2016, 4, 1011-1020.	0.5	0
68	Advances in primary sclerosing cholangitis. The Lancet Gastroenterology and Hepatology, 2016, 1, 68-77.	3.7	18
69	Obeticholic acid for the treatment of primary biliary cholangitis. Expert Opinion on Pharmacotherapy, 2016, 17, 1809-1815.	0.9	39
70	A Placebo-Controlled Trial of Obeticholic Acid in Primary Biliary Cholangitis. New England Journal of Medicine, 2016, 375, 631-643.	13.9	817
71	Distinguishing immunoglobulin G4–related disease from its pancreatobiliary mimics: Are we there now?. Hepatology, 2016, 64, 340-343.	3.6	11
72	Stratification of hepatocellular carcinoma risk in primary biliary cirrhosis: a multicentre international study. Gut, 2016, 65, 321-329.	6.1	139

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73	Long-term outcomes in antimitochondrial antibody negative primary biliary cirrhosis. Scandinavian Journal of Gastroenterology, 2016, 51, 745-752.	0.6	36
74	Emerging drugs for the treatment of Primary Biliary Cholangitis. Expert Opinion on Emerging Drugs, 2016, 21, 39-56.	1.0	16
75	The management of autoimmunity in patients with cholestatic liver diseases. Expert Review of Gastroenterology and Hepatology, 2016, 10, 73-91.	1.4	12
76	ÂAlkaline phosphatase normalization is a biomarker of improved survival in primary sclerosing cholangitis. Annals of Hepatology, 2016, 15, 246-53.	0.6	39
77	Editorial: betaretrovirus in biliary epithelia of patients with autoimmune and cryptogenic liver disease. Alimentary Pharmacology and Therapeutics, 2015, 41, 490-490.	1.9	3
78	Changing nomenclature for PBC: From â€~cirrhosis' to â€~cholangitis'. Hepatology, 2015, 62, 1620-1622.	3.6	125
79	Current research on the treatment of primary sclerosing cholangitis. Intractable and Rare Diseases Research, 2015, 4, 1-6.	0.3	26
80	Changing nomenclature for PBC: From â€~cirrhosis' to â€~cholangitis'. Clinics and Research in Hepatology and Gastroenterology, 2015, 39, e57-e59.	0.7	36
81	Efficacy of Obeticholic Acid in Patients With Primary Biliary Cirrhosis and Inadequate Response to Ursodeoxycholic Acid. Gastroenterology, 2015, 148, 751-761.e8.	0.6	470
82	Novel therapeutic targets in primary biliary cirrhosis. Nature Reviews Gastroenterology and Hepatology, 2015, 12, 147-158.	8.2	110
83	Primary biliary cirrhosis: safety and benefits of established and emerging therapies. Expert Opinion on Drug Safety, 2015, 14, 1435-1444.	1.0	8
84	ACG Clinical Guideline: Primary Sclerosing Cholangitis. American Journal of Gastroenterology, 2015, 110, 646-659.	0.2	400
85	Development and Validation of a Scoring System to Predict Outcomes of Patients With Primary Biliary Cirrhosis Receiving Ursodeoxycholic Acid Therapy. Gastroenterology, 2015, 149, 1804-1812.e4.	0.6	330
86	Changing nomenclature for PBC: From â€~cirrhosis' to â€~cholangitis'. Journal of Hepatology, 2015, 63, 1285-1287.	1.8	85
87	Primary biliary cirrhosis. Lancet, The, 2015, 386, 1565-1575.	6.3	502
88	Unmet clinical need in autoimmune liver diseases. Journal of Hepatology, 2015, 62, 208-218.	1.8	56
89	Recent advances in the development of farnesoid X receptor agonists. Annals of Translational Medicine, 2015, 3, 5.	0.7	115
90	Commentary: Primary Sclerosing Cholangitis. , 2015, , 61-63.		0

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91	Association between serum IgE level and adverse clinical endpoints in primary sclerosing cholangitis. Annals of Hepatology, 2014, 13, 384-389.	0.6	14
92	Primary sclerosing cholangitis and the microbiota: current knowledge and perspectives on etiopathogenesis and emerging therapies. Scandinavian Journal of Gastroenterology, 2014, 49, 901-908.	0.6	77
93	The Natural History of Primary Biliary Cirrhosis. Seminars in Liver Disease, 2014, 34, 329-333.	1.8	33
94	An overview of current and future therapeutic strategies for the treatment of primary sclerosing cholangitis. Expert Opinion on Orphan Drugs, 2014, 2, 545-556.	0.5	1
95	Levels of Alkaline Phosphatase and Bilirubin Are Surrogate End Points of Outcomes of Patients With Primary Biliary Cirrhosis: An International Follow-up Study. Gastroenterology, 2014, 147, 1338-1349.e5.	0.6	365
96	Ursodeoxycholic acid in primary sclerosing cholangitis: If withdrawal is bad, then administration is good (right?). Hepatology, 2014, 60, 785-788.	3.6	40
97	Low risk of HCC in patients who have PSC and cirrhosis. Nature Reviews Gastroenterology and Hepatology, 2014, 11, 276-277.	8.2	8
98	Neoplasia in the ileoanal pouch following colectomy in patients with ulcerative colitis and primary sclerosing cholangitis. Journal of Crohn's and Colitis, 2014, 8, 1294-1299.	0.6	23
99	Obeticholic acid and budesonide for the treatment of primary biliary cirrhosis. Expert Opinion on Pharmacotherapy, 2014, 15, 365-372.	0.9	38
100	Primary biliary cirrhosis in adults. Expert Review of Gastroenterology and Hepatology, 2014, 8, 427-433.	1.4	31
101	Association between serum IgE level and adverse clinical endpoints in primary sclerosing cholangitis. Annals of Hepatology, 2014, 13, 384-9.	0.6	7
102	Pathogenesis of Primary Sclerosing Cholangitis and Advances in Diagnosis and Management. Gastroenterology, 2013, 145, 521-536.	0.6	359
103	Primary sclerosing cholangitis. Lancet, The, 2013, 382, 1587-1599.	6.3	484
104	Randomised clinical trial: vancomycin or metronidazole in patients with primary sclerosing cholangitis ―a pilot study. Alimentary Pharmacology and Therapeutics, 2013, 37, 604-612.	1.9	212
105	Clinical management of autoimmune biliary diseases. Journal of Autoimmunity, 2013, 46, 88-96.	3.0	18
106	Primary sclerosing cholangitis: a review and update on therapeutic developments. Expert Review of Gastroenterology and Hepatology, 2013, 7, 103-114.	1.4	55
107	Role of the Microbiota and Antibiotics in Primary Sclerosing Cholangitis. BioMed Research International, 2013, 2013, 1-7.	0.9	79
108	Likelihood of Malignancy in Gallbladder Polyps and Outcomes Following Cholecystectomy in Primary Sclerosing Cholangitis. American Journal of Gastroenterology, 2012, 107, 431-439.	0.2	77

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109	Long-term Outcomes of Patients With Primary Biliary Cirrhosis and Hepatocellular Carcinoma. Clinical Gastroenterology and Hepatology, 2012, 10, 182-185.	2.4	24
110	Optimizing biochemical markers as endpoints for clinical trials in primary biliary cirrhosis. Liver International, 2012, 32, 790-795.	1.9	62
111	The safety and efficacy of oral docosahexaenoic acid supplementation for the treatment of primary sclerosing cholangitis – a pilot study. Alimentary Pharmacology and Therapeutics, 2012, 35, 255-265.	1.9	37
112	Pathogenesis and management of pruritus in cholestatic liver disease. Journal of Gastroenterology and Hepatology (Australia), 2012, 27, 1150-1158.	1.4	50
113	Biochemical and immunologic effects of rituximab in patients with primary biliary cirrhosis and an incomplete response to ursodeoxycholic acid. Hepatology, 2012, 55, 512-521.	3.6	130
114	Challenges of Cholangiocarcinoma Detection in Patients with Primary Sclerosing Cholangitis. Journal of Analytical Oncology, 2012, 1, 50-55.	0.1	20
115	Recent developments in the management of idiopathic cholestatic liver disease. Annals of Gastroenterology, 2012, 25, 317-326.	0.4	1
116	Colon Neoplasms Develop Early in the Course of Inflammatory Bowel Disease and Primary Sclerosing Cholangitis. Clinical Gastroenterology and Hepatology, 2011, 9, 52-56.	2.4	53
117	Bone Disease in Patients With Primary Sclerosing Cholangitis. Gastroenterology, 2011, 140, 180-188.	0.6	102
118	Alkaline phosphatase normalization is associated with better prognosis in primary sclerosing cholangitis. Digestive and Liver Disease, 2011, 43, 309-313.	0.4	138
119	Many Patients With Primary Sclerosing Cholangitis and Increased Serum Levels of Carbohydrate Antigen 19-9 Do Not Have Cholangiocarcinoma. Clinical Gastroenterology and Hepatology, 2011, 9, 434-439.e1.	2.4	108
120	Varices in Early Histological Stage Primary Biliary Cirrhosis. Journal of Clinical Gastroenterology, 2011, 45, e66-e71.	1.1	40
121	Antibiotics for the Treatment of Primary Sclerosing Cholangitis. American Journal of Therapeutics, 2011, 18, 261-265.	0.5	24
122	Primary Sclerosing Cholangitis Associated with Elevated ImmunoglobulinG4: Clinical Characteristics and Response to Therapy. American Journal of Therapeutics, 2011, 18, 198-205.	0.5	119
123	High-dose ursodeoxycholic acid increases risk of adverse outcomes in patients with early stage primary sclerosing cholangitis. Alimentary Pharmacology and Therapeutics, 2011, 34, 1185-1192.	1.9	7 5
124	Cancer surveillance in patients with primary sclerosing cholangitis. Hepatology, 2011, 54, 1842-1852.	3.6	248
125	High-Dose Ursodeoxycholic Acid Is Associated With the Development of Colorectal Neoplasia in Patients With Ulcerative Colitis and Primary Sclerosing Cholangitis. American Journal of Gastroenterology, 2011, 106, 1638-1645.	0.2	223
126	New Treatment Strategies for Primary Sclerosing Cholangitis. Digestive Diseases, 2011, 29, 113-116.	0.8	15

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127	Primary Sclerosing Cholangitis Patients With Serial Polysomy Fluorescence In Situ Hybridization Results Are at Increased Risk of Cholangiocarcinoma. American Journal of Gastroenterology, 2011, 106, 2023-2028.	0.2	101
128	Pregnancy in primary sclerosing cholangitis. Gut, 2011, 60, 1027-1028.	6.1	3
129	Moexipril for Treatment of Primary Biliary Cirrhosis in Patients with an Incomplete Response to Ursodeoxycholic Acid. Digestive Diseases and Sciences, 2010, 55, 476-483.	1.1	11
130	Long-term outcomes of positive fluorescence in situ hybridization tests in primary sclerosing cholangitis. Hepatology, 2010, 51, 174-180.	3.6	159
131	Reply: Diagnostic Utility of Chromosome 17 and p16 Abnormalities in Fluorescence In Situ Hybridization Tests in Primary Sclerosing Cholangitis. Hepatology, 2010, 52, 394-395.	3.6	0
132	Fatigue measurements in patients with primary biliary cirrhosis and the risk of mortality during follow-up. Liver International, 2010, 30, 251-258.	1.9	25
133	The possible link between the thyroid and autoimmune liver diseases: reply. Liver International, 2010, 30, 1240-1241.	1.9	1
134	Autoimmune Hepatitis–PBC Overlap Syndrome: A Simplified Scoring System May Assist in the Diagnosis. American Journal of Gastroenterology, 2010, 105, 345-353.	0.2	99
135	Fatigue in primary biliary cirrhosis. Nature Reviews Gastroenterology and Hepatology, 2010, 7, 313-319.	8.2	37
136	Is there a role for tetrathiomolybdate in the treatment of primary biliary cirrhosis?. Translational Research, 2010, 155, 120-122.	2.2	2
137	Linking medical education and patient care. Minnesota Medicine, 2010, 93, 32, 34.	0.1	0
138	Minocycline in the Treatment of Patients With Primary Sclerosing Cholangitis: Results of a Pilot Study. American Journal of Gastroenterology, 2009, 104, 83-88.	0.2	114
139	Primary biliary cirrhosis. Hepatology, 2009, 50, 291-308.	3.6	1,020
140	High-dose ursodeoxycholic acid for the treatment of primary sclerosing cholangitis. Hepatology, 2009, 50, 808-814.	3.6	603
141	B-cell depletion with anti-CD20 ameliorates autoimmune cholangitis but exacerbates colitis in transforming growth factor- \hat{l}^2 receptor II dominant negative mice. Hepatology, 2009, 50, 1893-1903.	3.6	88
142	Review article: nuclear receptors and liver disease – current understanding and new therapeutic implications. Alimentary Pharmacology and Therapeutics, 2009, 30, 816-825.	1.9	15
143	Silymarin in the Treatment of Patients with Primary Sclerosing Cholangitis: An Open-Label Pilot Study. Digestive Diseases and Sciences, 2008, 53, 1716-1720.	1.1	38
144	Conflict of interest policy. Liver Transplantation, 2008, 14, 1-1.	1.3	11

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145	Utility of serum tumor markers, imaging, and biliary cytology for detecting cholangiocarcinoma in primary sclerosing cholangitis. Hepatology, 2008, 48, 1106-1117.	3.6	329
146	Surveillance for hepatocellular carcinoma in patients with primary biliary cirrhosis. Hepatology, 2008, 48, 1149-1156.	3.6	62
147	Clinical trial: randomized controlled study of zidovudine and lamivudine for patients with primary biliary cirrhosis stabilized on ursodiol. Alimentary Pharmacology and Therapeutics, 2008, 28, 886-894.	1.9	52
148	Immunoglobulin G4–Associated Cholangitis: Clinical Profile and Response to Therapy. Gastroenterology, 2008, 134, 706-715.	0.6	807
149	Antimitochondrial Antibody–Negative Primary Biliary Cirrhosis. Gastroenterology Clinics of North America, 2008, 37, 479-484.	1.0	35
150	Clinical features and management of primary sclerosing Cholangitis. World Journal of Gastroenterology, 2008, 14, 3338.	1.4	50
151	Primary sclerosing cholangitis. Canadian Journal of Gastroenterology & Hepatology, 2008, 22, 689-698.	1.8	73
152	Overlap of Autoimmune Hepatitis and Primary Biliary Cirrhosis: Long-Term Outcomes. American Journal of Gastroenterology, 2007, 102, 1244-1250.	0.2	139
153	Ursodeoxycholic Acid for the Treatment of Primary Biliary Cirrhosis. New England Journal of Medicine, 2007, 357, 1524-1529.	13.9	98
154	Clinical Predictors for Hepatocellular Carcinoma in Patients With Primary Biliary Cirrhosis. Clinical Gastroenterology and Hepatology, 2007, 5, 259-264.	2.4	51
155	Immunoglobulin G4 associated cholangitis: Description of an emerging clinical entity based on review of the literature. Hepatology, 2007, 45, 1547-1554.	3.6	224
156	Increased prevalence of antimitochondrial antibodies in first-degree relatives of patients with primary biliary cirrhosis. Hepatology, 2007, 46, 785-792.	3.6	125
157	Impact of inflammatory bowel disease and ursodeoxycholic acid therapy on small-duct primary sclerosing cholangitis. Hepatology, 2007, 47, 133-142.	3.6	35
158	Long-term survival and impact of ursodeoxycholic acid treatment for recurrent primary biliary cirrhosis after liver transplantation. Liver Transplantation, 2007, 13, 1236-1245.	1.3	159
159	Development of autoimmune hepatitis in primary biliary cirrhosis. Liver International, 2007, 27, 1086-1090.	1.9	51
160	Fluoxetine for the Treatment of Fatigue in Primary Biliary Cirrhosis: A Randomized, Double-Blind Controlled Trial. Digestive Diseases and Sciences, 2006, 51, 1985-1991.	1.1	46
161	Elevated Serum IgG4 Concentration in Patients with Primary Sclerosing Cholangitis. American Journal of Gastroenterology, 2006, 101, 2070-2075.	0.2	327
162	Mycophenolate Mofetil for the Treatment of Primary Biliary Cirrhosis in Patients with an Incomplete Response to Ursodeoxycholic Acid. Journal of Clinical Gastroenterology, 2005, 39, 838.	1.1	41

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163	Primary Sclerosing Cholangitis. Inflammatory Bowel Diseases, 2005, 11, 62-72.	0.9	112
164	Alendronate improves bone mineral density in primary biliary cirrhosis: A randomized placebo-controlled trial. Hepatology, 2005, 42, 762-771.	3.6	138
165	Risk factors and comorbidities in primary biliary cirrhosis: A controlled interview-based study of 1032 patients. Hepatology, 2005, 42, 1194-1202.	3.6	560
166	The Value of Serum CA 19-9 in Predicting Cholangiocarcinomas in Patients with Primary Sclerosing Cholangitis. Digestive Diseases and Sciences, 2005, 50, 1734-1740.	1.1	300
167	Mycophenolate Mofetil for the Treatment of Primary Sclerosing Cholangitis. American Journal of Gastroenterology, 2005, 100, 308-312.	0.2	69
168	Mycophenolate mofetil for the treatment of primary biliary cirrhosis in patients with an incomplete response to ursodeoxycholic acid. Journal of Clinical Gastroenterology, 2005, 39, 168-71.	1.1	28
169	Incidence and Risk Factors for Cholangiocarcinoma in Primary Sclerosing Cholangitis. American Journal of Gastroenterology, 2004, 99, 523-526.	0.2	503
170	Ursodeoxycholic acid for treatment of nonalcoholic steatohepatitis: Results of a randomized trial. Hepatology, 2004, 39, 770-778.	3.6	651
171	Cost-minimization analysis of MRC versus ERCP for the diagnosis of primary sclerosing cholangitis. Hepatology, 2004, 40, 39-45.	3.6	117
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