

Gideon Hirschfield

List of Publications by Citations

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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.

296
papers

21,437
citations

62
h-index

142
g-index

330
ext. papers

25,950
ext. citations

9
avg, IF

6.96
L-index

#	Paper	IF	Citations
296	C-reactive protein: a critical update. <i>Journal of Clinical Investigation</i> , 2003 , 111, 1805-1812	15.9	2388
295	C-reactive protein and other circulating markers of inflammation in the prediction of coronary heart disease. <i>New England Journal of Medicine</i> , 2004 , 350, 1387-97	59.2	2317
294	The gut microbiota and host health: a new clinical frontier. <i>Gut</i> , 2016 , 65, 330-9	19.2	1182
293	C-reactive protein: a critical update. <i>Journal of Clinical Investigation</i> , 2003 , 111, 1805-12	15.9	1134
292	A Placebo-Controlled Trial of Obeticholic Acid in Primary Biliary Cholangitis. <i>New England Journal of Medicine</i> , 2016 , 375, 631-43	59.2	574
291	Targeting C-reactive protein for the treatment of cardiovascular disease. <i>Nature</i> , 2006 , 440, 1217-21	50.4	525
290	EASL Clinical Practice Guidelines: The diagnosis and management of patients with primary biliary cholangitis. <i>Journal of Hepatology</i> , 2017 , 67, 145-172	13.4	512
289	Primary biliary cirrhosis associated with HLA, IL12A, and IL12RB2 variants. <i>New England Journal of Medicine</i> , 2009 , 360, 2544-55	59.2	494
288	Genome-wide association study identifies loci influencing concentrations of liver enzymes in plasma. <i>Nature Genetics</i> , 2011 , 43, 1131-8	36.3	415
287	Efficacy of obeticholic acid in patients with primary biliary cirrhosis and inadequate response to ursodeoxycholic acid. <i>Gastroenterology</i> , 2015 , 148, 751-61.e8	13.3	381
286	Primary sclerosing cholangitis. <i>Lancet, The</i> , 2013 , 382, 1587-99	40	370
285	Overlap syndromes: the International Autoimmune Hepatitis Group (IAIHG) position statement on a controversial issue. <i>Journal of Hepatology</i> , 2011 , 54, 374-85	13.4	356
284	Genome-wide meta-analyses identify three loci associated with primary biliary cirrhosis. <i>Nature Genetics</i> , 2010 , 42, 658-60	36.3	337
283	Dense genotyping of immune-related disease regions identifies nine new risk loci for primary sclerosing cholangitis. <i>Nature Genetics</i> , 2013 , 45, 670-5	36.3	267
282	Levels of alkaline phosphatase and bilirubin are surrogate end points of outcomes of patients with primary biliary cirrhosis: an international follow-up study. <i>Gastroenterology</i> , 2014 , 147, 1338-49.e5; quiz e15	13.3	265
281	Development and Validation of a Scoring System to Predict Outcomes of Patients With Primary Biliary Cirrhosis Receiving Ursodeoxycholic Acid Therapy. <i>Gastroenterology</i> , 2015 , 149, 1804-1812.e4	13.3	235
280	Patient Age, Sex, and Inflammatory Bowel Disease Phenotype Associate With Course of Primary Sclerosing Cholangitis. <i>Gastroenterology</i> , 2017 , 152, 1975-1984.e8	13.3	219

279	Baseline ductopenia and treatment response predict long-term histological progression in primary biliary cirrhosis. <i>American Journal of Gastroenterology</i> , 2010 , 105, 2186-94	0.7	213
278	The immunobiology and pathophysiology of primary biliary cirrhosis. <i>Annual Review of Pathology: Mechanisms of Disease</i> , 2013 , 8, 303-30	34	208
277	Simeprevir increases rate of sustained virologic response among treatment-experienced patients with HCV genotype-1 infection: a phase IIb trial. <i>Gastroenterology</i> , 2014 , 146, 430-41.e6	13.3	193
276	Variants at IRF5-TNPO3, 17q12-21 and MMEL1 are associated with primary biliary cirrhosis. <i>Nature Genetics</i> , 2010 , 42, 655-7	36.3	186
275	International genome-wide meta-analysis identifies new primary biliary cirrhosis risk loci and targetable pathogenic pathways. <i>Nature Communications</i> , 2015 , 6, 8019	17.4	185
274	The UK-PBC risk scores: Derivation and validation of a scoring system for long-term prediction of end-stage liver disease in primary biliary cholangitis. <i>Hepatology</i> , 2016 , 63, 930-50	11.2	184
273	Pathogenesis of cholestatic liver disease and therapeutic approaches. <i>Gastroenterology</i> , 2010 , 139, 1481-96	13.6	179
272	C-reactive protein and cardiovascular disease: new insights from an old molecule. <i>QJM - Monthly Journal of the Association of Physicians</i> , 2003 , 96, 793-807	2.7	173
271	Transgenic human C-reactive protein is not proatherogenic in apolipoprotein E-deficient mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005 , 102, 8309-14	11.5	172
270	The evolution of cellular deficiency in GATA2 mutation. <i>Blood</i> , 2014 , 123, 863-74	2.2	153
269	IgG4-related sclerosing disease: autoimmune pancreatitis and extrapancreatic manifestations. <i>Radiographics</i> , 2011 , 31, 1379-402	5.4	152
268	NOTCH2 mutations in Alagille syndrome. <i>Journal of Medical Genetics</i> , 2012 , 49, 138-44	5.8	152
267	Inflammation and endothelial function: direct vascular effects of human C-reactive protein on nitric oxide bioavailability. <i>Circulation</i> , 2005 , 111, 1530-6	16.7	150
266	Genome-wide association study of primary sclerosing cholangitis identifies new risk loci and quantifies the genetic relationship with inflammatory bowel disease. <i>Nature Genetics</i> , 2017 , 49, 269-273	36.3	140
265	A randomized trial of obeticholic acid monotherapy in patients with primary biliary cholangitis. <i>Hepatology</i> , 2018 , 67, 1890-1902	11.2	139
264	norUrsodeoxycholic acid improves cholestasis in primary sclerosing cholangitis. <i>Journal of Hepatology</i> , 2017 , 67, 549-558	13.4	138
263	The British Society of Gastroenterology/UK-PBC primary biliary cholangitis treatment and management guidelines. <i>Gut</i> , 2018 , 67, 1568-1594	19.2	122
262	ImmunoChip analyses identify a novel risk locus for primary biliary cirrhosis at 13q14, multiple independent associations at four established risk loci and epistasis between 1p31 and 7q32 risk variants. <i>Human Molecular Genetics</i> , 2012 , 21, 5209-21	5.6	122

261	The genetics of complex cholestatic disorders. <i>Gastroenterology</i> , 2013 , 144, 1357-74	13.3	111
260	Effect of ileal bile acid transporter inhibitor GSK2330672 on pruritus in primary biliary cholangitis: a double-blind, randomised, placebo-controlled, crossover, phase 2a study. <i>Lancet, The</i> , 2017 , 389, 1114-1123	14.3	110
259	OX40, OX40L and Autoimmunity: a Comprehensive Review. <i>Clinical Reviews in Allergy and Immunology</i> , 2016 , 50, 312-32	12.3	109
258	Stratification of hepatocellular carcinoma risk in primary biliary cirrhosis: a multicentre international study. <i>Gut</i> , 2016 , 65, 321-9	19.2	107
257	PBC screen: an IgG/IgA dual isotype ELISA detecting multiple mitochondrial and nuclear autoantibodies specific for primary biliary cirrhosis. <i>Journal of Autoimmunity</i> , 2010 , 35, 436-42	15.5	103
256	Characterization of animal models for primary sclerosing cholangitis (PSC). <i>Journal of Hepatology</i> , 2014 , 60, 1290-303	13.4	96
255	Role of endoscopy in primary sclerosing cholangitis: European Society of Gastrointestinal Endoscopy (ESGE) and European Association for the Study of the Liver (EASL) Clinical Guideline. <i>Endoscopy</i> , 2017 , 49, 588-608	3.4	94
254	Novel therapeutic targets in primary biliary cirrhosis. <i>Nature Reviews Gastroenterology and Hepatology</i> , 2015 , 12, 147-58	24.2	94
253	Surrogate endpoints for clinical trials in primary sclerosing cholangitis: Review and results from an International PSC Study Group consensus process. <i>Hepatology</i> , 2016 , 63, 1357-67	11.2	94
252	Optimising risk stratification in primary biliary cirrhosis: AST/platelet ratio index predicts outcome independent of ursodeoxycholic acid response. <i>Journal of Hepatology</i> , 2014 , 60, 1249-58	13.4	92
251	The gut-adherent microbiota of PSC-IBD is distinct to that of IBD. <i>Gut</i> , 2017 , 66, 386-388	19.2	91
250	Daclatasvir plus peginterferon alfa and ribavirin for treatment-naive chronic hepatitis C genotype 1 or 4 infection: a randomised study. <i>Gut</i> , 2015 , 64, 948-56	19.2	90
249	Review article: overlap syndromes and autoimmune liver disease. <i>Alimentary Pharmacology and Therapeutics</i> , 2012 , 36, 517-33	6.1	89
248	The immunogenetics of primary biliary cirrhosis: A comprehensive review. <i>Journal of Autoimmunity</i> , 2015 , 64, 42-52	15.5	88
247	Transgenic human CRP is not pro-atherogenic, pro-atherothrombotic or pro-inflammatory in apoE ^{-/-} mice. <i>Atherosclerosis</i> , 2008 , 196, 248-255	3.1	87
246	Seladelpar (MBX-8025), a selective PPAR- δ agonist, in patients with primary biliary cholangitis with an inadequate response to ursodeoxycholic acid: a double-blind, randomised, placebo-controlled, phase 2, proof-of-concept study. <i>The Lancet Gastroenterology and Hepatology</i> , 2017 , 2, 716-726	18.8	81
245	Ustekinumab for patients with primary biliary cholangitis who have an inadequate response to ursodeoxycholic acid: A proof-of-concept study. <i>Hepatology</i> , 2016 , 64, 189-99	11.2	81
244	Ursodeoxycholic acid therapy and liver transplant-free survival in patients with primary biliary cholangitis. <i>Journal of Hepatology</i> , 2019 , 71, 357-365	13.4	80

243	Toronto HCC risk index: A validated scoring system to predict 10-year risk of HCC in patients with cirrhosis. <i>Journal of Hepatology</i> , 2017 ,	13.4	78
242	Effect of NGM282, an FGF19 analogue, in primary sclerosing cholangitis: A multicenter, randomized, double-blind, placebo-controlled phase II trial. <i>Journal of Hepatology</i> , 2019 , 70, 483-493	13.4	77
241	Using GWAS to identify genetic predisposition in hepatic autoimmunity. <i>Journal of Autoimmunity</i> , 2016 , 66, 25-39	15.5	73
240	British Society of Gastroenterology and UK-PSC guidelines for the diagnosis and management of primary sclerosing cholangitis. <i>Gut</i> , 2019 , 68, 1356-1378	19.2	73
239	Long-term efficacy and safety of obeticholic acid for patients with primary biliary cholangitis: 3-year results of an international open-label extension study. <i>The Lancet Gastroenterology and Hepatology</i> , 2019 , 4, 445-453	18.8	73
238	Prospective evaluation of ursodeoxycholic acid withdrawal in patients with primary sclerosing cholangitis. <i>Hepatology</i> , 2014 , 60, 931-40	11.2	72
237	Combined ursodeoxycholic acid (UDCA) and fenofibrate in primary biliary cholangitis patients with incomplete UDCA response may improve outcomes. <i>Alimentary Pharmacology and Therapeutics</i> , 2016 , 43, 283-93	6.1	70
236	Association of primary biliary cirrhosis with variants in the CLEC16A, SOCS1, SPIB and SIAE immunomodulatory genes. <i>Genes and Immunity</i> , 2012 , 13, 328-35	4.4	67
235	X Chromosome Dose and Sex Bias in Autoimmune Diseases: Increased Prevalence of 47,XXX in Systemic Lupus Erythematosus and Sjögren's Syndrome. <i>Arthritis and Rheumatology</i> , 2016 , 68, 1290-1300	9.5	65
234	Role of endoscopy in primary sclerosing cholangitis: European Society of Gastrointestinal Endoscopy (ESGE) and European Association for the Study of the Liver (EASL) Clinical Guideline. <i>Journal of Hepatology</i> , 2017 , 66, 1265-1281	13.4	62
233	Loss of CD28 expression by liver-infiltrating T cells contributes to pathogenesis of primary sclerosing cholangitis. <i>Gastroenterology</i> , 2014 , 147, 221-232.e7	13.3	61
232	CTLA4/ICOS gene variants and haplotypes are associated with rheumatoid arthritis and primary biliary cirrhosis in the Canadian population. <i>Arthritis and Rheumatism</i> , 2009 , 60, 931-7		61
231	Pretreatment prediction of response to ursodeoxycholic acid in primary biliary cholangitis: development and validation of the UDCA Response Score. <i>The Lancet Gastroenterology and Hepatology</i> , 2018 , 3, 626-634	18.8	60
230	Cellular and Molecular Mechanisms of Autoimmune Hepatitis. <i>Annual Review of Pathology: Mechanisms of Disease</i> , 2018 , 13, 247-292	34	59
229	Primary biliary cholangitis: pathogenesis and therapeutic opportunities. <i>Nature Reviews Gastroenterology and Hepatology</i> , 2020 , 17, 93-110	24.2	58
228	The IRF5-TNPO3 association with systemic lupus erythematosus has two components that other autoimmune disorders variably share. <i>Human Molecular Genetics</i> , 2015 , 24, 582-96	5.6	57
227	A randomized, placebo-controlled, phase II study of obeticholic acid for primary sclerosing cholangitis. <i>Journal of Hepatology</i> , 2020 , 73, 94-101	13.4	57
226	Progress in the genetics of primary biliary cirrhosis. <i>Seminars in Liver Disease</i> , 2011 , 31, 147-56	7.3	57

225	Intestinal CCL25 expression is increased in colitis and correlates with inflammatory activity. <i>Journal of Autoimmunity</i> , 2016 , 68, 98-104	15.5	53
224	Mesenchymal stromal cells and liver fibrosis: a complicated relationship. <i>FASEB Journal</i> , 2016 , 30, 3905-3908	3.9	53
223	Factors Associated With Recurrence of Primary Biliary Cholangitis After Liver Transplantation and Effects on Graft and Patient Survival. <i>Gastroenterology</i> , 2019 , 156, 96-107.e1	13.3	52
222	The phenotypic expression of inflammatory bowel disease in patients with primary sclerosing cholangitis differs in the distribution of colitis. <i>Digestive Diseases and Sciences</i> , 2013 , 58, 2608-14	4	51
221	High-throughput T-cell receptor sequencing across chronic liver diseases reveals distinct disease-associated repertoires. <i>Hepatology</i> , 2016 , 63, 1608-19	11.2	51
220	Validation of the prognostic value of histologic scoring systems in primary sclerosing cholangitis: An international cohort study. <i>Hepatology</i> , 2017 , 65, 907-919	11.2	50
219	Utility and cost evaluation of multiparametric magnetic resonance imaging for the assessment of non-alcoholic fatty liver disease. <i>Alimentary Pharmacology and Therapeutics</i> , 2018 , 47, 631-644	6.1	49
218	Expert clinical management of autoimmune hepatitis in the real world. <i>Alimentary Pharmacology and Therapeutics</i> , 2017 , 45, 723-732	6.1	48
217	Anti-kelch-like 12 and anti-hexokinase 1: novel autoantibodies in primary biliary cirrhosis. <i>Liver International</i> , 2015 , 35, 642-51	7.9	48
216	Comparative MRI analysis of morphologic patterns of bile duct disease in IgG4-related systemic disease versus primary sclerosing cholangitis. <i>American Journal of Roentgenology</i> , 2014 , 202, 536-43	5.4	46
215	The challenges of primary biliary cholangitis: What is new and what needs to be done. <i>Journal of Autoimmunity</i> , 2019 , 105, 102328	15.5	45
214	The spectrum of sclerosing cholangitis and the relevance of IgG4 elevations in routine practice. <i>American Journal of Gastroenterology</i> , 2012 , 107, 56-63	0.7	45
213	Unmet clinical need in autoimmune liver diseases. <i>Journal of Hepatology</i> , 2015 , 62, 208-18	13.4	44
212	Mechanisms of tissue injury in autoimmune liver diseases. <i>Seminars in Immunopathology</i> , 2014 , 36, 553-68	2	44
211	Good maternal and fetal outcomes for pregnant women with primary biliary cirrhosis. <i>Clinical Gastroenterology and Hepatology</i> , 2014 , 12, 1179-1185.e1	6.9	44
210	Pathway-based analysis of primary biliary cirrhosis genome-wide association studies. <i>Genes and Immunity</i> , 2013 , 14, 179-86	4.4	44
209	Amyloidosis: new strategies for treatment. <i>International Journal of Biochemistry and Cell Biology</i> , 2003 , 35, 1608-13	5.6	44
208	Major Hepatic Complications in Ursodeoxycholic Acid-Treated Patients With Primary Biliary Cholangitis: Risk Factors and Time Trends in Incidence and Outcome. <i>American Journal of Gastroenterology</i> , 2018 , 113, 254-264	0.7	44

207	Risk stratification in autoimmune cholestatic liver diseases: Opportunities for clinicians and trialists. <i>Hepatology</i> , 2016 , 63, 644-59	11.2	43
206	PR3-ANCA: a promising biomarker in primary sclerosing cholangitis (PSC). <i>PLoS ONE</i> , 2014 , 9, e112877	3.7	43
205	The human lymph node microenvironment unilaterally regulates T-cell activation and differentiation. <i>PLoS Biology</i> , 2018 , 16, e2005046	9.7	43
204	Low-dose interleukin-2 promotes STAT-5 phosphorylation, T survival and CTLA-4-dependent function in autoimmune liver diseases. <i>Clinical and Experimental Immunology</i> , 2017 , 188, 394-411	6.2	42
203	Obeticholic acid for the treatment of primary biliary cirrhosis. <i>Expert Review of Clinical Pharmacology</i> , 2016 , 9, 13-26	3.8	42
202	Autoantibodies and liver disease: uses and abuses. <i>Canadian Journal of Gastroenterology & Hepatology</i> , 2010 , 24, 225-31		42
201	Long-term follow-up of patients with difficult to treat type 1 autoimmune hepatitis on Tacrolimus therapy. <i>Scandinavian Journal of Gastroenterology</i> , 2016 , 51, 329-36	2.4	41
200	Klinefelter syndrome (47,XXY) is in excess among men with Sjögren syndrome. <i>Clinical Immunology</i> , 2016 , 168, 25-29	9	41
199	The inter-relationship of symptom severity and quality of life in 2055 patients with primary biliary cholangitis. <i>Alimentary Pharmacology and Therapeutics</i> , 2016 , 44, 1039-1050	6.1	41
198	Effects of Primary Sclerosing Cholangitis on Risks of Cancer and Death in People With Inflammatory Bowel Disease, Based on Sex, Race, and Age. <i>Gastroenterology</i> , 2020 , 159, 915-928	13.3	37
197	A Pilot Integrative Analysis of Colonic Gene Expression, Gut Microbiota, and Immune Infiltration in Primary Sclerosing Cholangitis-Inflammatory Bowel Disease: Association of Disease With Bile Acid Pathways. <i>Journal of Crohn's and Colitis</i> , 2020 , 14, 935-947	1.5	36
196	Identifying opportunities to improve management of autoimmune hepatitis: evaluation of drug adherence and psychosocial factors. <i>Journal of Hepatology</i> , 2012 , 57, 1299-304	13.4	36
195	The specificity of fatigue in primary biliary cirrhosis: evaluation of a large clinic practice. <i>Hepatology</i> , 2010 , 52, 562-70	11.2	36
194	Vascular adhesion protein-1 is elevated in primary sclerosing cholangitis, is predictive of clinical outcome and facilitates recruitment of gut-tropic lymphocytes to liver in a substrate-dependent manner. <i>Gut</i> , 2018 , 67, 1135-1145	19.2	35
193	Carriage of a tumor necrosis factor polymorphism amplifies the cytotoxic T-lymphocyte antigen 4 attributed risk of primary biliary cirrhosis: evidence for a gene-gene interaction. <i>Hepatology</i> , 2010 , 52, 223-9	11.2	35
192	Milder disease stage in patients with primary biliary cholangitis over a 44-year period: A changing natural history. <i>Hepatology</i> , 2018 , 67, 1920-1930	11.2	35
191	A comprehensive assessment of environmental exposures among 1000 North American patients with primary sclerosing cholangitis, with and without inflammatory bowel disease. <i>Alimentary Pharmacology and Therapeutics</i> , 2015 , 41, 980-90	6.1	34
190	Treatment of autoimmune liver disease: current and future therapeutic options. <i>Therapeutic Advances in Chronic Disease</i> , 2013 , 4, 119-41	4.9	34

189	Gilbert's syndrome: an overview for clinical biochemists. <i>Annals of Clinical Biochemistry</i> , 2006 , 43, 340-3	2.2	34
188	Primary biliary cholangitis. <i>Lancet, The</i> , 2020 , 396, 1915-1926	4.0	33
187	Preprocedural inflammatory markers do not predict restenosis after successful coronary stenting. <i>American Heart Journal</i> , 2004 , 147, 1071-7	4.9	32
186	Goals of Treatment for Improved Survival in Primary Biliary Cholangitis: Treatment Target Should Be Bilirubin Within the Normal Range and Normalization of Alkaline Phosphatase. <i>American Journal of Gastroenterology</i> , 2020 , 115, 1066-1074	0.7	31
185	Human C-reactive protein does not protect against acute lipopolysaccharide challenge in mice. <i>Journal of Immunology</i> , 2003 , 171, 6046-51	5.3	31
184	Diagnostic accuracy of non-invasive tests for advanced fibrosis in patients with NAFLD: an individual patient data meta-analysis. <i>Gut</i> , 2021 ,	19.2	31
183	Multiparametric magnetic resonance imaging for quantitation of liver disease: a two-centre cross-sectional observational study. <i>Scientific Reports</i> , 2018 , 8, 9189	4.9	31
182	A Randomized, Controlled, Phase 2 Study of Maralixibat in the Treatment of Itching Associated With Primary Biliary Cholangitis. <i>Hepatology Communications</i> , 2019 , 3, 365-381	6	30
181	Biliary atresia and survival into adulthood without transplantation: a collaborative multicentre clinic review. <i>Liver International</i> , 2012 , 32, 510-8	7.9	30
180	Amyloidosis: a clinico-pathophysiological synopsis. <i>Seminars in Cell and Developmental Biology</i> , 2004 , 15, 39-44	7.5	30
179	Factors that Influence Health-Related Quality of Life in Patients with Primary Sclerosing Cholangitis. <i>Digestive Diseases and Sciences</i> , 2016 , 61, 1692-9	4	29
178	Fibrosis stage is an independent predictor of outcome in primary biliary cholangitis despite biochemical treatment response. <i>Alimentary Pharmacology and Therapeutics</i> , 2019 , 50, 1127-1136	6.1	29
177	Rare X Chromosome Abnormalities in Systemic Lupus Erythematosus and Sjögren's Syndrome. <i>Arthritis and Rheumatology</i> , 2017 , 69, 2187-2192	9.5	29
176	Phenotyping and auto-antibody production by liver-infiltrating B cells in primary sclerosing cholangitis and primary biliary cholangitis. <i>Journal of Autoimmunity</i> , 2017 , 77, 45-54	15.5	29
175	Insights into the management of Wilson's disease. <i>Therapeutic Advances in Gastroenterology</i> , 2017 , 10, 889-905	4.7	29
174	A validated clinical tool for the prediction of varices in PBC: the Newcastle Varices in PBC Score. <i>Journal of Hepatology</i> , 2013 , 59, 327-35	13.4	28
173	Diagnosis of primary biliary cirrhosis. <i>Baillieres's Best Practice and Research in Clinical Gastroenterology</i> , 2011 , 25, 701-12	2.5	28
172	Recipient HLA-DR3, tumour necrosis factor-alpha promoter allele-2 (tumour necrosis factor-2) and cytomegalovirus infection are interrelated risk factors for chronic rejection of liver grafts. <i>Journal of Hepatology</i> , 2001 , 34, 711-5	13.4	28

171	Factors Associated With Outcomes of Patients With Primary Sclerosing Cholangitis and Development and Validation of a Risk Scoring System. <i>Hepatology</i> , 2019 , 69, 2120-2135	11.2	28
170	Genetic association analysis identifies variants associated with disease progression in primary sclerosing cholangitis. <i>Gut</i> , 2018 , 67, 1517-1524	19.2	28
169	Effects of Age and Sex of Response to Ursodeoxycholic Acid and Transplant-free Survival in Patients With Primary Biliary Cholangitis. <i>Clinical Gastroenterology and Hepatology</i> , 2019 , 17, 2076-2084.e2	6.9	27
168	Twenty-Year Comparative Analysis of Patients With Autoimmune Liver Diseases on Transplant Waitlists. <i>Clinical Gastroenterology and Hepatology</i> , 2018 , 16, 278-287.e7	6.9	27
167	Gut and liver T-cells of common clonal origin in primary sclerosing cholangitis-inflammatory bowel disease. <i>Journal of Hepatology</i> , 2017 , 66, 116-122	13.4	27
166	Autoimmune hepatitis: an approach to disease understanding and management. <i>British Medical Bulletin</i> , 2015 , 114, 181-91	5.4	26
165	Increased sensitivity of Treg cells from patients with PBC to low dose IL-12 drives their differentiation into IFN- β -secreting cells. <i>Journal of Autoimmunity</i> , 2018 , 94, 143-155	15.5	25
164	Improvement of ischemic cholangiopathy in three patients with hereditary hemorrhagic telangiectasia following treatment with bevacizumab. <i>Journal of Hepatology</i> , 2013 , 59, 186-9	13.4	25
163	Patients with autoimmune hepatitis who have antimitochondrial antibodies need long-term follow-up to detect late development of primary biliary cirrhosis. <i>Clinical Gastroenterology and Hepatology</i> , 2012 , 10, 682-4	6.9	25
162	Clinical outcomes of donation after circulatory death liver transplantation in primary sclerosing cholangitis. <i>Journal of Hepatology</i> , 2017 , 67, 957-965	13.4	24
161	Comparative analysis of portal cell infiltrates in antimitochondrial autoantibody-positive versus antimitochondrial autoantibody-negative primary biliary cirrhosis. <i>Hepatology</i> , 2012 , 55, 1495-506	11.2	24
160	Prevalence and risk factors for liver biochemical abnormalities in Canadian patients with systemic lupus erythematosus. <i>Journal of Rheumatology</i> , 2012 , 39, 254-61	4.1	24
159	Efficacy of rituximab in difficult-to-manage autoimmune hepatitis: Results from the International Autoimmune Hepatitis Group. <i>JHEP Reports</i> , 2019 , 1, 437-445	10.3	24
158	Effects of Vedolizumab in Patients With Primary Sclerosing Cholangitis and Inflammatory Bowel Diseases. <i>Clinical Gastroenterology and Hepatology</i> , 2020 , 18, 179-187.e6	6.9	24
157	Grand round: Autoimmune hepatitis. <i>Journal of Hepatology</i> , 2019 , 70, 773-784	13.4	23
156	The Pathogenesis of Primary Biliary Cholangitis: A Comprehensive Review. <i>Seminars in Liver Disease</i> , 2020 , 40, 34-48	7.3	23
155	Primary biliary cirrhosis: one disease with many faces. <i>Israel Medical Association Journal</i> , 2011 , 13, 55-9	0.9	23
154	A liver mass post-Fontan operation. <i>QJM - Monthly Journal of the Association of Physicians</i> , 2014 , 107, 571-2	2.7	22

153	Antimitochondrial antibody-negative primary biliary cirrhosis. <i>Clinics in Liver Disease</i> , 2008 , 12, 323-31; viii-ix	4.6	22
152	Long-term impact of preventive UDCA therapy after transplantation for primary biliary cholangitis. <i>Journal of Hepatology</i> , 2020 , 73, 559-565	13.4	21
151	Changes in natural killer cells and exhausted memory regulatory T Cells with corticosteroid therapy in acute autoimmune hepatitis. <i>Hepatology Communications</i> , 2018 , 2, 421-436	6	21
150	BAT117213: Ileal bile acid transporter (IBAT) inhibition as a treatment for pruritus in primary biliary cirrhosis: study protocol for a randomised controlled trial. <i>BMC Gastroenterology</i> , 2016 , 16, 71	3	21
149	The Pathogenesis of Autoimmune Liver Disease. <i>Digestive Diseases</i> , 2016 , 34, 327-33	3.2	21
148	Liver homing of clinical grade Tregs after therapeutic infusion in patients with autoimmune hepatitis. <i>JHEP Reports</i> , 2019 , 1, 286-296	10.3	20
147	Genetics in PBC: what do the "risk genes" teach us?. <i>Clinical Reviews in Allergy and Immunology</i> , 2015 , 48, 176-81	12.3	20
146	Inequity of care provision and outcome disparity in autoimmune hepatitis in the United Kingdom. <i>Alimentary Pharmacology and Therapeutics</i> , 2018 , 48, 951-960	6.1	20
145	Apical Sodium-Dependent Transporter Inhibitors in Primary Biliary Cholangitis and Primary Sclerosing Cholangitis. <i>Digestive Diseases</i> , 2017 , 35, 267-274	3.2	19
144	The impact of ileal pouch-anal anastomosis on graft survival following liver transplantation for primary sclerosing cholangitis. <i>Alimentary Pharmacology and Therapeutics</i> , 2018 , 48, 322-332	6.1	19
143	Proximity to transplant center and outcome among liver transplant patients. <i>American Journal of Transplantation</i> , 2019 , 19, 208-220	8.7	19
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