



Review

Primary Biliary Cholangitis and Primary Sclerosing Cholangitis: Current Knowledge of Pathogenesis and Therapeutics

Ji-Won Park ^{1,2,†}, Jung-Hee Kim ^{1,2,†}, Sung-Eun Kim ^{1,2}, Jang Han Jung ^{1,2}, Myoung-Kuk Jang ^{1,2}, Sang-Hoon Park ¹, Myung-Seok Lee ¹, Hyoung-Su Kim ^{1,2}, Ki Tae Suk ^{1,2} and Dong Joon Kim ^{1,2,*}

¹ Department of Internal Medicine, College of Medicine, Hallym University, Chuncheon-si 24252, Korea; miunorijw@hallym.or.kr (J.-W.P.); jungheekim@hallym.or.kr (J.-H.K.); sekim@hallym.or.kr (S.-E.K.); con2000@hallym.or.kr (J.H.J.); mkjang@kdh.or.kr (M.-K.J.); sanghoon@hallym.or.kr (S.-H.P.); leemsm@hallym.or.kr (M.-S.L.); hskim@kdh.or.kr (H.-S.K.); ktsuk@hallym.or.kr (K.T.S.)

² Institute for Liver and Digestive Diseases, Hallym University, Chuncheon 200-010, Korea

* Correspondence: djkim@hallym.ac.kr; Tel.: +82-33-240-5646

† These authors contributed equally to this work.

Citation: Park, J.-W.; Kim, J.-H.; Kim, S.-E.; Jung, J.H.; Jang, M.-K.; Park, S.-H.; Lee, M.-S.; Kim, H.-S.; Suk, K.T.; Kim, D.J. Primary Biliary Cholangitis and Primary Sclerosing Cholangitis: Current Knowledge of Pathogenesis and Therapeutics. *Biomedicines* **2022**, *10*, 1288. <https://doi.org/10.3390/biomedicines10061288>

Academic Editor: Jeongeun Hyun

Received: 29 April 2022

Accepted: 28 May 2022

Published: 31 May 2022

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2022 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<https://creativecommons.org/licenses/by/4.0/>).

Abstract: Cholangiopathies encompass various biliary diseases affecting the biliary epithelium, resulting in cholestasis, inflammation, fibrosis, and ultimately liver cirrhosis. Primary biliary cholangitis (PBC) and primary sclerosing cholangitis (PSC) are the most important progressive cholangiopathies in adults. Much research has broadened the scope of disease biology to genetic risk, epigenetic changes, dysregulated mucosal immunity, altered biliary epithelial cell function, and dysbiosis, all of which interact and arise in the context of ill-defined environmental triggers. An in-depth understanding of the molecular pathogenesis of these cholestatic diseases will help clinicians better prevent and treat diseases. In this review, we focus on the main underlying mechanisms of disease initiation and progression, and novel targeted therapeutics beyond currently approved treatments.

Keywords: cholangiopathy; primary biliary cholangitis; primary sclerosing cholangitis

1. Introduction

Cholangiocytes are mature epithelial cells lining the biliary tree. Their functions include the secretion and modification of bile components and the transport of bile to the intestine. Dysfunction of cholangiocytes can lead to the development of various biliary diseases with a chronic–progressive course and often invalidating outcomes. This disease entity affecting the biliary epithelium is known as cholangiopathies. It is caused by diverse etiologies, including genetic, immune-mediated, infectious, drug-induced, idiopathic, malignant and vascular diseases [1]. The perturbed structure and function of cholangiocytes result in impaired bile formation and secretion. These syndromes comprising cholangiopathies have unresolved pathophysiological problems and significant unmet needs in clinical practice [2].

Primary biliary cholangitis (PBC) and primary sclerosing cholangitis (PSC) belong to the main subgroup of chronic cholestatic liver diseases. PBC is a progressive fibrosing cholangiopathy of the small intrahepatic bile ductules (diameter < 100 µm). The prevalence of PBC ranges from 1.91 to 40.2 per 100,000 inhabitants in different geographic areas [3]. In 2015, the nomenclature of PBC was changed from primary biliary cirrhosis to remove the “cirrhosis stigma” and more accurately reflect the disorder and its natural history [4,5]. Although a predisposing genetic background along with infective, immu-

nological, and environmental triggers have been proposed to elucidate the onset of the disease, the etiology of PBC is still unclear [6–8].

PSC is also characterized by the progressive idiopathic stricturing of the biliary system, typically leading to end-stage liver diseases such as cirrhosis and colonic or hepatobiliary malignancy. The prevalence of PSC ranges from one to 16 per 100,000 worldwide [3,9], with an annual incidence of 0.5–0.75 cases per 100,000 [9,10]. The pathogenesis of PSC remains uncertain; however, similar to PBC, it appears to be multifactorial, with environmental triggers leading to cholangiocyte damage and an aberrant and exaggerated cholangiocyte immune response promoting clinical disease in immunogenetically susceptible persons.

Both PBC and PSC are considered immune-mediated cholangiopathies and have features of portal inflammation, biliary tract injury, and sequential fibrosis and cirrhosis development, leading to end-stage liver failure. Although these disorders present common clinical features, PSC is characterized by damage of medium to large extrahepatic and intrahepatic bile ducts, whereas PBC chiefly targets small intrahepatic bile ducts [11,12]. They are uncommon or even rare; however, these diseases often cause considerable morbidity and mortality. The largely unknown etiology and disease mechanisms of PBC and PSC make curative therapies unavailable.

Thus, in this review, we address the mechanism of functional loss of cholangiocytes in the initiation and progression of PBC and PSC, together with the treatments based on currently ongoing studies.

2. Cholangiocyte Pathobiology

Cholangiocytes line a complex network of interconnecting tubes extending from the canals of Hering (CoH) in the liver to the duodenum. CoH begins in the lobules, consists partially of hepatocytes and cholangiocytes, and carries bile from the bile canaliculi to terminal bile ducts in portal tracts [13]. Even though cholangiocytes compose a minor part (3–5% of all liver cells) of the liver, they are essential in bile generation, a life-sustaining function of the liver [14]. The fact that cholangiocytes account for 3–5% of the total liver cell population and generate 25–40% of the total bile flow in humans, which indicates how active these cells are when transporting water and HCO_3^- into the bile duct [15–17].

Cholangiocytes represent heterogeneity with regard to morphological, biochemical, and functional aspects [18]. Immature cholangiocytes within the CoH, and from the intrahepatic and extrahepatic peribiliary glands, are poorly differentiated, and are considered progenitor cells that participate in epithelium restoration and tissue regeneration; however, cholangiocytes gradually obtain a greater degree of differentiation along the biliary tree (from small to large bile ducts) regarding cell polarity, expression of receptors and transporters, and response to hormones [18,19]. Cholangiocytes are classified into small and large cholangiocytes, and each consists of small and large intrahepatic bile ducts, respectively. Large cholangiocytes take part in the alteration of bile composition and volume through secretory and absorptive processes which thoroughly regulated by molecules such as hormones, peptides, and neurotransmitters [20]. In contrast, small cholangiocytes can change their phenotype in response to exogenous or endogenous -stimuli, including microorganisms, toxic material, drugs, and hormones, which consequently participate in the inflammatory response during biliary tract injury [20,21], and function as liver progenitor cells in certain circumstances [22].

Bile, consisting mainly of bile salt, phospholipids, cholesterol, conjugated bilirubin, electrolytes, and water, is a physiological aqueous solution and secretory fluid produced by the hepatobiliary system. Bile contains various components which contribute to health by aiding digestion, maintaining enterohepatic circulation, and supporting the elimination of harmful molecules from the body. Within the ductal lumen, bile is modified via activities at the apical plasma membrane domain of cholangiocytes. Additional-

ly, via tight junctions and immunoglobulin, A (IgA) secretion, and barrier formation, cholangiocytes protect themselves from potentially damaging molecules and microorganisms in bile, and can access the immune and vascular systems through the basolateral plasma membrane domain. These complicated processes are regulated by extracellular signals, biliary constituents (bile acids, glucose, vesicles), and physical forces such as flow and pressure [15].

Cholangiocytes can be affected during liver injury and participate in various liver diseases' pathobiology. Additionally, cholangiocytes can also have a role in liver regeneration when hepatocyte regeneration is impaired [23]. Detailed molecular mechanisms responsible for cholangiocyte dysfunction in PBC and PSC will be reviewed in the following sections.

3. Primary Biliary Cholangitis

3.1. Pathogenesis of PBC

3.1.1. Genetic Factor

PBC can be triggered by environmental factors such as infectious diseases and harmful chemicals in genetically susceptible individuals [6]; therefore, genetic factors are thought to play a substantial role in disease onset. There was a higher concordance rate in monozygotic twins than in dizygotic twins, and familial clustering of patients with PBC (relative risk of 9.13–10.5 in first-degree relatives compared with 1.66 in fifth-degree relatives), which suggests that genetic factors contribute to the occurrence of PBC [24–26]. Although the genetic correlations are lower than those seen in other autoimmune diseases, large-scale genome-wide association studies have recently indicated multiple genes affecting the susceptibility to PBC in human leukocyte antigen (HLA) and non-HLA loci. The HLA complex at chromosomal position 6p21 includes the most polymorphic genes in the human genome [27]. The products of the classical HLA class I (A, B, and C) and class II (DR, DQ, and DP) genes contain cell surface glycoproteins participating in the binding and presentation of self or non-self peptides to T-cell receptors. Class I molecules are recognized by CD8⁺ cytotoxic T cells presenting as endogenous peptides, whereas class II molecules are recognized by CD4⁺ helper T cells presenting as processed peptides from exogenous pathogens [28]. The extent to which endogenous and exogenous peptides bind to HLA molecules is determined by allelic polymorphisms.

HLA class II alleles have been associated with PBC onset for decades. Previous studies have shown that PBC is related to *HLA DR*08* as a predisposing allele and *HLA DRB1*11* and **13* alleles as protective alleles [29,30]. Although HLA alleles are crucial in determining the susceptibility of PBC via alteration of autoantigen presentation, HLA alone does not clarify the whole genetic predisposition to PBC. In fact, 80–90% of patients with PBC do not represent the most common HLA susceptibility alleles [28]. Genome-wide association study (GWAS) findings denoted risk loci including *interleukin (IL)12A*, *IL12RB*, *interferon regulatory Factor 5 (IRF5)*, *transportin 3 (TNPO3)*, *transcription factor Spi-B (SPIB)*, *tumor necrosis factor superfamily, member 15 (TNFSF15)*, and *POU domain class 2-associating Factor 1 (POU2AF1)* [31–33]. Through interaction with death receptor 3 (DR3), TNFSF15 promotes T-cell expansion and induces apoptosis in cells that overexpress DR3. Additionally, synergistic interaction with IL-12 and IL-18 ultimately promotes interferon-gamma production. The reason why much of the heritability of PBC remains uncertain may be explained by the “missing heritability”, such as DNA methylation, histone modification, and noncoding RNAs (i.e., miRNA and lncRNA) that cannot be captured by GWAS. For example, polymorphisms of solute carrier family 4 member 2 (SLC4A2)/anion exchanger 2 (AE2) genes were associated with disease progression in a case-control study [34] but not in GWAS. In addition, telomere dysregulation in biliary epithelial cells (BECs) may be involved in disease onset. A deficiency in sex chromosomes and skewed gene expression in the X chromosome may explain the female pre-

dominance in PBC [35]. A previous study demonstrated that through X-chromosome profiling data of PBC patients, there are distinct DNA methylation patterns in CD4, CD8, and CD14 cells; moreover, demethylation of the C-X-C chemokine receptor 3 (CXCR3) promoter leads to elevated expression of CXCR3 in CD4 T cells in early-stage PBC patients [36]. CXCR3, which is expressed mainly on activated T lymphocytes, natural killer (NK) cells, and some epithelial cells and endothelial cells, plays a key role in inducing leukocyte trafficking, cytoskeletal changes, and chemotactic migration. To investigate the genetic variation underlying the progression of the disease and therapeutic response, further studies are necessary.

3.1.2. Anion Exchanger Deficiency in PBC

Similar to other epithelial cells, cholangiocytes are polarized cells that enable the secretion of bicarbonate (HCO_3^-) and water to the ductular lumen. Thus, fluidizing and alkalizing canalicular bile can be formed [15,16]. The bile epithelium is protected from bile salts through the secretion of HCO_3^- ; therefore, a damaged “biliary bicarbonate umbrella” is considered a common pathology of fibrotic cholangiopathy [37]. Cholangiocytes react to pathogen associated molecular patterns (PAMPs) and damage associated molecular patterns (DAMPs) through the pattern recognition receptors (PRRs). This reaction leads to stimulation of the NF- κ B-dependent cytokines/chemokines secretion. Pro-inflammatory cytokines and chemokines inhibit biliary fluid and bicarbonate secretion by interfering with cAMP-dependent ion transport mechanisms of cholangiocytes [38]. AE2, an electroneutral chloride (Cl^-)/ HCO_3^- exchanger, has been reported to play a central role in maintaining a “biliary bicarbonate umbrella” for protecting biliary mucosa. Bile salt is negatively charged on the outer leaflet of the plasma membrane, where it can be protonated by attracting protons. Consequently, the protonated bile salts are not polar and enter cells freely through diffusion, resulting in apoptosis. The main function of the biliary bicarbonate umbrella is that of the alkaline barrier, which is maintained by biliary HCO_3^- secretion, and sets bile salts in their polar, deprotonated, and membrane-impermeable state; therefore, dysfunctional AE2 may be related to the pathogenesis of PBC [39–41]. In addition to several studies indicating that AE2 was reduced in PBC, a recent study reported that downregulation of AE2 may sensitize BECs to apoptotic insults activating soluble adenylyl cyclase (sAC) [39]. sAC, an evolutionarily conserved bicarbonate sensor, plays a key role in regulating bile salt-induced apoptosis [42]. Reduced AE2 expression leads to a decrease in the bicarbonate secretion and bicarbonate accumulation in the cells. Increased intracellular bicarbonate modulates bile salt induced apoptosis by increasing sAC activity. Decreased bicarbonate secretion allows more bile salts to enter cells, releasing intracellular Ca^{2+} stores, thereby increasing sAC activity. In vitro studies reported that inhibition of sAC reversed sensitization to bile salt-induced apoptosis and prevented bile salt-induced apoptosis altogether [39,40]. Motoko et al., have previously reported that uncontrolled autophagy may play an important role in the pathogenesis of PBC by causing autoimmune processes through aberrant expression of mitochondrial antigens such as the pyruvate dehydrogenase complex, E2 (PDC-E2) and promoting cellular senescence in BECs in the biliary tract in PBC [41,43,44]. More recently, it was disclosed that the decreased expression of AE2 was closely correlated with the abnormal expression of PDC-E2 and autophagy-related markers LC3 and p62 [45]. Furthermore, AE2 knockdown was reported to induce cellular senescence [45,46]. Interestingly, dysregulated autophagy can induce cellular senescence [47]; thus, it is likely that cellular senescence can be induced through unregulated autophagy due to AE2 downregulation in PBC. Senescent BECs express diverse chemokines and cytokines with a senescence-associated secretory phenotypes in PBC and PSC, which are involved in inflammatory cell infiltration and fibrosis in cholangiopathies [43,48–50].

In addition, it is interesting that allelic variations in the AE2 gene have a meaningful association with the disease progression rate in PBC under ursodeoxycholic acid (UDCA) treatment [34].

3.1.3. Anti-Mitochondrial Antibodies (AMA) in PBC

The presence of disease-specific autoantibodies, that is, serum AMA, suggests that autoimmunity is one of the central mechanisms of PBC [51]. AMA was first detected for a non-organ-specific ATPase-associated antigen called M2, and it was primarily directed to PDC-E2 [52]. AMA is considered as the typical characteristic of PBC and is present in 90–95% of patients [53]; however, it can be found in less than 1% of healthy subjects. AMA targets lipoic acid-containing immunodominant epitopes, especially the E2 subunits of the 2-oxoacid dehydrogenase complex enzymes, including PDC-E2. For unknown reasons, PDC-E2 is abnormally expressed in the luminal surface of the bile duct epithelial cells of PBC patients, resulting in a pathogenetic process, so-called “autoimmune epithelitis” [54]. CD8⁺ T cells recognize this epitope, resulting in subsequent bile duct injury and accumulating bile acids to toxic concentrations [55,56]. The T-lymphocyte-mediated destruction of small bile ducts is followed by secondary damage of hepatocytes from the accumulation of potentially toxic molecules such as bile acids, which are normally secreted into the bile. Naturally, generated bile acids (cholic acid, chenodeoxycholic acid, and deoxycholic acid) are all detergents and can lyse cell membranes if present at a substantially high concentration, which is often reached in cholestasis [56].

Although AMA has diagnostic value, it has no prognostic value. In other words, AMA titer and subtypes are not associated with disease severity and outcome [57–59]. Although there was a report that treatment with UDCA may decrease the AMA titer, it is controversial whether the AMA titer is associated with treatment response [60,61].

AMA-negative PBC accounts for 5–10% of PBC cases [54,62]. PBC-specific anti-nuclear antibodies (ANA) (anti-GP210 and/or anti-SP100) might help to diagnose AMA-negative subjects. Interestingly, anti-SP100 and anti-GP210 are specific for PBC and correlate with disease severity [63].

3.1.4. Immune Response in PBC

The immune system plays a central role in PBC pathogenesis, and various immune cells have been shown to infiltrate the portal tract areas of PBC patients. The primary cause of PBC relates to the loss of immune tolerance to PDC-E2 [64]. PDC-E2 is located on the inner mitochondrial membrane and contains a lipoic acid–lysine bond, which is essential for antigen recognition and immune cell activation [65,66]. Disease-specific AMAs target immunodominant epitopes, chiefly PDC-E2. Aberrant modification of mitochondrial PDC-E2 occurs within apoptotic BECs, and characteristic apoptotic blebs containing immunologically intact PDC-E2 are released. The immunogenic complex is recognized by circulating AMAs; as a result, antigen–antibody complexes are formed [67]. Then, autoantigen–AMA complexes are recognized by innate immune cells such as macrophages, and the disrupted immunotolerance of the liver leads to the further recruitment of various immune cells into the liver [67,68].

BECs can also directly act as antigen-presenting cells to present CD1d-restricted antigens to invariant natural killer T (iNKT) cells, resulting in the activation of this key immune cell subset [69]. In addition to enhancing immune activation, BEC can sustain an inflammatory profile in PBC, which is achieved by secreting several chemokines and recruiting the corresponding immune cells in the liver [48,70]. Once the immune response is set up against aberrant autoantigens expressed on cholangiocytes, PDC-E2-specific autoreactive CD4 T and CD8 T cells are selectively abundant in the livers of PBC patients [71]. Additionally, autoreactive B cells may act as important antigen-presenting cells via the uptake and presentation of autoantigens to T cells [72], which ultimately activate autoreactive B cells to release more AMA [73], thus developing a positive feedback loop that induces BEC injury in PBC.

During PBC progression, a shift from the Th1 to Th17 response occurs. Th17 cell activation increases with disease progression; thus, maximal Th17 activation can be a feature of the progressive disease stage [74,75]. In addition to promoting inflammation,

Th17 cells are known to perform profibrotic functions such as interleukin 17A (IL-17A), as a mechanism to promote the proliferation of hepatic stellate cells; however, this pathologic mechanism has not been proven in PBC [76]. An in vitro study demonstrated that mucosal-associated invariant T (MAIT) cells, another main source of IL-17 in later stages of PBC, induced a profibrotic and activated phenotype of human hepatic stellate cells (HSCs) [77–79]. Subsequently, immune-mediated mechanisms involving autoreactive T cells contribute to the development of chronic liver inflammation and direct bile duct injury by secreting a wide variety of pathologic factors.

3.1.5. Gut Microbial Profile in PBC

The human gut harbors a complicated ecosystem of trillions of microbial cells, called the gut microbiome, contributing to essential functions such as regulating metabolism and immunity [80,81]. Accumulating evidence implies potential links between the gut microbiome and PBC [82–84]. PBC can change the gut microbiome by causing intestinal motility disorders, immunologic derangement, bile secretory defects, and portal hypertension [84,85]. The interactive relationship between the gut and the liver, called the “gut-liver axis”, is established by the portal vein, which enables the transport of gut-derived products directly to the liver [86].

The biliary epithelium expresses toll-like receptors (TLRs). Once various ligands, including microbial products such as lipopolysaccharide (LPS), bind to TLRs, cellular injury occurs through the proinflammatory nuclear factor- κ B pathway and IL-8 and CX3C-chemokine ligand 1 (CX3CL1) release, which facilitates the recruitment of effector lymphoid cells into portal tracts in the livers of PBC patients [87,88]. In contrast, MAIT cells, a novel subset of innate-like T cells, have a critical role in protecting the biliary tree from microbial triggers; thus, MAIT cells are referred to as a “biliary firewall” [89]. Under healthy conditions, MAIT cells that are abundant in the portal vein are activated by antigen exposure and enhance the local immune response to suppress pathogens [77]; however, an amount of intrahepatic MAIT cells is decreased in PBC, therefore, the protective role of MAIT cells in the maintenance of biliary integrity is limited, especially after exposure to bacterial pathogens [90]. Interestingly, a recent study showed that even after response to UDCA treatment, the amount of MAIT cells did not normalize, which suggested the mechanism of the disease progresses in spite of the treatment response in PBC [79,89].

Additionally, portal plasma cells produce liver-derived secretory IgA, which is secreted into the lumen via bile. Secretory IgA is also able to contribute to protecting the biliary tree from microorganisms by agglutinating or entrapping bacteria, neutralizing bacterial toxins, altering the virulence of bacteria via disruption of gene expression and inhibiting bacterial access to enterocytes [91]. Compared with healthy individuals, PBC patients have reduced secreted IgA from duodenal enterocytes. Moreover, relevant localization of IgA to the basolateral membrane and expansion of tight junctions potentially serving as entrances to the bile tree were observed [92,93].

Distinct fecal microbial features in patients with PBC include increased colonization of *Enterobacteriaceae*, *Pseudomonas*, *Veillonella*, and *Clostridium*, and decreased *Oscillospira* and *Sutterella* [94,95]. In a previous experimental study, long-term bacterial exposure in normal mice was associated with autoantibody production and histological signatures resembling PBC [96]. Recently, Tang et al., reported differences in the fecal microbiota from PBC patients naïve to UDCA compared with healthy controls [97]. In PBC patients, the gut microbiota richness was significantly reduced. At the phylum level, Bacteroidetes spp. were significantly decreased, whereas Fusobacteria and Proteobacteria spp. were overrepresented. At the genus level, Bacteroidetes spp., *Sutterella*, *Oscillospira* and *Faecalibacterium* were significantly decreased in patients, whereas eight genera (*Haemophilus*, *Veillonella*, *Clostridium*, *Lactobacillus*, *Streptococcus*, *Pseudomonas*, *Klebsiella*, and an unknown genus in the family *Enterobacteriaceae*) were significantly increased [98].

Another study, including early-stage PBC patients and healthy controls, demonstrated that the guts of PBC patients were depleted of some potentially beneficial bacteria, such as Acidobacteria, *Lachnobacterium* spp., *Bacteroides eggerthii*, and *Ruminococcus bromii*, but were plentiful in some bacterial taxa including opportunistic pathogens, such as γ -Proteobacteria. [99]. As a result of that study, it was suggested that the alteration of the gut microbiome could be critical for the onset or development of PBC by interacting with metabolism and immunity. Interestingly, Tang et al. found that PBC-associated dysbiosis was partially reversed during UDCA treatment [97]. This finding suggests that modifications of specific bacterial species—for example, those sensitive to bile salt concentration and those involved in bile acid metabolism—may occur after UDCA treatment. Bile acids, which are important metabolites of the microbiome, can modify the composition of the gut microbiota directly or indirectly through activation of the innate immune system [100,101]. Considering the results of recent studies, once PBC occurs together with dysbiosis of the gut microbiota, a vicious cycle may be established, leading to harmful and even fatal outcomes in patients.

3.2. Currently Approved Disease-Modifying Therapies for PBC

3.2.1. UDCA

Until recently, UDCA was an accepted first-line treatment for PBC. UDCA targets bile secretion and bile acid synthesis to improve biochemical markers, histological findings, complication rates, and transplant-free survival [102].

UDCA consists of a hydrophilic dihydroxy bile acid originating in the colon via bacterial 7 β epimerization of chenodeoxycholic acid (CDCA) and is present in low concentrations in human bile acids. UDCA is passively absorbed through the colonic mucosa and then enters the circulating bile acid pool [103]. Although UDCA does not alter total bile acid amount, serum cholesterol levels, or bile phospholipid production, UDCA promotes the metabolic conversion of cholesterol to bile acids and reduces the cholesterol fraction of biliary lipids [104]. The suggested mechanisms of UDCA are as follows: (1) enrichment of the hydrophilic bile acid pool with exogenous UDCA that replaces endogenous hydrophobic toxic bile acids, which often occurs in cholestasis; (2) stimulation of bile secretion via upregulation of AE2 expression on the surface of BECs; and (3) immune modulation through a reduction in hepatocellular and biliary expression of major histocompatibility complex (MHC) class I and MHC class II proteins, possibly reducing adaptive immunity-mediated injury [105]. Additionally, reduced toxic bile salt disruption of cholesterol-rich membranes, especially of highly exposed cholangiocytes, leads to the stabilization of cell structures such as plasma membranes and mitochondria; thus, cytoprotective effects can be induced by UDCA [103]. Additionally, the subcellular anti-apoptotic pathway might be activated by UDCA, which possibly promotes mitochondrial membrane stability via the inhibition of the Bcl-2-associated X protein (BAX) translocation, a pro-apoptotic protein related to the initiation of core apoptotic pathways within mitochondria in hepatocyte cell lines [103]. UDCA might also suppress deregulated cholangiocyte autophagy pathways related to intense endoplasmic reticulum stress caused by toxic hydrophobic bile acids, as observed in cultured BECs and livers of PBC patients [43,44].

UDCA at a dose of 13–15 mg/kg/day has been recommended to decrease the progression of PBC [102,106]. To define the biochemical response, several criteria have been proposed (Table 1). Most studies have indicated that alkaline phosphatase (ALP) and total bilirubin are the two most meaningful variables in evaluating the UDCA response. Above this, prediction of the response before the commencement of UDCA is attempted in the UK and Italy [107]. Bilirubin, ALP, transaminase, age, and lag time from diagnosis to treatment are suggested as parameters predicting the response to UDCA. Up to 40% of patients with PBC will denote a suboptimal biochemical response to UDCA [108]; therefore, other treatment options are necessary.

Table 1. Response criteria of UDCA therapy in patients with PBC.

Criteria	Definition of Response (Time to Evaluation, Months)	Ref.
Barcelona	> 40% decrease or normalization of ALP (12)	[109]
Mayo	ALP < 2 × ULN (6)	[110]
Paris I	ALP ≤ 3 × ULN and AST ≤ 2 × ULN and normalization of bilirubin (12)	[111]
Rotterdam	Normalization of bilirubin and/or albumin (12)	[112]
Ehime	≥ 70% decrease or normalization of GGT (6)	[113]
Toronto	ALP ≤ 1.67 × ULN (24)	[114]
Paris II	ALP and AST ≤ 1.5 × ULN and normalization of bilirubin (12)	[115]

ALP: alkaline phosphatase, ULN: upper limit of normal, GGT: γ -glutamyl transferase, AST: aspartate aminotransferase.

3.2.2. Obeticholic Acid (OCA)

There is no consensus on therapies for patients with suboptimal biochemical responses to UDCA. OCA is the first novel adjunctive licensed agent beyond the traditional use of UDCA alone in PBC. OCA, a semisynthetic hydrophobic bile acid analog, is highly selective for the farnesoid X receptor (FXR) because bile acid becomes more hydrophobic and a more potent FXR activator [116]. OCA with enhanced hydrophobicity through modification of CDCA showed 100-fold improved FXR potency compared with CDCA [117]. FXR is abundantly expressed in the liver and enterocytes. Understanding the mechanism of action of FXR agonism in chronic cholestasis has resulted in the development of OCA.

Primary bile acids such as cholic acid and CDCA bind FXR and downstream signaling pathways, ultimately inhibit the transcription of *CYP7A1* and the potent suppression of bile acid synthesis [4]. In reabsorption from the intestinal lumen, bile acids also activate FXR in ileal enterocytes. FXR activation in these cells results in the expression of fibroblast growth factor 19 (FGF19), which enters the portal circulation and binds to cell surface fibroblast growth factor receptor 4 (FGFR4) on hepatocytes, resulting in the suppression of *CYP7A1* expression and bile acid synthesis [118–120]. In addition, activation of FXR promotes the export of bile acids out of hepatocytes and enterocytes and hepatocyte and ileal reuptake [121,122].

A double-blind phase III clinical trial from the PBC OCA International Study of Efficacy (POISE) group demonstrated that 12 months of OCA therapy (add-on to UDCA or as monotherapy) led to a better biochemical response than the placebo group in nearly half of the PBC patients who were prior biochemical non-responders or intolerant to UDCA. A recent 3-year interim analysis showed significantly decreased ALP and bilirubin after 48 months of OCA treatment compared with the baseline [123,124].

3.3. Novel Therapies Currently under Investigation for PBC

3.3.1. Peroxisome Proliferator-Activated Receptor (PPAR) Agonists

PPAR, a nuclear hormone receptor, participates in multiple metabolic processes, including the regulation of bile acid homeostasis [125]. PPAR agonists have been known to induce beta-oxidation gene expression, decreasing oxidative stress and inflammation in the liver while increasing the secretion of favorable adipokines [126]. PPAR agonists also inhibit hepatic bile-acid transportation [127]. PPAR has three distinct isoforms, α , δ and γ . PPAR α , which is abundantly present in the liver, induces the expression of numerous genes related to lipid and bile-acid metabolism and the downregulation of genes in immune-related pathways [128]. In cholangiocytes, PPAR δ agonism has been known to coordinate cholesterol flux and bile acid metabolism and has a role in apoptotic cell

elimination via macrophages, reducing the autoimmune response against self-antigens released by dying cells [129,130]. PPAR γ , found in the intrahepatic biliary epithelium, suppresses proinflammatory cytokine production and contributes to bile acid homeostasis [131]. PPAR γ levels are reduced with bile duct injury. A previous study reported that administration of PPAR γ ligands significantly reduced portal inflammation and the number of T cells in a mouse model of PBC [132]. Reports from the USA, Europe, and Asia demonstrated good efficacy of fenofibrate, an oral medication of the fibrate class (PPAR α ligands), in PBC patients with a suboptimal response to UDCA [133,134]. Although the FDA approves fibrates only as lipid-lowering agents, their effect on lowering biochemical markers of cholestasis, such as ALP and γ -glutamyltransferase (GGT) levels, has been reported [135].

Compared with fenofibrate, which is specific to PPAR α , bezafibrate is a pan (α , β/δ , γ) PPAR activator. Previous studies demonstrated that bezafibrate combined with UDCA could significantly decrease ALP, GGT, alanine aminotransferase (ALT), IgM, triglyceride, and total cholesterol levels [136,137]. In recent studies published after 2018, bezafibrate, combined with UDCA in PBC patients unresponsive to UDCA monotherapy, improved liver biochemistry and prognostic scores (e.g., UK-PBC and GLOBE scores) for long-term prognoses [94,138]; however, unfortunately, adverse reactions with bezafibrate-UDCA combination therapy are more frequent than UDCA monotherapy, including polydipsia, exacerbation of itching, arthritis, elevated serum creatinine levels and muscle pain, leg edema and gastrointestinal discomfort [136,139].

Recently, data on the efficacy and safety of elafibranor, a novel PPAR α and PPAR δ agonist, for treating noncirrhotic patients with PBC and an incomplete response to UDCA were published [140]. Patients receiving a dose of 80–120 mg of elafibranor showed a significant decrease in ALP compared with the placebo group. Additionally, adding the dual PPAR α and PPAR γ agonist, saroglitazar, to UDCA resulted in a significant decrease in ALP levels, showing an acceptable toxicity profile [141].

As a second-line treatment, seladelpar, a potent and selective PPAR δ agonist, has been investigated [142,143]. The phase 3 ENHANCE study was conducted in patients with PBC who did not respond to first-line treatment. This clinical trial was terminated early because of an unexpected histologic finding (nonalcoholic steatohepatitis); however, the finding was revealed to be unrelated to seladelpar. When a blind analysis following termination was performed, the treatment response rate was significantly higher in the seladelpar group. Additionally, seladelpar improved PBC patients' pruritus and quality of life [144]; therefore, ongoing studies on the efficacy and safety in PBC patients are noteworthy.

3.3.2. Budesonide

Budesonide, the second generation of corticosteroids, has a high first-pass metabolism within the liver; thus, fewer systemic adverse effects is reported than conventional glucocorticosteroids. Previous studies demonstrated that budesonide (6–9 mg/day) added to UDCA (15 mg/kg/day) in PBC patients showed better biochemical and histological improvement than UDCA monotherapy [145]; however, a recent small-scale randomized clinical trial demonstrated that budesonide (9 mg/day) added to UDCA (12–16 mg/kg/day) for 36 months failed to attain histological improvement in PBC patients with a suboptimal response to UDCA monotherapy [146]. Nevertheless, a combination of budesonide and UDCA might be an effective therapy for PBC patients; thus, further investigation is essential to evaluate the efficacy on long-term clinical outcomes, including mortality and the requirement for liver transplantation.

3.3.3. Fibroblast Growth Factor (FGF) 19 Analog

FGF19, an endocrine hormone, is induced in the intestine with the activation of FXR [147]. In the liver, FGF19 suppresses the expression of CYP7A1, the gene encoding cholesterol 7 α -hydroxylase, which catalyzes the first and rate-limiting step in the classic pathway of bile acid synthesis [148]. Administration of FGF19 has been reported to reduce liver damage in mouse models of intrahepatic and extrahepatic cholestasis [149,150]; however, the therapeutic potential of FGF19 is limited because of concerns about tumorigenicity, such as the development of hepatocellular carcinoma induced by ectopic overexpression of FGF19 in mice [151]. To overcome this defect, NGM282, a nontumorigenic engineered analog of FGF19, was developed to treat PBC, and MGM282 administration for 28 days showed a significant response in ALP and transaminase levels compared with placebos in PBC patients with an inadequate response to UDCA [152].

3.3.4. Other Farnesoid X Receptor Agonists

Despite the therapeutic potential of OCA, severe hepatotoxicity recorded in patients with advanced liver disease or drug-related side effects, such as pruritus, are factors limiting the therapeutic use of OCA. Some of these side effects can be caused by the bile acid-like structure and behavior of the OCA molecule. Namely, enterohepatic circulation of OCA and its metabolites and significant G protein-coupled bile acid receptor 1 (TGR5) agonistic properties are challenging pharmacokinetics [153]. Thus, new non-bile acid FXR agonists are in various stages of preclinical and clinical development [154]. EDP-305, a novel non-bile acid FXR agonist, has minimal activity against TGR5. A recent experimental study reported the therapeutic efficacy of EDP-305 in direct comparison with OCA in mouse models of liver disease [155]. Phase 2 clinical trials assessing the efficacy and safety of non-bile acid-type FXR agonists, including EDP-305, tropifexor (LJN452), and cilofexor (GS-9674), in PBC patients have recently been completed; however, the results have not yet been published.

3.3.5. Baricitinib

Baricitinib, a novel small molecule approved in 2018 for the treatment of moderate to severe rheumatoid arthritis, is a Janus kinase (JAK) 1 and 2 inhibitor [156]. JAK, an intracellular enzyme, responds to cytokine and growth factor receptor stimulation to affect downstream hematopoiesis and immune cell function. JAK activates signal transducers and activators of transcription (STAT) through phosphorylation, and the JAK-STAT pathway is responsible for key immune signals. A recent whole-genome linkage study suggested that JAK-STAT pathway proteins have a potential role in developing PBC [157]. A randomized, double-blind placebo-controlled trial was initiated to evaluate the efficacy and safety of baricitinib, but the study was terminated early because of low enrollment. This proof-of-concept study demonstrated that in PBC patients with a suboptimal response to UDCA, baricitinib showed a 30% decrease in ALP and improvement in inflammation and liver fibrosis markers [158].

3.3.6. S-adenosyl-L-methionine (SAME)

S-adenosyl-L-methionine (SAME), an endogenous molecule with hepatoprotective properties linked to redox regulation and methylation, is synthesized from methionine and adenosine triphosphate (ATP). SAME not only maintains mitochondrial function by raising glutathione levels, but also reduces fibrosis through inhibiting collagen secretion in activated HSCs [159–161]. In various chronic liver diseases, the SAME biosynthesis can be decreased [162]. A prospective, open-label pilot study to evaluate the effect of SAME and UDCA reported a significant improvement in liver biochemistry, such as ALP, GGT, total cholesterol, fatigue, and pruritus, in addition to noncirrhotic patients

with PBC [163]. Through analyzing serum samples of patients treated with SAME and an in vitro study, Kilanczyk et al., suggested that SAME may inhibit autoimmune events in PBC patients via its antioxidant and S-glutathionylation properties [164].

3.3.7. Probiotics

The use of *L. rhamnosus* GG for hepatitis, cholestasis, and fibrosis, following common bile duct ligation in mice, improved biochemical and histological indications. This can probably be explained by increased FXR activity by probiotics in the intestine. Activated FXR enhances the formation of FGF15, which reduces the production of bile acids in the liver through negative feedback. The intake of this probiotic resulted in increased activity of FXR in the intestine and increased levels of FGF15 in the blood. The use of a potent FXR antagonist blocks the positive effect of probiotics. In tissue cultures, the culture supernatant containing probiotics increases the activity of FXR and supports the relationship between probiotics and FXR activity [165].

Additionally, *L. rhamnosus* GG increases the content of Firmicutes and Actinobacteria in the gut microbiota, which converts primary bile acids into secondary bile acids, which are poorly absorbed and are consequently removed via excretion. Until now, clinical trials of probiotics in PBC have been very sparse. Considering the very encouraging results of the experimental study, further studies on this topic would potentially be very interesting.

3.3.8. Mesenchymal Stem Cells

Liver transplantation (LT) is still the most essential treatment for patients with advanced, end-stage PBC. Mesenchymal stem cell (MSC) transplantation has been proposed as an effective alternative therapy for PBC patients. MSCs are fibroblast-like, multipotent cells, which can be present in almost all postnatal organs and tissues, including the liver [166]. Few studies using umbilical cord (UC) and bone marrow (BM) MSCs have been reported [167,168]. The clinical study conducted by Wang et al., demonstrated that after BM-MSC transplantation, patients' quality of life was improved, and ALT, aspartate aminotransferase (AST), GGT and IgM significantly decreased. Additionally, histological deterioration, such as fibrosis, was not observed in BM-MSC-treated patients [168]. In addition, the beneficial effects of BM-MSCs included an increase in Tregs, a decrease in inflammatory cytotoxic CD8⁺ T cells, and elevated IL-10 and anti-inflammatory cytokines. Although MSC-derived immunomodulation can play a crucial role in attenuating PBC, further studies are necessary to decide the optimal frequency of MSC infusion and assess the safety of MSC-based therapy in long-term follow-ups.

3.3.9. Anti-Fractalkin Antibody E6011

Fractalkine (CX3CL1) is a chemokine with both chemoattractant and cell-adhesive functions, which interacts with its receptor CX3CR1 in the chemoattraction and recruitment of intraepithelial lymphocytes. Previous studies have reported that CX3CL1 in PBC could be important in the development and maintenance of portal lymphocyte infiltration in PBC [169]. Injured bile ducts resulting from PBC lead to the upregulation of CX3CL1 expression in BECs, followed by the chemoattraction of CX3CR1-expressing mononuclear cells, including CD4⁺ and CD8⁺ T cells, their adhesion to BECs, and the accumulation of biliary intraepithelial lymphocytes. E6011, a novel humanized anti-fractalkine monoclonal antibody, was developed to treat various inflammatory diseases, including Crohn's disease, rheumatoid arthritis, and PBC [170]. Unfortunately, a clinical trial (NCT03092765) to assess the efficacy and safety of E6011 was terminated early. Although the reason for early termination is not a safety issue, it is unclear whether a new clinical trial using E6011 will be initiated.

The main results relating to the novel therapeutics mentioned above are summarized in Table 2.

Table 2. Novel therapies currently under investigation in PBC.

Agent	Mechanism	Clinical Trial Stage	Main Results	Ref.
Bezafibrate	Panspecific PPAR agonist	Phase III (BEZURSO trial)	<ul style="list-style-type: none"> Complete biochemical response: 31% vs. 0%; ALP normalization: 67% vs. 2%; Improvement in liver stiffness: 15% decrease vs. 22% increase. 	[94]
Elafibranor	PPAR α and PPAR δ agonist	Phase II (NCT03124108)	<ul style="list-style-type: none"> Significant decrease in ALP: 41–48% vs. 3% increase; ALP < 1.67 \times ULN, ALP decrease > 15%, total bilirubin < ULN: 67–79% vs. 6.7%. 	[140]
Saroglitazar	PPAR α and PPAR γ agonist	Phase II (EPICS)	<ul style="list-style-type: none"> Mean percentage reduction in ALP: 48.9–50.6% vs. 3.3%; ALP < 1.67 \times ULN, ALP decrease > 15%, total bilirubin < ULN: 69–71% vs. 10%. 	[141]
Tropifexor	FXR agonist	Phase II (NCT02516605)	Not yet published	
Cilofexor	FXR agonist	Phase II (NCT02943447)	Not yet published	
EDP-305	FXR agonist	Phase II (NCT03394924)	Not yet published	
Baricitinib	JAK 1 and 2 inhibitor	Proof-of-concept study	<ul style="list-style-type: none"> Thirty percent decrease in ALP; Improvement in the itch NRS; Increase in the fatigue NRS; Improvement of inflammation and liver fibrosis marker. 	[158]
S-adenosyl-L-methionine	17-beta-estradiol glucuronide-induced cholestasis reversal agent	Pilot, open-label study	<ul style="list-style-type: none"> A positive effect of adding SAME to UDCA in noncirrhotic PBC patients. 	[163]
Probiotics	Regulation of bile acid homeostasis	Phase II (NCT03521297)	Data yet to be collected	
Mesenchymal Stem Cells	Immunoregulation	NCT03668145	Data yet to be collected	

PPAR: Peroxisome proliferator-activated receptor, ALP: alkaline phosphatase, ULN: upper limit of normal, SAME: S-adenosyl-L-methionine, UDCA: Ursodeoxycholic acid, PBC: primary biliary cholangitis, FXR: farnesoid X receptor.

4. Primary Sclerosing Cholangitis

PSC is a chronic hepatobiliary disease affecting the intra- and extrahepatic bile duct by multifocal fibrotic bile duct stricture, dilatation, and cholestasis. Gradual injury of the biliary tract of the liver advances to liver fibrosis, cirrhosis, and finally, end-stage liver disease [171]. The symptoms fluctuate across a broad spectrum, from asymptomatic, general symptoms (pruritus, fatigue), cholangitis, and inflammatory bowel disease (IBD), to hepatobiliary malignancies [171,172].

However, the treatment of patients with PSC is difficult, given the limited supporting data for management. Current treatment relies on individual symptom control ac-

cording to the patient's condition. The etiology affecting the intrahepatic and extrahepatic bile ducts is still unclear, even from various theories of pathogenesis and research [173]. The understanding of the pathogenesis of PSC is the first step to the development of effective therapies. New challenges for clarifying the pathogenesis and developing new target drugs are still under investigation.

4.1. Mechanism of Chronic Bile Duct Injury in the Bile Duct in PSC

The pathological lesion of PSC is “onion skin” scars, which appear as obliterated concentric periductal fibrosis in the lining of the bile duct cells leading to biliary stricture. How bile duct cells and immune cells (mostly T cells, neutrophils and macrophages) work together with hepatic stellate cells and portal myofibroblasts in this fibrillary production is important but unclear [174]; however, a major theory of pathogenesis is the exposure of genetically sensitive patients to environmental triggers causing bile duct injury [174,175].

4.1.1. Genetics of PSC

For the predisposing factor for PSC, genetic factors contribute approximately 10% and may be explained by sibling patients who have an enhanced risk of developing PSC [176,177]. According to GWASs, the genetic architecture of PSC shares features with both autoimmune diseases and IBD [176,178]. A total of 34–60% of patients with PSC have concurrent IBD in several Western and Asian populations [179–181]. In addition, in up to 25% of cases, patients with PSC may have other autoimmune diseases [181]; however, PSC predisposing genes show little evidence of an association with IBD and overlap with other autoimmune diseases, such as type 1 diabetes, celiac disease, rheumatoid arthritis, sarcoidosis, multiple sclerosis, and psoriasis [175,182]. This suggests that the genetic susceptibility to PSC extends into autoimmune pathophysiology beyond that represented by IBD.

Genes in the HLA class II region encode molecules that present extracellular sources antigen to CD8+ and CD4+ T lymphocytes, supporting the adaptive immune response in disease pathogenesis of PSC as an autoimmune disease. Polymorphisms in these genes are associated with most autoimmune diseases because they contribute to the specificity of immune responses [183–185].

More than 20 risk genes expressed by HLA haplotypes affect the development or decrease the risk of PSC.

4.1.2. Bile Acid Toxicity to Cholangiocytes and Hepatocytes

Cholangiocytes are exposed to hydrophobic bile acid in a physiological environment without cell toxicity [186]; however, hydrophobic bile acid-induced cell injury in various cell types, including hepatocytes, even at low concentrations [187]. The toxic effects of bile acids are explained by cholestasis, changes in the bile composition of disease progression in the bile ducts and colon, or impaired protective mechanisms. Bile acid is abundant in the gut, and it undergoes a bacteria-mediated transformation into bioactive molecules. Its metabolites control the host immune response by modulating the balance of Th17 and regulatory T cells (Tregs) [188]. In an observational study, conjugated primary bile acids and their derivatives were increased in patients with PSC compared with healthy controls; however, secondary bile acid did not differ between the groups [189].

The genetic variation in TGR5, promoting Cl^- and HCO_3^- secretion, induced down-regulation of the TGR5 protein in cholangiocytes of PSC patients [190]. The other gene variation stabilizing the apical cholangiocyte membrane also induced the impairment of the “biliary bicarbonate umbrella” in PSC [176]. The protective mechanism of hepatocytes against harmful bile acid accumulation is explained by FGF19, which is a negative feedback regulator of bile synthesis produced in the ileum after FXR activation by bile

acids [191]. Abnormal hepatic FGF19 expression was observed in the livers of PSC patients but not in healthy controls. It induced the pathologic accumulation of bile acids in the livers of PSC patients [191,192]. Defects in homeostasis for the regulation and control of bile acid leads to the chronic progression of fibrosing cholangiopathies and hepatopathy.

4.1.3. Fibrosis Development Related to Cholangiocyte and Hepatic Stellate Cell/Portal Myofibroblast

Cholangiocytes display an activated phenotype in the PSC by recognizing cytokines, hormones, and bile acids expressed by environmental damage, autoantigens, or the gut microbiome. Activated cholangiocytes are correlated with activated biliary tree stem cells that induce biliary fibrosis and progression of the bile duct [193,194]. TLR and nucleotide oligomerization domain-like receptors aid in detecting pathogens and activating bile duct cells, resulting in the secretion of proinflammatory cytokines [195]. IL-2 is a key factor in the regulation and programming of the immune system in PSC [196]. TNF- α , TGF β 1, IL-1 β , and IL-6, along with CD8+ and CD4+ T cells, cause myofibroblast activation and fibrosis. These induced the peribiliary fibrosis development and subsequent cirrhosis through interactions with HSCs [197,198].

Chronic injury can lead to cholangiocyte senescence and differentiation of matrix-depositing HSCs from myofibroblasts and portal fibroblasts, resulting in tissue scarring and bile duct strictures [199,200]. During chronic senescence, the surrounding tissues are susceptible to senescence associated secretory phenotype (SASP) related damage, resulting in persistent inflammatory and fibrosis responses. Moreover, destructive SASP not only maintains the inflammatory response, but can also activate the senescent phenotype in surrounding non-senescent cells [201].

4.1.4. Gut-Liver Immunity of PSC

The PSC is considered a part of the hepatobiliary manifestation of IBD, and gut-derived adaptive and innate immune responses contribute to chronic and progressive biliary inflammation. The liver biopsy with PSC showed predominant T-cell infiltration with portal inflammation. In addition, other neutrophils and macrophages have also been observed to release TGF β chronically and contribute to chronic inflammation, fibrosis, and cirrhosis [202]. In PSCs, hepatic inflammation by nutrition or bacterial inflow through the portal tract results in aberrant hepatic expression of the mucosal vascular address in cell adhesion molecule 1 (MADCAM1) and the C-C motif chemokine ligand 25 (CCL25), which results in the recruitment of mucosal T cells to the liver [175]. Other studies of recruited T cells in the portal area reported results regarding gut and liver relationships in PSC patients with or without IBD. The normal colon expresses the endothelial adhesion molecule MADCAM1 and the chemokine CCL25, which recruit mucosal lymphocytes with receptors for MADCAM1 and CCL25 (α 4 β 7 integrin and C-C chemokine receptor 9 (CCR9), respectively) during activation by gut dendritic cells [203,204].

The intestinal and biliary epithelia share many properties, including the expression of tight junction proteins such as E-cadherin, pattern recognition receptors (PRRs), and the ability to release secretory IgA [205,206]. The PRR, in terms of allowing cells into gut-derived bacterial products, was expressed in Kupffer cells, sinusoidal endothelial cells, and cholangiocytes; however, in PSC patients, genetic polymorphisms reduced the threshold of PRR signaling and changed the gut microbiota, leading to liver injury [206]. Another study suggested that cellular antigen(s), which are shared by the human colon and biliary epithelium by molecular mimicry, induced immune-mediated chronic inflammation [207].

4.1.5. Gut Microbial Profile in PSC

The microbiota has been considered a crucial factor for the pathogenesis of PSC in recent studies. The analysis of the gut microbiota in PSC compared with healthy controls or IBD patients has been described using 16S rRNA sequencing technologies with mucosal or fecal sources [195]. The composition of the gut microbial community is altered, with an overall reduction in bacterial diversity and altered abundance of certain bacteria compared with PSC/PSC-IBD vs. healthy controls and IBD in fecal microbiota [208–211]. Although diverse heterogeneity was observed, some bacterial taxa were consistently altered in the feces of patients with PSC compared to healthy controls. In particular, *Veillonella* was higher in the stool of PSC patients than in healthy controls in all the studies [212–214]. *Veillonella* discriminated PSC from healthy controls with an area under the receiver operator characteristic curve (AUC) of 0.64 [209]. They have genes that encode amine oxidases and are producers of primary amines that can act as vascular adhesion protein-1 (VAP-1) substrates, which are critical for effector cell recruitment to the liver [212,213]. An increase in the genus *Sphingomonas* expressing amine oxidase is associated with the abnormal return of intestinal lymphocytes to the liver as the basis of the gut-liver axis [212,215].

As with members of the Proteobacteria phylum, such as *E. coli*, other microbiomes, such as *Enterococcus*, *Streptococcus*, and *Lactobacillus*, are frequently enriched in PSCs [214–216]. The unit of the *Enterococcus* genus correlated with elevated levels of ALP, a marker of disease severity [211]. *Fusobacterium* was associated with intestinal inflammation severity, whereas *Enterococcus* was associated with biliary pathology [210]. Furthermore, a similar fecal microbiota composition (AUC 0.88) was analyzed in geographically different cohorts in Germany and Norway in PSC [208]. These studies presumed the possible utility of microbial components as prognostic and/or diagnostic markers in PSC.

In addition, the microbiota is being considered as a potential biological treatment option of PSC to ameliorate cholestasis and hepatic fibrosis, although with an in murine model. The PSC mice have reduced the abundance of *Prevotella copri*. The gavage *P. copri* daily in these mice induced the improvement of cholestasis in enterohepatic circulation caused by affecting bile acid level [217].

Recently, studies on the pathophysiology of PSC extended to integrative analysis between gut microbiota, gene expression, and immunologic response. Quraishi et al. insisted on microbial alteration and differential gene expression (colonic transcriptome) in PSC/IBD and ulcerative colitis patients and implicated the dysregulation of bile acid metabolism [216]. The relationship between the specific microbiome and the colonic transcriptome impacts metabolism, such as bile acid, bile salt, and fatty acid metabolism. Genomics, microbiota, and functional analysis will be considered future challenges for single microbes in disease pathogenesis; however, the microbiome is influenced by environmental factors such as diet, drugs, physical activity, hygiene, and race [214,218]. Several studies have attempted to reduce bias by controlling for antibiotic use or classifying geographic differences, but more multicenter and systematic studies are needed to control for confounding factors [210,211].

Other studies have focused on fungal dysbiosis in feces. The fungus *Exophiala*, a fungal genus of the *Herpotrichiellaceae* family, increased PSC and increased the fungal biodiversity and altered the composition [219]. Another case reported that *exophiala* increased liver cirrhosis through cholestasis and dilation of intrahepatic bile ducts [220].

4.2. Symptomatic Treatment and Biliary Complications of PSC

PSC patients are subject to several significant events throughout a fluctuating and highly variable disease course. The absence of robust data limits the standardization of treatment recommendations.

4.2.1. Pruritus

Pruritus is a concomitant manifestation of cholestatic disease. More than two-thirds of these patients experience itching during the course of PSC or PBC. Increased concentrations of bile salts, histamine, serotonin, progesterone metabolites, and endogenous opioids have been controversially discussed as potential pruritogens. The pathogenesis of cholestasis pruritus remains largely unknown [221]. Recently, a randomized, placebo-controlled FITCH trial clearly showed the beneficial effect of the PPAR agonist bezafibrate (400 mg/day) on moderate to severe cholestasis-associated pruritus in PSC and PBC [222]. As another cause, the imbalance between the μ -opioid receptors (MOR) and κ -opioid receptors (KOR) has been proposed to modulate the transmission and enhancement of pruritus signals in the central nervous system. In a randomized, double-blind study of nalfurafine hydrochloride, the KOR agonist nalfurafine hydrochloride (2.5 or 5 μ g daily) was effective without significant side effects in the treatment of intractable pruritus in patients with chronic liver disease [223]; however, endogenous opioid levels and MOR/KOR ligands did not differ between the levels of pruritus and nonpruritic patients with chronic liver disease and did not correlate with itch intensity [224]. The previously recommended first-line medical treatment is cholestyramine (4–16 g/day). This anion exchange resin has been reported to alleviate pruritus in several small uncontrolled case series [221]. Rifampicin, naltrexone, sertraline, and KOR agonists/MOR antagonists may be considered in the following steps; however, only a few well-designed, randomized, placebo-controlled trials and several cohort studies have evaluated these medications. Further studies to understand the correlation between pruritogen, medication and symptoms are needed in the future.

4.2.2. Bacterial Cholangitis and Dominant Stricture

The risk of bacterial cholangitis is increased after endoscopic or surgical manipulations (including liver biopsy) in patients with PSC, but cholangitis can also occur spontaneously. Bile cultures from patients with PSC show a broad spectrum of bacteria in patients with and without prior biliary intervention. Although cholangitis occurs frequently, symptoms can be atypical [222]. Empirical antibiotics are typically effective, and prophylactic antibiotics should be administered before and after biliary interventions. In more severe cases, hospitalization is necessary for intravenous treatment, including broad-spectrum antibiotics [225].

In a meta-analysis study, treatment with antibiotics (metronidazole, vancomycin, rifaximin and minocycline) for PSC patients was also associated with a significant reduction in ALP and total serum bilirubin levels regardless of cholangitis [213]. Long-term antibiotic treatment (vancomycin) has been focused upon as a management method for controlling cholangitis and modulating the intestinal microflora caused by the bacterial dysbiosis of PSCs [226]. To judge the effectiveness of antibiotic treatment beyond the treatment of cholangitis, a longer treatment duration and follow-up with a powered placebo-controlled studies are needed. Furthermore, an integrative analysis should be applied to properly compare the genetic profile, microbiome, immune system, and antibiotic therapy.

4.2.3. Dominant Stricture

Dominant stenosis (extrahepatic bile duct less than 1.5 mm or narrowing of less than 1 mm in the area within 2 cm of the main biliary junction) results in advanced focal stenosis in 40–58% of patients with PSC [227]. Patients with dominant stenosis may be asymptomatic and present with various clinical manifestations or with worsening liver function tests, abdominal pain, and/or cholangitis [228]. Dominant stenosis may also exacerbate symptoms of cholestasis, including worsening pruritus or developing cholangitis; therefore, endoscopic treatment with balloon dilatation or stent placement would be beneficial in dominant stenosis, however, a single retrospective observational study re-

ported that 45% of patients with biliary stenosis showed similar changes in biochemistry for 1 year after diagnosis, with or without dominant stenosis [229]. Therefore, it has been recommended that endoscopic treatment of dominant stenosis should be performed only in the presence of clinical or biochemical abnormalities [230]; however, a recent large-scale retrospective study reported that regular endoscopic retrograde cholangiopancreatography (ERCP) with endoscopic balloon dilatation has significant benefits in PSC patients with dominant stenosis, affecting asymptomatic survival and the incidence of recurrent cholangitis [231]. In the endoscopic treatment of this dominant stenosis, balloon expansion is preferred over stent insertion because the incidence of related side effects (pancreatitis, cholangitis, bacteremia) is low [232].

4.2.4. Cholangiocarcinoma

The prevalence of cholangiocarcinoma (CCA) ranges from 7 to 15% of PSC patients over the patient's lifetime, with half of the cases being diagnosed in the first year after diagnosis of PSC, probably due to the development of CCA-related symptoms [171]. The pathogenesis of CCA in PSC is inflammation-driven carcinogenesis concomitant with various genetic and epigenetic abnormalities, but the pathogenesis of CCA in PSC is still controversial [233]. CCA may remain asymptomatic for a long time, but when symptoms appear, and CCA is found, an advanced tumor stage is usually present. Unfortunately, most cases are unresectable, and no effective medications are available when CCA is diagnosed; therefore, early recognition of CCA in PSC remains a major challenge. The surveillance and management of PSC patients at an increased risk of developing CCA are important. Several factors increase the risk of CCA in PSC patients, including old age, abnormalities related to the large duct, male sex, comorbidity of ulcerative colitis, and infections with hepatitis B or hepatitis C. Magnetic resonance cholangiopancreatography (MRCP) combined with serum CA 19-9 should be used for screening and surveillance. In contrast, ERCP-based brush cytology, biopsy, and fluorescence in situ hybridization (FISH) should be used for diagnosis [233].

4.2.5. Gallbladder Carcinoma

Gallbladder disease, including gallstones or cholecystitis, is relatively common in 25% of PSC patients. A gallbladder mass lesion is found in 6–15% of PSC patients, and over half of those had adenocarcinoma after cholecystectomy [234,235]. Surveillance with regular ultrasonography of the gallbladder may be of value in these patients. Adenocarcinoma can also be found in polyps smaller than 5 mm; therefore, cholecystectomy should be considered regardless of the size of the polyp in PSC patients [236].

4.2.6. Liver Cirrhosis and Hepatocellular Carcinoma, and Liver Transplantation

Histological characteristics of PSC with an “onion-skin” pattern mimicking concentric periductal fibrosis with lymphocyte infiltration and portal edema progressed to portal and bridging fibrosis. Finally, liver cirrhosis with regenerative nodules and extensive fibrosis occurred [199,200]. LT is the only curative treatment and lifesaving intervention for PSC. Indications for LT are similar to other liver diseases transplanted, according to the eligible model for end-stage liver disease (MELD) scores in cirrhotic patients. Specific indications for LT in PSC patients include recurrent uncontrollable cholangitis, decompensated secondary biliary cirrhosis, and intractable pruritus [175]; however, a high rate of recurrent PSC (more than 20%), less than 5 years after LT was observed, negatively impacted patient survival [237]. Therefore, it is important to distinguish between recurrent PSC and post-transplant biliary strictures, which can occur even more often (36%) for various reasons (ischemia, infection, treatment induction).

4.2.7. Inflammatory Bowel Disease and Colorectal Cancer

In patients with PSC, IBD was observed in 60% of all patients. Among these PSC-IBD patients, colonic involvement is most often identified as ulcerative colitis. Patients with PSC-IBD have an increased risk of colorectal cancer and colorectal dysplasia, increased hepatic–pancreatic biliary cancer and mortality, and an earlier onset than patients with IBD alone [238]; therefore, PSC and PSC-IBD patients under regular surveillance have better outcomes. In patients with PSC with IBD, annual colonoscopy, including chromoendoscopy and histological sampling, is recommended regardless of the presence of symptoms of colitis [230]. In PSC patients with IBD, colonic dysplasia should be endoscopically resected, similar to management in other IBD patients. Proctocolectomy should be considered if high-grade dysplasia is discovered [230].

4.3. Combination Treatment and Predicting Therapies for PSC

4.3.1. UDCA

UDCA is considered the first-line therapy with an excellent safety profile when administered at moderate doses in the treatment of PBC (13–15 mg/kg/day), which leads to improved LT-free survival in all patients regardless of the disease stage and biochemical response [108,239]; however, its effectiveness as monotherapy is insufficient to prevent PSC progression in most patients. Nevertheless, discontinuation of UDCA has been shown to cause worsening of liver biochemistry and symptoms [240]. This justified the maintenance of UDCA treatment. To overcome the therapeutic limitations of UDCA, a study is also underway to evaluate whether there is improvement in biliary lesions and the safety evaluation of intra-arterial injection of stem cells using UDCA together [NCT03516006].

4.3.2. FXR Agonist

A poor prognosis is expected if there is no response to UDCA treatment; therefore, continuing research for drugs with a new mechanism of action is necessary. OCA acts as an FXR agonist and affects bile acid synthesis, inflammation, and liver fibrosis [190]. It has been approved as a combination therapy with UDCA when there is no response to UDCA treatment due to evidence of the improvement of biochemical profiles in a long-term phase III randomized study [108,123]. In a phase II study of single therapy OCA, there was a reduced serum ALP level in patients with PSC [241]; however, OCA-induced dose-dependent pruritus was a side effect in this study. Cilofexor, a non-steroidal FXR agonist, also improved cholestasis and fibrosis biochemical markers in a phase II trial. A phase III trial will evaluate the reduction in the risk of fibrosis progression among noncirrhotic adults with PSC. [NCT0389012] FXR agonists induce endogenous FGF19 synthesis and affect the proliferation of hepatobiliary malignancies [242]. In this respect, long-term use of FXR agonists in PSC should require the evaluation of concomitant hepatobiliary malignancies.

4.3.3. PPAR Agonist

Bezafibrate, an intranuclear receptor, acts as a ligand of PPAR α . PPAR α promotes the multidrug resistance 3 (MDR3) gene expression and increases P-glycoprotein levels in bile duct canaliculi, which, in turn, causes buffer damage to the bile duct by forming micelles with hydrophobic toxic bile acids [243]. Bezafibrate, which has antioxidative and anti-inflammatory effects, may reduce the damage to the bile duct epithelium seen in cholestatic diseases by inhibiting TNF- α and inducing superoxide dismutase expression [244]. In a small, randomized study, bezafibrate significantly improved the biochemistry profile in PSC patients and showed 64% efficacy. A phase III randomized trial on the effect of the biochemical profile and LT-free survival of UDCA, and the bezafi-

brate combination in PSC with persistent cholestasis, despite UDCA therapy, is ongoing [NCT04309773].

4.3.4. 3-Hydroxy-3-methylglutaryl Coenzyme A (HMG-CoA) Reductase Inhibitor

Statins acting by inhibiting HMG-CoA reductase reduce lipid levels, leading to the inhibition of cholesterol synthesis and lowering cholesterol levels in serum and bile [245]. The effect on PSC would be decreased cholesterol levels in serum and bile and improved inflammation, bile stone formation, and ameliorated cholestasis. In a recent population-based cohort of PSC patients with IBD, statins reduced the risk for death or liver transplantation by 50%. In this study, treatment with UDCA was not associated with reduced mortality [241]. A phase III randomized trial has been ongoing for 5 years to evaluate the long-term survival and efficacy of simvastatin in improving the biochemistry profile of PSC patients [NCT04133792].

4.3.5. Oral Vancomycin

Oral vancomycin, an antibiotic that is poorly absorbed into the body from the bowel and concentrates in the intestine, is also considered a treatment option for PSC. The dysbiosis of the gut microbiome affects the gut immune system, attracting white cells, which damage the bile ducts, increases the delivery of toxins to the liver through a leaky bowel wall, and directly causes damage through toxic bile acids [195,246]. From this point of view, oral vancomycin greatly modified intestinal bacteria, and in particular, caused a decrease in the number of intestinal bacteria of the *Bacteroides* and *Prevotella* species. Recently, one randomized controlled trial showed that oral vancomycin was well tolerated and was associated with improvements in liver chemistry [247]. A phase III study will evaluate the efficacy of oral vancomycin in treating biochemistry and fibrosis in PSC [NCT03710122].

In addition to these studies, various novel therapies are being conducted to inhibit the progression of PSC through chemokine receptor antagonists, FGF19 analogs or fecal microbiota transplantation (Table 3).

Table 3. Novel therapies currently under investigation in PSC.

Agent	Mechanism	Clinical Trial Stage	Main Results	Ref.
24-norursodeoxycholic acid (norUDCA)	Side chain-shortened C23 homolog of UDCA	Phase II (NUC-3) Phase III (NCT03872921)	• Reduced ALP levels by −12.3%, −17.3%, −26.0% in the 500, 1000, 1500 mg/d groups. • Data yet to be collected	[248]
Berberine ursodeoxycholate (BUDCA)	Ionic salt of two active moieties, berberine and UDCA	Phase II	• Not yet published	
Obeticholic acid	FXR agonist	Phase II (AESOP)	• Reduced serum ALP.	[249]
Cilofexor	FXR agonist	Phase III (PRIMIS)	• Data yet to be collected	
Vidofludimus calcium	FXR agonists + dihydroorotate dehydrogenase inhibitor	Phase II	• Normalization of ALP occurred in 27.7%.	[250]
Benzafibrate (+UDCA)	PPAR agonist	Phase III (BEZASCLER)	• Data yet to be collected	
Seladelpar	PPAR δ agonist	Phase II	• Not yet published	

(NCT04024813)			
Simvastatin	HMGCoA reductase inhibitors	Phase III (PiSCATIN)	Data yet to be collected
Vancomycin	Antibiotics	Phase III (NCT03710122)	Data yet to be collected
CM-101	Monoclonal Ab blocking CCL24	Phase II (SPRING)	Data yet to be collected
Cenicriviroc	Dual antagonist of CCR2 and CCR5	Phase II (PER-SEUS)	Median 18% reduction in ALP. [251]
Timolumab	Monoclonal anti-VAP-1 antibody	Phase II (BU-TEO)	Not yet published
NGM282	FGF19 analog	Phase II	Enhanced Liver Fibrosis score and inhibited bile acid synthesis without affecting ALP levels. [252]
Sulfasalazine	Aminosalicylates	Phase II (NCT02177136)	Data yet to be collected
Fecal Microbiota Transplantation	Restore the microbiome	Phase II	Thirty percent experienced a $\geq 50\%$ decrease in ALP. [253]
Umbilical Cord Mesenchymal Stem Cells	Repair of damaged tissue and immuno-modulation	Phase II (NCT03516006)	Data yet to be collected

UDCA: Ursodeoxycholic acid, PPAR: Peroxisome proliferator-activated receptor, ALP: alkaline phosphatase, FXR: farnesoid X receptor, HMG-CoA: β -hydroxy β -methylglutaryl-CoA, CCL: C-C motif chemokine ligand, CCR: C-C motif chemokine receptor.

Hitherto, we have covered the pathophysiological mechanisms, clinical features, and treatment of PBC and PSC. Figure 1 shows the differences between PBC and PSC.

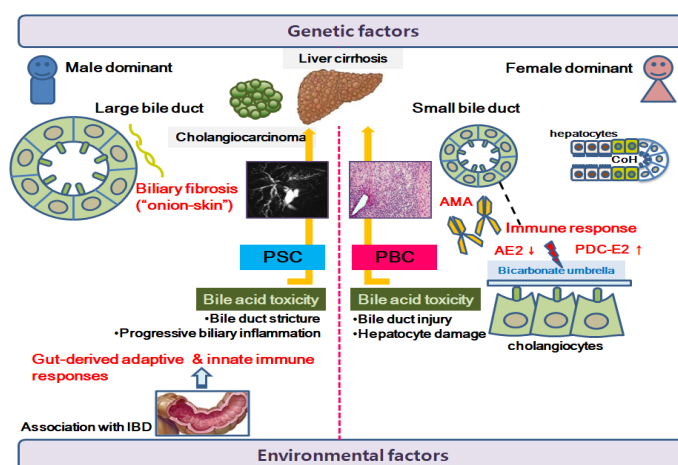


Figure 1. The comparison between PBC and PSC. This is a figure highlighting the differences between two diseases. The picture is adapted from Gidwaney et al [254] and Libre Pathology (<https://librepathology.org>, accessed on 28 April 2022). CoH: canals of Hering, PSC: primary sclerosing cholangitis, PBC: primary biliary cholangitis, IBD: inflammatory bowel disease, AMA: anti-mitochondrial antibody, AE2: anion exchanger 2, PDC-E2: pyruvate dehydrogenase complex, E2.

5. Conclusions

Despite extensive research to understand the molecular pathogenic features of PBC and PSC, there are still significant gaps between knowledge and its application to clinical practice. As medical treatments, UDCA and OCA have been approved for PBC; however, patients who are not responsive to these treatments remain a challenge. Moreover, medical therapy for PSC remains limited, with no effective or approved pharmacologic treatment identified. Fortunately, various therapeutics targeting pathogenic molecules in these cholestatic liver diseases are under investigation. Combination therapy composed of anti-biliary, anti-inflammatory, antioxidative, or anti-fibrotic treatment may be a viable solution. It is necessary to identify patients at high risk for disease progression or malignancy. Through continual research, we will improve treatment options for these complex and perplexing chronic progressive hepatobiliary diseases.

Author Contributions: Conceptualization, J.-W.P., J.-H.K. and D.J.K.; investigation, S.-E.K., J.H.J. and M.-K.J.; resources, H.-S.K., M.-S.L., K.T.S. and S.-H.P.; writing—original draft preparation, J.-W.P. and J.-H.K.; supervision, D.J.K. All authors have read and agreed to the published version of the manuscript.

Funding: This research was supported by the Basic Science Research Program through the National Research Foundation of Korea (NRF), funded by the Ministry of Education (2020R1A6A1A03043026).

Conflicts of Interest: The authors declare no conflict of interest.

References

- Alvaro, D.; Bragazzi, M.C.; Ridola, L. Inflammatory and neoplastic cholangiopathies. *Recenti Progress. Med.* **2018**, *109*, 595–599. <https://doi.org/10.1701/3082.30744>.
- Lazaridis, K.N.; Strazzabosco, M.; Larusso, N.F. The cholangiopathies: Disorders of biliary epithelia. *Gastroenterology* **2004**, *127*, 1565–1577. <https://doi.org/10.1053/j.gastro.2004.08.006>.
- Boonstra, K.; Beuers, U.; Ponsioen, C.Y. Epidemiology of primary sclerosing cholangitis and primary biliary cirrhosis: A systematic review. *J. Hepatol.* **2012**, *56*, 1181–1188. <https://doi.org/10.1016/j.jhep.2011.10.025>.
- Beuers, U.; Trauner, M.; Jansen, P.; Poupon, R. New paradigms in the treatment of hepatic cholestasis: From UDCA to FXR, PXR and beyond. *J. Hepatol.* **2015**, *62*, S25–S37. <https://doi.org/10.1016/j.jhep.2015.02.023>.
- Tanaka, A.; Takikawa, H.; Mochida, S.; Koike, K.; Miwa, H.; Shimosegawa, T. Changing Nomenclature for PBC from “Primary Biliary Cirrhosis” to “Primary Biliary Cholangitis”. *J. Gastroenterol.* **2016**, *51*, 748–749. <https://doi.org/10.1007/s00535-016-1223-z>.
- Juran, B.D.; Lazaridis, K.N. Environmental factors in primary biliary cirrhosis. *Semin. Liver Dis.* **2014**, *34*, 265–272. <https://doi.org/10.1055/s-0034-1383726>.
- Webb, G.J.; Hirschfield, G.M. Using GWAS to identify genetic predisposition in hepatic autoimmunity. *J. Autoimmun.* **2016**, *66*, 25–39. <https://doi.org/10.1016/j.jaut.2015.08.016>.
- Bianchi, I.; Carbone, M.; Lleo, A.; Invernizzi, P. Genetics and epigenetics of primary biliary cirrhosis. *Semin. Liver Dis.* **2014**, *34*, 255–264. <https://doi.org/10.1055/s-0034-1383725>.
- Molodecky, N.A.; Kareemi, H.; Parab, R.; Barkema, H.W.; Quan, H.; Myers, R.P.; Kaplan, G.G. Incidence of primary sclerosing cholangitis: A systematic review and meta-analysis. *Hepatology* **2011**, *53*, 1590–1599. <https://doi.org/10.1002/hep.24247>.
- Boonstra, K.; Weersma, R.K.; van Erpecum, K.J.; Rauws, E.A.; Spanier, B.W.; Poen, A.C.; van Nieuwkerk, K.M.; Drenth, J.P.; Witteman, B.J.; Tuynman, H.A.; et al. Population-based epidemiology, malignancy risk, and outcome of primary sclerosing cholangitis. *Hepatology* **2013**, *58*, 2045–2055. <https://doi.org/10.1002/hep.26565>.
- Sclear, S.N.; Little, E.; Levy, C. Current Concepts in Primary Biliary Cirrhosis and Primary Sclerosing Cholangitis. *Clin. Transl. Gastroenterol.* **2015**, *6*, e109. <https://doi.org/10.1038/ctg.2015.33>.
- Marchioni Beery, R.M.; Vaziri, H.; Forouhar, F. Primary Biliary Cirrhosis and Primary Sclerosing Cholangitis: A Review Featuring a Women’s Health Perspective. *J. Clin. Transl. Hepatol.* **2014**, *2*, 266–284. <https://doi.org/10.14218/JCTH.2014.00024>.
- Saxena, R.; Theise, N. Canals of Hering: Recent insights and current knowledge. *Semin. Liver Dis.* **2004**, *24*, 43–48. <https://doi.org/10.1055/s-2004-823100>.
- Boyer, J.L. Bile formation and secretion. *Compr. Physiol.* **2013**, *3*, 1035–1078. <https://doi.org/10.1002/cphy.c120027>.
- Banales, J.M.; Huebert, R.C.; Karlsen, T.; Strazzabosco, M.; LaRusso, N.F.; Gores, G.J. Cholangiocyte pathobiology. *Nat. Rev. Gastroenterol. Hepatol.* **2019**, *16*, 269–281. <https://doi.org/10.1038/s41575-019-0125-y>.
- Banales, J.M.; Prieto, J.; Medina, J.F. Cholangiocyte anion exchange and biliary bicarbonate excretion. *World J. Gastroenterol.* **2006**, *12*, 3496–3511. <https://doi.org/10.3748/wjg.v12.i22.3496>.
- Hundt, M.; Basit, H.; John, S. Physiology, Bile Secretion. In *StatPearls*; StatPearls Publishing: Treasure Island, FL, USA, 2022.

18. Han, Y.; Glaser, S.; Meng, F.; Francis, H.; Marzioni, M.; McDaniel, K.; Alvaro, D.; Venter, J.; Carpino, G.; Onori, P.; et al. Recent advances in the morphological and functional heterogeneity of the biliary epithelium. *Exp. Biol. Med.* **2013**, *238*, 549–565. <https://doi.org/10.1177/1535370213489926>.
19. Kanno, N.; LeSage, G.; Glaser, S.; Alvaro, D.; Alpini, G. Functional heterogeneity of the intrahepatic biliary epithelium. *Hepatology* **2000**, *31*, 555–561. <https://doi.org/10.1002/hep.510310302>.
20. Maroni, L.; Haibo, B.; Ray, D.; Zhou, T.; Wan, Y.; Meng, F.; Marzioni, M.; Alpini, G. Functional and structural features of cholangiocytes in health and disease. *Cell. Mol. Gastroenterol. Hepatol.* **2015**, *1*, 368–380. <https://doi.org/10.1016/j.jcmgh.2015.05.005>.
21. Franchitto, A.; Onori, P.; Renzi, A.; Carpino, G.; Mancinelli, R.; Alvaro, D.; Gaudio, E. Recent advances on the mechanisms regulating cholangiocyte proliferation and the significance of the neuroendocrine regulation of cholangiocyte pathophysiology. *Ann. Transl. Med.* **2013**, *1*, 27. <https://doi.org/10.3978/j.issn.2305-5839.2012.10.03>.
22. Sell, S. Heterogeneity and plasticity of hepatocyte lineage cells. *Hepatology* **2001**, *33*, 738–750. <https://doi.org/10.1053/jhep.2001.21900>.
23. Alvaro, D.; Mancino, M.G.; Glaser, S.; Gaudio, E.; Marzioni, M.; Francis, H.; Alpini, G. Proliferating cholangiocytes: A neuroendocrine compartment in the diseased liver. *Gastroenterology* **2007**, *132*, 415–431. <https://doi.org/10.1053/j.gastro.2006.07.023>.
24. Selmi, C.; Mayo, M.J.; Bach, N.; Ishibashi, H.; Invernizzi, P.; Gish, R.G.; Gordon, S.C.; Wright, H.I.; Zweiban, B.; Podda, M.; et al. Primary biliary cirrhosis in monozygotic and dizygotic twins: Genetics, epigenetics, and environment. *Gastroenterology* **2004**, *127*, 485–492. <https://doi.org/10.1053/j.gastro.2004.05.005>.
25. Lazaridis, K.N.; Juran, B.D.; Boe, G.M.; Slusser, J.P.; de Andrade, M.; Homburger, H.A.; Ghosh, K.; Dickson, E.R.; Lindor, K.D.; Petersen, G.M. Increased prevalence of antimitochondrial antibodies in first-degree relatives of patients with primary biliary cirrhosis. *Hepatology* **2007**, *46*, 785–792. <https://doi.org/10.1002/hep.21749>.
26. Gulamhusein, A.F.; Lazaridis, K.N. Primary biliary cholangitis, DNA, and beyond: The Relative contribution of genes. *Hepatology* **2018**, *68*, 19–21. <https://doi.org/10.1002/hep.29783>.
27. Mungall, A.J.; Palmer, S.A.; Sims, S.K.; Edwards, C.A.; Ashurst, J.L.; Wilming, L.; Jones, M.C.; Horton, R.; Hunt, S.E.; Scott, C.E.; et al. The DNA sequence and analysis of human chromosome 6. *Nature* **2003**, *425*, 805–811. <https://doi.org/10.1038/nature02055>.
28. Joshita, S.; Umemura, T.; Tanaka, E.; Ota, M. Genetics and epigenetics in the pathogenesis of primary biliary cholangitis. *Clin. J. Gastroenterol.* **2018**, *11*, 11–18. <https://doi.org/10.1007/s12328-017-0799-z>.
29. Donaldson, P.T.; Baragiotta, A.; Heneghan, M.A.; Floreani, A.; Venturi, C.; Underhill, J.A.; Jones, D.E.; James, O.F.; Bassendine, M.F. HLA class II alleles, genotypes, haplotypes, and amino acids in primary biliary cirrhosis: A large-scale study. *Hepatology* **2006**, *44*, 667–674. <https://doi.org/10.1002/hep.21316>.
30. Invernizzi, P.; Selmi, C.; Poli, F.; Frison, S.; Floreani, A.; Alvaro, D.; Almasio, P.; Rosina, F.; Marzioni, M.; Fabris, L.; et al. Human leukocyte antigen polymorphisms in Italian primary biliary cirrhosis: A multicenter study of 664 patients and 1992 healthy controls. *Hepatology* **2008**, *48*, 1906–1912. <https://doi.org/10.1002/hep.22567>.
31. Hirschfield, G.M.; Liu, X.; Xu, C.; Lu, Y.; Xie, G.; Lu, Y.; Gu, X.; Walker, E.J.; Jing, K.; Juran, B.D.; et al. Primary biliary cirrhosis associated with HLA, IL12A, and IL12RB2 variants. *N. Engl. J. Med.* **2009**, *360*, 2544–2555. <https://doi.org/10.1056/NEJMoa0810440>.
32. Liu, X.; Invernizzi, P.; Lu, Y.; Kosoy, R.; Lu, Y.; Bianchi, I.; Podda, M.; Xu, C.; Xie, G.; Macchiardi, F.; et al. Genome-wide meta-analyses identify three loci associated with primary biliary cirrhosis. *Nat. Genet.* **2010**, *42*, 658–660. <https://doi.org/10.1038/ng.627>.
33. Cordell, H.J.; Han, Y.; Mells, G.F.; Li, Y.; Hirschfield, G.M.; Greene, C.S.; Xie, G.; Juran, B.D.; Zhu, D.; Qian, D.C.; et al. International genome-wide meta-analysis identifies new primary biliary cirrhosis risk loci and targetable pathogenic pathways. *Nat. Commun.* **2015**, *6*, 8019. <https://doi.org/10.1038/ncomms9019>.
34. Poupon, R.; Ping, C.; Chretien, Y.; Corpechot, C.; Chazouilleres, O.; Simon, T.; Heath, S.C.; Matsuda, F.; Poupon, R.E.; Housset, C.; et al. Genetic factors of susceptibility and of severity in primary biliary cirrhosis. *J. Hepatol.* **2008**, *49*, 1038–1045. <https://doi.org/10.1016/j.jhep.2008.07.027>.
35. Li, Y.; Tang, R.; Ma, X. Epigenetics of Primary Biliary Cholangitis. *Adv. Exp. Med. Biol.* **2020**, *1253*, 259–283. https://doi.org/10.1007/978-981-15-3449-2_10.
36. Lleo, A.; Zhang, W.; Zhao, M.; Tan, Y.; Bernuzzi, F.; Zhu, B.; Liu, Q.; Tan, Q.; Malinverno, F.; Valenti, L.; et al. DNA methylation profiling of the X chromosome reveals an aberrant demethylation on CXCR3 promoter in primary biliary cirrhosis. *Clin. Epigenet.* **2015**, *7*, 61. <https://doi.org/10.1186/s13148-015-0098-9>.
37. Beuers, U.; Hohenester, S.; de Buy Wenniger, L.J.; Kremer, A.E.; Jansen, P.L.; Elferink, R.P. The biliary HCO₃⁽⁻⁾ umbrella: A unifying hypothesis on pathogenetic and therapeutic aspects of fibrosing cholangiopathies. *Hepatology* **2010**, *52*, 1489–1496. <https://doi.org/10.1002/hep.23810>.
38. Spirli, C.; Nathanson, M.H.; Fiorotto, R.; Duner, E.; Denson, L.A.; Sanz, J.M.; Di Virgilio, F.; Okolicsanyi, L.; Casagrande, F.; Strazzabosco, M. Proinflammatory cytokines inhibit secretion in rat bile duct epithelium. *Gastroenterology* **2001**, *121*, 156–169. <https://doi.org/10.1053/gast.2001.25516>.
39. Chang, J.C.; Go, S.; de Waart, D.R.; Munoz-Garrido, P.; Beuers, U.; Paulusma, C.C.; Oude Elferink, R. Soluble Adenylyl Cyclase Regulates Bile Salt-Induced Apoptosis in Human Cholangiocytes. *Hepatology* **2016**, *64*, 522–534. <https://doi.org/10.1002/hep.28550>.

40. Hohenester, S.; Wenniger, L.M.; Paulusma, C.C.; van Vliet, S.J.; Jefferson, D.M.; Elferink, R.P.; Beuers, U. A biliary HCO₃[−] umbrella constitutes a protective mechanism against bile acid-induced injury in human cholangiocytes. *Hepatology* **2012**, *55*, 173–183. <https://doi.org/10.1002/hep.24691>.
41. Sasaki, M.; Miyakoshi, M.; Sato, Y.; Nakanuma, Y. Increased expression of mitochondrial proteins associated with autophagy in biliary epithelial lesions in primary biliary cirrhosis. *Liver Int.* **2013**, *33*, 312–320. <https://doi.org/10.1111/liv.12049>.
42. Chang, J.C.; Beuers, U.; Oude Elferink, R.P. The Emerging Role of Soluble Adenylyl Cyclase in Primary Biliary Cholangitis. *Dig. Dis.* **2017**, *35*, 217–223. <https://doi.org/10.1159/000450914>.
43. Sasaki, M.; Nakanuma, Y. Bile Acids and Deregulated Cholangiocyte Autophagy in Primary Biliary Cholangitis. *Dig. Dis.* **2017**, *35*, 210–216. <https://doi.org/10.1159/000450913>.
44. Sasaki, M.; Yoshimura-Miyakoshi, M.; Sato, Y.; Nakanuma, Y. A possible involvement of endoplasmic reticulum stress in biliary epithelial autophagy and senescence in primary biliary cirrhosis. *J. Gastroenterol.* **2015**, *50*, 984–995. <https://doi.org/10.1007/s00535-014-1033-0>.
45. Sasaki, M.; Sato, Y.; Nakanuma, Y. An impaired biliary bicarbonate umbrella may be involved in dysregulated autophagy in primary biliary cholangitis. *Lab. Investig.* **2018**, *98*, 745–754. <https://doi.org/10.1038/s41374-018-0045-4>.
46. Hisamoto, S.; Shimoda, S.; Harada, K.; Iwasaka, S.; Onohara, S.; Chong, Y.; Nakamura, M.; Bekki, Y.; Yoshizumi, T.; Ikegami, T.; et al. Hydrophobic bile acids suppress expression of AE2 in biliary epithelial cells and induce bile duct inflammation in primary biliary cholangitis. *J. Autoimmun.* **2016**, *75*, 150–160. <https://doi.org/10.1016/j.jaut.2016.08.006>.
47. Sasaki, M.; Miyakoshi, M.; Sato, Y.; Nakanuma, Y. Autophagy mediates the process of cellular senescence characterizing bile duct damages in primary biliary cirrhosis. *Lab. Investig.* **2010**, *90*, 835–843. <https://doi.org/10.1038/labinvest.2010.56>.
48. Sasaki, M.; Miyakoshi, M.; Sato, Y.; Nakanuma, Y. Modulation of the microenvironment by senescent biliary epithelial cells may be involved in the pathogenesis of primary biliary cirrhosis. *J. Hepatol.* **2010**, *53*, 318–325. <https://doi.org/10.1016/j.jhep.2010.03.008>.
49. Tabibian, J.H.; O'Hara, S.P.; Splinter, P.L.; Trussoni, C.E.; LaRusso, N.F. Cholangiocyte senescence by way of N-ras activation is a characteristic of primary sclerosing cholangitis. *Hepatology* **2014**, *59*, 2263–2275. <https://doi.org/10.1002/hep.26993>.
50. Sasaki, M.; Miyakoshi, M.; Sato, Y.; Nakanuma, Y. Chemokine-chemokine receptor CCL2-CCR2 and CX3CL1-CX3CR1 axis may play a role in the aggravated inflammation in primary biliary cirrhosis. *Dig. Dis. Sci.* **2014**, *59*, 358–364. <https://doi.org/10.1007/s10620-013-2920-6>.
51. Selmi, C.; Bowlus, C.L.; Gershwin, M.E.; Coppel, R.L. Primary biliary cirrhosis. *Lancet* **2011**, *377*, 1600–1609. [https://doi.org/10.1016/S0140-6736\(10\)61965-4](https://doi.org/10.1016/S0140-6736(10)61965-4).
52. Bogdanos, D.P.; Pares, A.; Baum, H.; Caballeria, L.; Rigopoulou, E.I.; Ma, Y.; Burroughs, A.K.; Rodes, J.; Vergani, D. Disease-specific cross-reactivity between mimicking peptides of heat shock protein of *Mycobacterium gordonae* and dominant epitope of E2 subunit of pyruvate dehydrogenase is common in Spanish but not British patients with primary biliary cirrhosis. *J. Autoimmun.* **2004**, *22*, 353–362. <https://doi.org/10.1016/j.jaut.2004.03.002>.
53. Arbour, L.; Rupps, R.; Field, L.; Ross, P.; Erikson, A.; Henderson, H.; Hill, W.; Yoshida, E. Characteristics of primary biliary cirrhosis in British Columbia's First Nations population. *Can. J. Gastroenterol.* **2005**, *19*, 305–310. <https://doi.org/10.1155/2005/203028>.
54. Selmi, C.; Gershwin, M.E. Chronic Autoimmune Epithelitis in Sjogren's Syndrome and Primary Biliary Cholangitis: A Comprehensive Review. *Rheumatol. Ther.* **2017**, *4*, 263–279. <https://doi.org/10.1007/s40744-017-0074-2>.
55. Lleo, A.; Maroni, L.; Glaser, S.; Alpini, G.; Marziani, M. Role of cholangiocytes in primary biliary cirrhosis. *Semin. Liver Dis.* **2014**, *34*, 273–284. <https://doi.org/10.1055/s-0034-1383727>.
56. Lleo, A.; Leung, P.S.C.; Hirschfield, G.M.; Gershwin, E.M. The Pathogenesis of Primary Biliary Cholangitis: A Comprehensive Review. *Semin. Liver Dis.* **2020**, *40*, 34–48. <https://doi.org/10.1055/s-0039-1697617>.
57. Dahlqvist, G.; Gaouar, F.; Carrat, F.; Meurisse, S.; Chazouilleres, O.; Poupon, R.; Johanet, C.; Corpechot, C.; French Network of Immunology Laboratories. Large-scale characterization study of patients with antimitochondrial antibodies but nonestablished primary biliary cholangitis. *Hepatology* **2017**, *65*, 152–163. <https://doi.org/10.1002/hep.28859>.
58. Yamagiwa, S.; Kamimura, H.; Takamura, M.; Aoyagi, Y. Autoantibodies in primary biliary cirrhosis: Recent progress in research on the pathogenetic and clinical significance. *World J. Gastroenterol.* **2014**, *20*, 2606–2612. <https://doi.org/10.3748/wjg.v20.i10.2606>.
59. Joshi, S.; Cauch-Dudek, K.; Heathcote, E.J.; Lindor, K.; Jorgensen, R.; Klein, R. Antimitochondrial antibody profiles: Are they valid prognostic indicators in primary biliary cirrhosis? *Am. J. Gastroenterol.* **2002**, *97*, 999–1002. <https://doi.org/10.1111/j.1572-0241.2002.05620.x>.
60. Tang, L.; Zhong, R.; He, X.; Wang, W.; Liu, J.; Zhu, Y.; Li, Y.; Hou, J. Evidence for the association between IgG-antimitochondrial antibody and biochemical response to ursodeoxycholic acid treatment in primary biliary cholangitis. *J. Gastroenterol. Hepatol.* **2017**, *32*, 659–666. <https://doi.org/10.1111/jgh.13534>.
61. Zandanell, S.; Strasser, M.; Feldman, A.; Streibinger, G.; Aigner, G.; Niederseer, D.; Laimer, M.; Mussnig, B.; Paulweber, B.; Datz, C.; et al. Similar clinical outcome of AMA immunoblot-M2-negative compared to immunoblot-positive subjects over six years of follow-up. *Postgrad. Med.* **2021**, *133*, 291–298. <https://doi.org/10.1080/00325481.2021.1885945>.
62. Muratori, P.; Muratori, L.; Gershwin, M.E.; Czaja, A.J.; Pappas, G.; MacCariello, S.; Granito, A.; Cassani, F.; Loria, P.; Lenzi, M.; et al. 'True' antimitochondrial antibody-negative primary biliary cirrhosis, low sensitivity of the routine assays, or both? *Clin. Exp. Immunol.* **2004**, *135*, 154–158. <https://doi.org/10.1111/j.1365-2249.2004.02332.x>.

63. Granito, A.; Yang, W.H.; Muratori, L.; Lim, M.J.; Nakajima, A.; Ferri, S.; Pappas, G.; Quarneti, C.; Bianchi, F.B.; Bloch, D.B.; et al. PML nuclear body component Sp140 is a novel autoantigen in primary biliary cirrhosis. *Am. J. Gastroenterol.* **2010**, *105*, 125–131. <https://doi.org/10.1038/ajg.2009.596>.
64. Hirschfield, G.M.; Gershwin, M.E. The immunobiology and pathophysiology of primary biliary cirrhosis. *Annu. Rev. Pathol.* **2013**, *8*, 303–330. <https://doi.org/10.1146/annurev-pathol-020712-164014>.
65. Kita, H.; Lian, Z.X.; Van de Water, J.; He, X.S.; Matsumura, S.; Kaplan, M.; Luketic, V.; Coppel, R.L.; Ansari, A.A.; Gershwin, M.E. Identification of HLA-A2-restricted CD8(+) cytotoxic T cell responses in primary biliary cirrhosis: T cell activation is augmented by immune complexes cross-presented by dendritic cells. *J. Exp. Med.* **2002**, *195*, 113–123. <https://doi.org/10.1084/jem.20010956>.
66. Lleo, A.; Shimoda, S.; Ishibashi, H.; Gershwin, M.E. Primary biliary cirrhosis and autoimmune hepatitis: Apotopes and epitopes. *J. Gastroenterol.* **2011**, *46* (Suppl. S1), 29–38. <https://doi.org/10.1007/s00535-010-0303-8>.
67. Lleo, A.; Bowlus, C.L.; Yang, G.X.; Invernizzi, P.; Podda, M.; Van de Water, J.; Ansari, A.A.; Coppel, R.L.; Worman, H.J.; Gores, G.J.; et al. Biliary apotopes and anti-mitochondrial antibodies activate innate immune responses in primary biliary cirrhosis. *Hepatology* **2010**, *52*, 987–998. <https://doi.org/10.1002/hep.23783>.
68. Lleo, A.; Selmi, C.; Invernizzi, P.; Podda, M.; Coppel, R.L.; Mackay, I.R.; Gores, G.J.; Ansari, A.A.; Van de Water, J.; Gershwin, M.E. Apotopes and the biliary specificity of primary biliary cirrhosis. *Hepatology* **2009**, *49*, 871–879. <https://doi.org/10.1002/hep.22736>.
69. Schrupf, E.; Tan, C.; Karlsen, T.H.; Sponheim, J.; Bjorkstrom, N.K.; Sundnes, O.; Alfsnes, K.; Kaser, A.; Jefferson, D.M.; Ueno, Y.; et al. The biliary epithelium presents antigens to and activates natural killer T cells. *Hepatology* **2015**, *62*, 1249–1259. <https://doi.org/10.1002/hep.27840>.
70. Shimoda, S.; Harada, K.; Niino, H.; Yoshizumi, T.; Soejima, Y.; Taketomi, A.; Maehara, Y.; Tsuneyama, K.; Nakamura, M.; Komori, A.; et al. Biliary epithelial cells and primary biliary cirrhosis: The role of liver-infiltrating mononuclear cells. *Hepatology* **2008**, *47*, 958–965. <https://doi.org/10.1002/hep.22102>.
71. Kita, H.; Matsumura, S.; He, X.S.; Ansari, A.A.; Lian, Z.X.; Van de Water, J.; Coppel, R.L.; Kaplan, M.M.; Gershwin, M.E. Quantitative and functional analysis of PDC-E2-specific autoreactive cytotoxic T lymphocytes in primary biliary cirrhosis. *J. Clin. Investig.* **2002**, *109*, 1231–1240. <https://doi.org/10.1172/JCI14698>.
72. Yan, J.; Harvey, B.P.; Gee, R.J.; Shlomchik, M.J.; Mamula, M.J. B cells drive early T cell autoimmunity in vivo prior to dendritic cell-mediated autoantigen presentation. *J. Immunol.* **2006**, *177*, 4481–4487. <https://doi.org/10.4049/jimmunol.177.7.4481>.
73. Vinuesa, C.G.; Tangye, S.G.; Moser, B.; Mackay, C.R. Follicular B helper T cells in antibody responses and autoimmunity. *Nat. Rev. Immunol.* **2005**, *5*, 853–865. <https://doi.org/10.1038/nri1714>.
74. Yang, C.Y.; Ma, X.; Tsuneyama, K.; Huang, S.; Takahashi, T.; Chalasani, N.P.; Bowlus, C.L.; Yang, G.X.; Leung, P.S.; Ansari, A.A.; et al. IL-12/Th1 and IL-23/Th17 biliary microenvironment in primary biliary cirrhosis: Implications for therapy. *Hepatology* **2014**, *59*, 1944–1953. <https://doi.org/10.1002/hep.26979>.
75. Qian, C.; Jiang, T.; Zhang, W.; Ren, C.; Wang, Q.; Qin, Q.; Chen, J.; Deng, A.; Zhong, R. Increased IL-23 and IL-17 expression by peripheral blood cells of patients with primary biliary cirrhosis. *Cytokine* **2013**, *64*, 172–180. <https://doi.org/10.1016/j.cyto.2013.07.005>.
76. Tan, Z.; Qian, X.; Jiang, R.; Liu, Q.; Wang, Y.; Chen, C.; Wang, X.; Ryffel, B.; Sun, B. IL-17A plays a critical role in the pathogenesis of liver fibrosis through hepatic stellate cell activation. *J. Immunol.* **2013**, *191*, 1835–1844. <https://doi.org/10.4049/jimmunol.1203013>.
77. Bottcher, K.; Rombouts, K.; Saffioti, F.; Roccarina, D.; Rosselli, M.; Hall, A.; Luong, T.; Tsochatzis, E.A.; Thorburn, D.; Pinzani, M. MAIT cells are chronically activated in patients with autoimmune liver disease and promote profibrogenic hepatic stellate cell activation. *Hepatology* **2018**, *68*, 172–186. <https://doi.org/10.1002/hep.29782>.
78. Jiang, X.; Lian, M.; Li, Y.; Zhang, W.; Wang, Q.; Wei, Y.; Zhang, J.; Chen, W.; Xiao, X.; Miao, Q.; et al. The immunobiology of mucosal-associated invariant T cell (MAIT) function in primary biliary cholangitis: Regulation by cholic acid-induced Interleukin-7. *J. Autoimmun.* **2018**, *90*, 64–75. <https://doi.org/10.1016/j.jaut.2018.01.007>.
79. Setsu, T.; Yamagiwa, S.; Tominaga, K.; Kimura, N.; Honda, H.; Kamimura, H.; Tsuchiya, A.; Takamura, M.; Terai, S. Persistent reduction of mucosal-associated invariant T cells in primary biliary cholangitis. *J. Gastroenterol. Hepatol.* **2018**, *33*, 1286–1294. <https://doi.org/10.1111/jgh.14076>.
80. Fan, Y.; Pedersen, O. Gut microbiota in human metabolic health and disease. *Nat. Rev. Microbiol.* **2021**, *19*, 55–71. <https://doi.org/10.1038/s41579-020-0433-9>.
81. Brussow, H. Microbiota and the human nature: Know thyself. *Environ. Microbiol.* **2015**, *17*, 10–15. <https://doi.org/10.1111/1462-2920.12693>.
82. Milosevic, I.; Vujovic, A.; Barac, A.; Djelic, M.; Korac, M.; Radovanovic Spurnic, A.; Gmizic, I.; Stevanovic, O.; Djordjevic, V.; Lekic, N.; et al. Gut-Liver Axis, Gut Microbiota, and Its Modulation in the Management of Liver Diseases: A Review of the Literature. *Int. J. Mol. Sci.* **2019**, *20*, 395. <https://doi.org/10.3390/ijms20020395>.
83. Trebicka, J.; Macnaughtan, J.; Schnabl, B.; Shawcross, D.L.; Bajaj, J.S. The microbiota in cirrhosis and its role in hepatic decompensation. *J. Hepatol.* **2021**, *75* (Suppl. S1), S67–S81. <https://doi.org/10.1016/j.jhep.2020.11.013>.
84. Liu Chen Kiow, J.; Vincent, C.; Sidani, S.; Bouin, M. High occurrence of small intestinal bacterial overgrowth in primary biliary cholangitis. *Neurogastroenterol. Motil.* **2019**, *31*, e13691. <https://doi.org/10.1111/nmo.13691>.

85. Dyson, J.K.; Hirschfield, G.M.; Adams, D.H.; Beuers, U.; Mann, D.A.; Lindor, K.D.; Jones, D.E. Novel therapeutic targets in primary biliary cirrhosis. *Nat. Rev. Gastroenterol. Hepatol.* **2015**, *12*, 147–158. <https://doi.org/10.1038/nrgastro.2015.12>.
86. Albillos, A.; de Gottardi, A.; Rescigno, M. The gut-liver axis in liver disease: Pathophysiological basis for therapy. *J. Hepatol.* **2020**, *72*, 558–577. <https://doi.org/10.1016/j.jhep.2019.10.003>.
87. Shimoda, S.; Harada, K.; Niino, H.; Taketomi, A.; Maehara, Y.; Tsuneyama, K.; Kikuchi, K.; Nakanuma, Y.; Mackay, I.R.; Gershwin, M.E.; et al. CX3CL1 (fractalkine): A signpost for biliary inflammation in primary biliary cirrhosis. *Hepatology* **2010**, *51*, 567–575. <https://doi.org/10.1002/hep.23318>.
88. Harada, K.; Isse, K.; Nakanuma, Y. Interferon γ accelerates NF- κ B activation of biliary epithelial cells induced by Toll-like receptor and ligand interaction. *J. Clin. Pathol.* **2006**, *59*, 184–190. <https://doi.org/10.1136/jcp.2004.023507>.
89. Atif, M.; Warner, S.; Oo, Y.H. Linking the gut and liver: Crosstalk between regulatory T cells and mucosa-associated invariant T cells. *Hepatol. Int.* **2018**, *12*, 305–314. <https://doi.org/10.1007/s12072-018-9882-x>.
90. Jeffery, H.C.; van Wilgenburg, B.; Kurioka, A.; Parekh, K.; Stirling, K.; Roberts, S.; Dutton, E.E.; Hunter, S.; Geh, D.; Braitch, M.K.; et al. Biliary epithelium and liver B cells exposed to bacteria activate intrahepatic MAIT cells through MR1. *J. Hepatol.* **2016**, *64*, 1118–1127. <https://doi.org/10.1016/j.jhep.2015.12.017>.
91. Wiest, R.; Albillos, A.; Trauner, M.; Bajaj, J.S.; Jalan, R. Targeting the gut-liver axis in liver disease. *J. Hepatol.* **2017**, *67*, 1084–1103. <https://doi.org/10.1016/j.jhep.2017.05.007>.
92. Inamine, T.; Schnabl, B. Immunoglobulin A and liver diseases. *J. Gastroenterol.* **2018**, *53*, 691–700. <https://doi.org/10.1007/s00535-017-1400-8>.
93. Floreani, A.; Baragiotta, A.; Pizzuti, D.; Martines, D.; Cecchetto, A.; Chiarelli, S. Mucosal IgA defect in primary biliary cirrhosis. *Am. J. Gastroenterol.* **2002**, *97*, 508–510. <https://doi.org/10.1111/j.1572-0241.2002.05521.x>.
94. Corpechot, C.; Chazouilleres, O.; Rousseau, A.; Le Gruyer, A.; Habersetzer, F.; Mathurin, P.; Gorla, O.; Potier, P.; Minello, A.; Silvain, C.; et al. A Placebo-Controlled Trial of Bezafibrate in Primary Biliary Cholangitis. *N. Engl. J. Med.* **2018**, *378*, 2171–2181. <https://doi.org/10.1056/NEJMoa1714519>.
95. Campisi, L.; Barbet, G.; Ding, Y.; Esplugues, E.; Flavell, R.A.; Blander, J.M. Apoptosis in response to microbial infection induces autoreactive TH17 cells. *Nat. Immunol.* **2016**, *17*, 1084–1092. <https://doi.org/10.1038/ni.3512>.
96. Haruta, I.; Kikuchi, K.; Hashimoto, E.; Nakamura, M.; Miyakawa, H.; Hirota, K.; Shibata, N.; Kato, H.; Arimura, Y.; Kato, Y.; et al. Long-term bacterial exposure can trigger nonsuppurative destructive cholangitis associated with multifocal epithelial inflammation. *Lab. Invest.* **2010**, *90*, 577–588. <https://doi.org/10.1038/labinvest.2010.40>.
97. Tang, R.; Wei, Y.; Li, Y.; Chen, W.; Chen, H.; Wang, Q.; Yang, F.; Miao, Q.; Xiao, X.; Zhang, H.; et al. Gut microbial profile is altered in primary biliary cholangitis and partially restored after UDCA therapy. *Gut* **2018**, *67*, 534–541. <https://doi.org/10.1136/gutjnl-2016-313332>.
98. Lemoine, S.; Marteau, P. Gut microbial profile in primary biliary cholangitis: Towards bioindicators. *Clin. Res. Hepatol. Gastroenterol.* **2017**, *41*, 507–508. <https://doi.org/10.1016/j.clinre.2017.06.002>.
99. Lv, L.X.; Fang, D.Q.; Shi, D.; Chen, D.Y.; Yan, R.; Zhu, Y.X.; Chen, Y.F.; Shao, L.; Guo, F.F.; Wu, W.R.; et al. Alterations and correlations of the gut microbiome, metabolism and immunity in patients with primary biliary cirrhosis. *Environ. Microbiol.* **2016**, *18*, 2272–2286. <https://doi.org/10.1111/1462-2920.13401>.
100. Wahlstrom, A.; Sayin, S.I.; Marschall, H.U.; Backhed, F. Intestinal Crosstalk between Bile Acids and Microbiota and Its Impact on Host Metabolism. *Cell Metab.* **2016**, *24*, 41–50. <https://doi.org/10.1016/j.cmet.2016.05.005>.
101. Ridlon, J.M.; Kang, D.J.; Hylemon, P.B.; Bajaj, J.S. Bile acids and the gut microbiome. *Curr. Opin. Gastroenterol.* **2014**, *30*, 332–338. <https://doi.org/10.1097/MOG.0000000000000057>.
102. European Association for the Study of the Liver. EASL Clinical Practice Guidelines: The Diagnosis and Management of Patients with Primary Biliary Cholangitis. *J. Hepatol.* **2017**, *67*, 145–172. <https://doi.org/10.1016/j.jhep.2017.03.022>.
103. Lazaridis, K.N.; Gores, G.J.; Lindor, K.D. Ursodeoxycholic acid ‘mechanisms of action and clinical use in hepatobiliary disorders’. *J. Hepatol.* **2001**, *35*, 134–146. [https://doi.org/10.1016/s0168-8278\(01\)00092-7](https://doi.org/10.1016/s0168-8278(01)00092-7).
104. Hofmann, A.F. Pharmacology of ursodeoxycholic acid, an enterohepatic drug. *Scand. J. Gastroenterol.* **1994**, *29* (Suppl. S204), 1–15. <https://doi.org/10.3109/00365529409103618>.
105. Trauner, M.; Graziadei, I.W. Review article: Mechanisms of action and therapeutic applications of ursodeoxycholic acid in chronic liver diseases. *Aliment. Pharmacol. Ther.* **1999**, *13*, 979–996. <https://doi.org/10.1046/j.1365-2036.1999.00596.x>.
106. You, H.; Ma, X.; Efe, C.; Wang, G.; Jeong, S.H.; Abe, K.; Duan, W.; Chen, S.; Kong, Y.; Zhang, D.; et al. APASL clinical practice guidance: The diagnosis and management of patients with primary biliary cholangitis. *Hepatol. Int.* **2022**, *16*, 1–23. <https://doi.org/10.1007/s12072-021-10276-6>.
107. Carbone, M.; Nardi, A.; Flack, S.; Carpino, G.; Varvaropoulou, N.; Gavrilu, C.; Spicer, A.; Badrock, J.; Bernuzzi, F.; Cardinale, V.; et al. Pretreatment prediction of response to ursodeoxycholic acid in primary biliary cholangitis: Development and validation of the UDCA Response Score. *Lancet Gastroenterol. Hepatol.* **2018**, *3*, 626–634. [https://doi.org/10.1016/S2468-1253\(18\)30163-8](https://doi.org/10.1016/S2468-1253(18)30163-8).
108. Lammers, W.J.; van Buuren, H.R.; Hirschfield, G.M.; Janssen, H.L.; Invernizzi, P.; Mason, A.L.; Ponsioen, C.Y.; Floreani, A.; Corpechot, C.; Mayo, M.J.; et al. 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. <https://doi.org/10.1053/j.gastro.2014.08.029>.
109. Pares, A.; Caballeria, L.; Rodes, J. Excellent long-term survival in patients with primary biliary cirrhosis and biochemical response to ursodeoxycholic Acid. *Gastroenterology* **2006**, *130*, 715–720. <https://doi.org/10.1053/j.gastro.2005.12.029>.

110. Angulo, P.; Lindor, K.D.; Therneau, T.M.; Jorgensen, R.A.; Malinchoc, M.; Kamath, P.S.; Dickson, E.R. Utilization of the Mayo risk score in patients with primary biliary cirrhosis receiving ursodeoxycholic acid. *Liver* **1999**, *19*, 115–121. <https://doi.org/10.1111/j.1478-3231.1999.tb00020.x>.
111. Corpechot, C.; Abenavoli, L.; Rabahi, N.; Chretien, Y.; Andreani, T.; Johanet, C.; Chazouilleres, O.; Poupon, R. Biochemical response to ursodeoxycholic acid and long-term prognosis in primary biliary cirrhosis. *Hepatology* **2008**, *48*, 871–877. <https://doi.org/10.1002/hep.22428>.
112. Kuiper, E.M.; Hansen, B.E.; de Vries, R.A.; den Ouden-Muller, J.W.; van Ditzhuijsen, T.J.; Haagsma, E.B.; Houben, M.H.; Witteman, B.J.; van Erpecum, K.J.; van Buuren, H.R.; et al. Improved prognosis of patients with primary biliary cirrhosis that have a biochemical response to ursodeoxycholic acid. *Gastroenterology* **2009**, *136*, 1281–1287. <https://doi.org/10.1053/j.gastro.2009.01.003>.
113. Azemoto, N.; Abe, M.; Murata, Y.; Hiasa, Y.; Hamada, M.; Matsuura, B.; Onji, M. Early biochemical response to ursodeoxycholic acid predicts symptom development in patients with asymptomatic primary biliary cirrhosis. *J. Gastroenterol.* **2009**, *44*, 630–634. <https://doi.org/10.1007/s00535-009-0051-9>.
114. Kumagi, T.; Guindi, M.; Fischer, S.E.; Arenovich, T.; Abdalian, R.; Coltescu, C.; Heathcote, E.J.; Hirschfield, G.M. Baseline ductopenia and treatment response predict long-term histological progression in primary biliary cirrhosis. *Am. J. Gastroenterol.* **2010**, *105*, 2186–2194. <https://doi.org/10.1038/ajg.2010.216>.
115. Corpechot, C.; Chazouilleres, O.; Poupon, R. Early primary biliary cirrhosis: Biochemical response to treatment and prediction of long-term outcome. *J. Hepatol.* **2011**, *55*, 1361–1367. <https://doi.org/10.1016/j.jhep.2011.02.031>.
116. Sepe, V.; Distrutti, E.; Fiorucci, S.; Zampella, A. Farnesoid X receptor modulators 2014–present: A patent review. *Expert Opin. Ther. Pat.* **2018**, *28*, 351–364. <https://doi.org/10.1080/13543776.2018.1459569>.
117. Pellicciari, R.; Fiorucci, S.; Camaioni, E.; Clerici, C.; Costantino, G.; Maloney, P.R.; Morelli, A.; Parks, D.J.; Willson, T.M. 6 α -ethyl-chenodeoxycholic acid (6-ECDCA), a potent and selective FXR agonist endowed with anticholestatic activity. *J. Med. Chem.* **2002**, *45*, 3569–3572. <https://doi.org/10.1021/jm025529g>.
118. Inagaki, T.; Choi, M.; Moschetta, A.; Peng, L.; Cummins, C.L.; McDonald, J.G.; Luo, G.; Jones, S.A.; Goodwin, B.; Richardson, J.A.; et al. Fibroblast growth factor 15 functions as an enterohepatic signal to regulate bile acid homeostasis. *Cell Metab.* **2005**, *2*, 217–225. <https://doi.org/10.1016/j.cmet.2005.09.001>.
119. Kim, I.; Ahn, S.H.; Inagaki, T.; Choi, M.; Ito, S.; Guo, G.L.; Kliewer, S.A.; Gonzalez, F.J. Differential regulation of bile acid homeostasis by the farnesoid X receptor in liver and intestine. *J. Lipid Res.* **2007**, *48*, 2664–2672. <https://doi.org/10.1194/jlr.M700330-JLR200>.
120. Triantis, V.; Saeland, E.; Bijl, N.; Oude-Elferink, R.P.; Jansen, P.L. Glycosylation of fibroblast growth factor receptor 4 is a key regulator of fibroblast growth factor 19-mediated down-regulation of cytochrome P450 7A1. *Hepatology* **2010**, *52*, 656–666. <https://doi.org/10.1002/hep.23708>.
121. Ananthanarayanan, M.; Balasubramanian, N.; Makishima, M.; Mangelsdorf, D.J.; Suchy, F.J. Human bile salt export pump promoter is transactivated by the farnesoid X receptor/bile acid receptor. *J. Biol. Chem.* **2001**, *276*, 28857–28865. <https://doi.org/10.1074/jbc.M011610200>.
122. Kast, H.R.; Goodwin, B.; Tarr, P.T.; Jones, S.A.; Anisfeld, A.M.; Stoltz, C.M.; Tontonoz, P.; Kliewer, S.; Willson, T.M.; Edwards, P.A. Regulation of multidrug resistance-associated protein 2 (ABCC2) by the nuclear receptors pregnane X receptor, farnesoid X-activated receptor, and constitutive androstane receptor. *J. Biol. Chem.* **2002**, *277*, 2908–2915. <https://doi.org/10.1074/jbc.M109326200>.
123. Nevens, F.; Andreone, P.; Mazzella, G.; Strasser, S.I.; Bowlus, C.; Invernizzi, P.; Drenth, J.P.; Pockros, P.J.; Regula, J.; Beuers, U.; et al. A Placebo-Controlled Trial of Obeticholic Acid in Primary Biliary Cholangitis. *N. Engl. J. Med.* **2016**, *375*, 631–643. <https://doi.org/10.1056/NEJMoa1509840>.
124. Trauner, M.; Nevens, F.; Shiffman, M.L.; Drenth, J.P.H.; Bowlus, C.L.; Vargas, V.; Andreone, P.; Hirschfield, G.M.; Pencek, R.; Malecha, E.S.; et al. Long-term efficacy and safety of obeticholic acid for patients with primary biliary cholangitis: 3-year results of an international open-label extension study. *Lancet Gastroenterol. Hepatol.* **2019**, *4*, 445–453. [https://doi.org/10.1016/S2468-1253\(19\)30094-9](https://doi.org/10.1016/S2468-1253(19)30094-9).
125. Hegade, V.S.; Kendrick, S.F.; Dobbins, R.L.; Miller, S.R.; Thompson, D.; Richards, D.; Storey, J.; Dukes, G.E.; Corrigan, M.; Oude Elferink, R.P.; et al. Effect of ileal bile acid transporter inhibitor GSK2330672 on pruritus in primary biliary cholangitis: A double-blind, randomised, placebo-controlled, crossover, phase 2a study. *Lancet* **2017**, *389*, 1114–1123. [https://doi.org/10.1016/S0140-6736\(17\)30319-7](https://doi.org/10.1016/S0140-6736(17)30319-7).
126. Liss, K.H.; Finck, B.N. PPARs and nonalcoholic fatty liver disease. *Biochimie* **2017**, *136*, 65–74. <https://doi.org/10.1016/j.biochi.2016.11.009>.
127. Li, T.; Chiang, J.Y. Regulation of bile acid and cholesterol metabolism by PPARs. *PPAR Res.* **2009**, *2009*, 501739. <https://doi.org/10.1155/2009/501739>.
128. Kersten, S.; Stienstra, R. The role and regulation of the peroxisome proliferator activated receptor alpha in human liver. *Biochimie* **2017**, *136*, 75–84. <https://doi.org/10.1016/j.biochi.2016.12.019>.
129. Xia, X.; Jung, D.; Webb, P.; Zhang, A.; Zhang, B.; Li, L.; Ayers, S.D.; Gabbi, C.; Ueno, Y.; Gustafsson, J.A.; et al. Liver X receptor beta and peroxisome proliferator-activated receptor delta regulate cholesterol transport in murine cholangiocytes. *Hepatology* **2012**, *56*, 2288–2296. <https://doi.org/10.1002/hep.25919>.

130. Mukundan, L.; Odegaard, J.I.; Morel, C.R.; Heredia, J.E.; Mwangi, J.W.; Ricardo-Gonzalez, R.R.; Goh, Y.P.; Eagle, A.R.; Dunn, S.E.; Awakuni, J.U.; et al. PPAR- δ senses and orchestrates clearance of apoptotic cells to promote tolerance. *Nat. Med.* **2009**, *15*, 1266–1272. <https://doi.org/10.1038/nm.2048>.
131. Harada, K.; Isse, K.; Kamihira, T.; Shimoda, S.; Nakanuma, Y. Th1 cytokine-induced downregulation of PPAR γ in human biliary cells relates to cholangitis in primary biliary cirrhosis. *Hepatology* **2005**, *41*, 1329–1338. <https://doi.org/10.1002/hep.20705>.
132. Nozaki, Y.; Harada, K.; Sanzen, T.; Nakanuma, Y. PPAR γ ligand attenuates portal inflammation in the MRL-lpr mouse: A new strategy to restrain cholangiopathy in primary biliary cirrhosis. *Med. Mol. Morphol.* **2013**, *46*, 153–159. <https://doi.org/10.1007/s00795-013-0017-0>.
133. Han, X.F.; Wang, Q.X.; Liu, Y.; You, Z.R.; Bian, Z.L.; Qiu, D.K.; Ma, X. Efficacy of fenofibrate in Chinese patients with primary biliary cirrhosis partially responding to ursodeoxycholic acid therapy. *J. Dig. Dis.* **2012**, *13*, 219–224. <https://doi.org/10.1111/j.1751-2980.2012.00574.x>.
134. Ohira, H.; Sato, Y.; Ueno, T.; Sata, M. Fenofibrate treatment in patients with primary biliary cirrhosis. *Am. J. Gastroenterol.* **2002**, *97*, 2147–2149. <https://doi.org/10.1111/j.1572-0241.2002.05944.x>.
135. Duan, W.; Ou, X.; Wang, X.; Wang, Y.; Zhao, X.; Wang, Q.; Wu, X.; Zhang, W.; Ma, H.; You, H.; et al. Efficacy and safety of fenofibrate add-on therapy for patients with primary biliary cholangitis and a suboptimal response to UDCA. *Rev. Esp. Enferm. Dig.* **2018**, *110*, 557–563. <https://doi.org/10.17235/reed.2018.5533/2018>.
136. Yin, Q.; Li, J.; Xia, Y.; Zhang, R.; Wang, J.; Lu, W.; Zhou, Y.; Zheng, Y.; Abudumijiti, H.; Chen, R.; et al. Systematic review and meta-analysis: Bezafibrate in patients with primary biliary cirrhosis. *Drug Des. Dev. Ther.* **2015**, *9*, 5407–5419. <https://doi.org/10.2147/DDDT.S92041>.
137. Reig, A.; Sese, P.; Pares, A. Effects of Bezafibrate on Outcome and Pruritus in Primary Biliary Cholangitis with Suboptimal Ursodeoxycholic Acid Response. *Am. J. Gastroenterol.* **2018**, *113*, 49–55. <https://doi.org/10.1038/ajg.2017.287>.
138. Honda, A.; Tanaka, A.; Kaneko, T.; Komori, A.; Abe, M.; Inao, M.; Namisaki, T.; Hashimoto, N.; Kawata, K.; Takahashi, A.; et al. Bezafibrate Improves GLOBE and UK-PBC Scores and Long-Term Outcomes in Patients with Primary Biliary Cholangitis. *Hepatology* **2019**, *70*, 2035–2046. <https://doi.org/10.1002/hep.30552>.
139. Bolier, R.; de Vries, E.S.; Pares, A.; Helder, J.; Kemper, E.M.; Zwinderman, K.; Elferink, R.P.O.; Beuers, U.; Netherlands Association for the Study of the Liver Cholestatic Liver Diseases Study Group. Fibrates for the treatment of cholestatic itch (FITCH): Study protocol for a randomized controlled trial. *Trials* **2017**, *18*, 230. <https://doi.org/10.1186/s13063-017-1966-8>.
140. Schattenberg, J.M.; Pares, A.; Kowdley, K.V.; Heneghan, M.A.; Caldwell, S.; Pratt, D.; Bonder, A.; Hirschfield, G.M.; Levy, C.; Vierling, J.; et al. A randomized placebo-controlled trial of elafibranor in patients with primary biliary cholangitis and incomplete response to UDCA. *J. Hepatol.* **2021**, *74*, 1344–1354. <https://doi.org/10.1016/j.jhep.2021.01.013>.
141. Vuppalanchi, R.; Caldwell, S.H.; Pyrsopoulos, N. Results of a Phase 2, Prospective, Multicenter, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate Safety, Tolerability, and Efficacy of Saroglitazar Magnesium in Patients with Primary Biliary Cholangitis (EPICS). *Gastroenterol. Hepatol.* **2021**, *17*, 8.
142. Jones, D.; Boudes, P.F.; Swain, M.G.; Bowlus, C.L.; Galambos, M.R.; Bacon, B.R.; Doerffel, Y.; Gitlin, N.; Gordon, S.C.; Odin, J.A.; et al. 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. *Lancet Gastroenterol. Hepatol.* **2017**, *2*, 716–726. [https://doi.org/10.1016/S2468-1253\(17\)30246-7](https://doi.org/10.1016/S2468-1253(17)30246-7).
143. Hirschfield, G.M.; Kowdley, K.V.; Shiffman, M.L. ENHANCE: Safety and Efficacy of Seladelpar in Patients with Primary Biliary Cholangitis—A Phase 3, International, Randomized, Placebo-Controlled Study. *Gastroenterol. Hepatol.* **2021**, *17*, 5–6.
144. Kremer, A.E.; Mayo, M.J.; Hirschfield, G.; Levy, C.; Bowlus, C.L.; Jones, D.E.; Steinberg, A.; McWherter, C.A.; Choi, Y.J. Seladelpar improved measures of pruritus, sleep, and fatigue and decreased serum bile acids in patients with primary biliary cholangitis. *Liver Int.* **2022**, *42*, 112–123. <https://doi.org/10.1111/liv.15039>.
145. Rautiainen, H.; Karkkainen, P.; Karvonen, A.L.; Nurmi, H.; Pikkarainen, P.; Nuutinen, H.; Farkkila, M. Budesonide combined with UDCA to improve liver histology in primary biliary cirrhosis: A three-year randomized trial. *Hepatology* **2005**, *41*, 747–752. <https://doi.org/10.1002/hep.20646>.
146. Hirschfield, G.M.; Beuers, U.; Kupcinskas, L.; Ott, P.; Bergquist, A.; Farkkila, M.; Manns, M.P.; Pares, A.; Spengler, U.; Stiess, M.; et al. A placebo-controlled randomised trial of budesonide for PBC following an insufficient response to UDCA. *J. Hepatol.* **2021**, *74*, 321–329. <https://doi.org/10.1016/j.jhep.2020.09.011>.
147. Kliewer, S.A.; Mangelsdorf, D.J. Bile Acids as Hormones: The FXR-FGF15/19 Pathway. *Dig. Dis.* **2015**, *33*, 327–331. <https://doi.org/10.1159/000371670>.
148. Russell, D.W. Fifty years of advances in bile acid synthesis and metabolism. *J. Lipid Res.* **2009**, *50*, S120–S125. <https://doi.org/10.1194/jlr.R800026-JLR200>.
149. Modica, S.; Petruzzelli, M.; Bellafante, E.; Murzilli, S.; Salvatore, L.; Celli, N.; Di Tullio, G.; Palasciano, G.; Moustafa, T.; Halilbasic, E.; et al. Selective activation of nuclear bile acid receptor FXR in the intestine protects mice against cholestasis. *Gastroenterology* **2012**, *142*, 355–365.e4. <https://doi.org/10.1053/j.gastro.2011.10.028>.
150. Luo, J.; Ko, B.; Elliott, M.; Zhou, M.; Lindhout, D.A.; Phung, V.; To, C.; Learned, R.M.; Tian, H.; DePaoli, A.M.; et al. A nontumorigenic variant of FGF19 treats cholestatic liver diseases. *Sci. Transl. Med.* **2014**, *6*, 247ra100. <https://doi.org/10.1126/scitranslmed.3009098>.

151. Nicholes, K.; Guillet, S.; Tomlinson, E.; Hillan, K.; Wright, B.; Frantz, G.D.; Pham, T.A.; Dillard-Telm, L.; Tsai, S.P.; Stephan, J.P.; et al. A mouse model of hepatocellular carcinoma: Ectopic expression of fibroblast growth factor 19 in skeletal muscle of transgenic mice. *Am. J. Pathol.* **2002**, *160*, 2295–2307. [https://doi.org/10.1016/S0002-9440\(10\)61177-7](https://doi.org/10.1016/S0002-9440(10)61177-7).
152. Mayo, M.J.; Wigg, A.J.; Leggett, B.A.; Arnold, H.; Thompson, A.J.; Weltman, M.; Carey, E.J.; Muir, A.J.; Ling, L.; Rossi, S.J.; et al. NGM282 for Treatment of Patients with Primary Biliary Cholangitis: A Multicenter, Randomized, Double-Blind, Placebo-Controlled Trial. *Hepatol. Commun.* **2018**, *2*, 1037–1050. <https://doi.org/10.1002/hep4.1209>.
153. Pellicciari, R.; Costantino, G.; Camaioni, E.; Sadeghpour, B.M.; Entrena, A.; Willson, T.M.; Fiorucci, S.; Clerici, C.; Gioiello, A. Bile acid derivatives as ligands of the farnesoid X receptor. Synthesis, evaluation, and structure-activity relationship of a series of body and side chain modified analogues of chenodeoxycholic acid. *J. Med. Chem.* **2004**, *47*, 4559–4569. <https://doi.org/10.1021/jm049904b>.
154. Gege, C.; Hambruch, E.; Hambruch, N.; Kinzel, O.; Kremoser, C. Nonsteroidal FXR Ligands: Current Status and Clinical Applications. *Handb. Exp. Pharmacol.* **2019**, *256*, 167–205. https://doi.org/10.1007/164_2019_232.
155. An, P.; Wei, G.; Huang, P.; Li, W.; Qi, X.; Lin, Y.; Vaid, K.A.; Wang, J.; Zhang, S.; Li, Y.; et al. A novel non-bile acid FXR agonist EDP-305 potently suppresses liver injury and fibrosis without worsening of ductular reaction. *Liver Int.* **2020**, *40*, 1655–1669. <https://doi.org/10.1111/liv.14490>.
156. Mogul, A.; Corsi, K.; McAuliffe, L. Baricitinib: The Second FDA-Approved JAK Inhibitor for the Treatment of Rheumatoid Arthritis. *Ann. Pharmacother.* **2019**, *53*, 947–953. <https://doi.org/10.1177/1060028019839650>.
157. Asuri, S.; McIntosh, S.; Taylor, V.; Rokeby, A.; Kelly, J.; Shumansky, K.; Field, L.L.; Yoshida, E.M.; Arbour, L. Primary Biliary Cholangitis in British Columbia First Nations: Clinical Features and Discovery of Novel Genetic Susceptibility Loci. *Liver Int.* **2018**, *38*, 940–948. <https://doi.org/10.1111/liv.13686>.
158. Gordon, S.C.; Trudeau, S.; Regev, A.; Uhas, J.M.; Chakladar, S.; Pinto-Correia, A.; Gottlieb, K.; Schlichting, D. Baricitinib and primary biliary cholangitis. *J. Transl. Autoimmun.* **2021**, *4*, 100107. <https://doi.org/10.1016/j.jtauto.2021.100107>.
159. Andringa, K.K.; King, A.L.; Eccleston, H.B.; Mantena, S.K.; Landar, A.; Jhala, N.C.; Dickinson, D.A.; Squadrito, G.L.; Bailey, S.M. Analysis of the liver mitochondrial proteome in response to ethanol and S-adenosylmethionine treatments: Novel molecular targets of disease and hepatoprotection. *Am. J. Physiol. Gastrointest. Liver Physiol.* **2010**, *298*, G732–G745. <https://doi.org/10.1152/ajpgi.00332.2009>.
160. Caballero, F.; Fernandez, A.; Matias, N.; Martinez, L.; Fucho, R.; Elena, M.; Caballeria, J.; Morales, A.; Fernandez-Checa, J.C.; Garcia-Ruiz, C. Specific contribution of methionine and choline in nutritional nonalcoholic steatohepatitis: Impact on mitochondrial S-adenosyl-L-methionine and glutathione. *J. Biol. Chem.* **2010**, *285*, 18528–18536. <https://doi.org/10.1074/jbc.M109.099333>.
161. Ko, K.; Yang, H.; Noureddin, M.; Iglesias-Ara, A.; Xia, M.; Wagner, C.; Luka, Z.; Mato, J.M.; Lu, S.C. Changes in S-adenosylmethionine and GSH homeostasis during endotoxemia in mice. *Lab. Investig.* **2008**, *88*, 1121–1129. <https://doi.org/10.1038/labinvest.2008.69>.
162. Mato, J.M.; Alvarez, L.; Ortiz, P.; Mingorance, J.; Duran, C.; Pajares, M.A. S-adenosyl-L-methionine synthetase and methionine metabolism deficiencies in cirrhosis. *Adv. Exp. Med. Biol.* **1994**, *368*, 113–117. https://doi.org/10.1007/978-1-4615-1989-8_11.
163. Wunsch, E.; Raszeja-Wyszomirska, J.; Barbier, O.; Milkiewicz, M.; Krawczyk, M.; Milkiewicz, P. Effect of S-adenosyl-L-methionine on liver biochemistry and quality of life in patients with primary biliary cholangitis treated with ursodeoxycholic acid. A prospective, open label pilot study. *J. Gastrointest. Liver Dis.* **2018**, *27*, 273–279. <https://doi.org/10.15403/jgld.2014.1121.273.icz>.
164. Kilanczyk, E.; Banales, J.M.; Wunsch, E.; Barbier, O.; Avila, M.A.; Mato, J.M.; Milkiewicz, M.; Milkiewicz, P. S-adenosyl-L-methionine (SAME) halts the autoimmune response in patients with primary biliary cholangitis (PBC) via anti-oxidant and S-glutathionylation processes in cholangiocytes. *Biochim. Biophys. Acta Mol. Basis Dis.* **2020**, *1866*, 165895. <https://doi.org/10.1016/j.bbdis.2020.165895>.
165. Liu, Y.; Chen, K.; Li, F.; Gu, Z.; Liu, Q.; He, L.; Shao, T.; Song, Q.; Zhu, F.; Zhang, L.; et al. Probiotic *Lactobacillus rhamnosus* GG Prevents Liver Fibrosis Through Inhibiting Hepatic Bile Acid Synthesis and Enhancing Bile Acid Excretion in Mice. *Hepatology* **2020**, *71*, 2050–2066. <https://doi.org/10.1002/hep.30975>.
166. Volarevic, V.; Ljubic, B.; Stojkovic, P.; Lukic, A.; Arsenijevic, N.; Stojkovic, M. Human stem cell research and regenerative medicine—present and future. *Br. Med. Bull.* **2011**, *99*, 155–168. <https://doi.org/10.1093/bmb/ldr027>.
167. Wang, L.; Li, J.; Liu, H.; Li, Y.; Fu, J.; Sun, Y.; Xu, R.; Lin, H.; Wang, S.; Lv, S.; et al. Pilot study of umbilical cord-derived mesenchymal stem cell transfusion in patients with primary biliary cirrhosis. *J. Gastroenterol. Hepatol.* **2013**, *28* (Suppl. S1), 85–92. <https://doi.org/10.1111/jgh.12029>.
168. Wang, L.; Han, Q.; Chen, H.; Wang, K.; Shan, G.L.; Kong, F.; Yang, Y.J.; Li, Y.Z.; Zhang, X.; Dong, F.; et al. Allogeneic bone marrow mesenchymal stem cell transplantation in patients with UDCA-resistant primary biliary cirrhosis. *Stem Cells Dev.* **2014**, *23*, 2482–2489. <https://doi.org/10.1089/scd.2013.0500>.
169. Isse, K.; Harada, K.; Zen, Y.; Kamihira, T.; Shimoda, S.; Harada, M.; Nakanuma, Y. Fractalkine and CX3CR1 are involved in the recruitment of intraepithelial lymphocytes of intrahepatic bile ducts. *Hepatology* **2005**, *41*, 506–516. <https://doi.org/10.1002/hep.20582>.

170. Tabuchi, H.; Katsurabara, T.; Mori, M.; Aoyama, M.; Obara, T.; Yasuda, N.; Kawano, T.; Imai, T.; Ieiri, I.; Kumagai, Y. Pharmacokinetics, Pharmacodynamics, and Safety of E6011, a Novel Humanized Antifractalkine (CX3CL1) Monoclonal Antibody: A Randomized, Double-Blind, Placebo-Controlled Single-Ascending-Dose Study. *J. Clin. Pharmacol.* **2019**, *59*, 688–701. <https://doi.org/10.1002/jcph.1361>.
171. Chapman, M.H.; Thorburn, D.; Hirschfield, G.M.; Webster, G.G.J.; Rushbrook, S.M.; Alexander, G.; Collier, J.; Dyson, J.K.; Jones, D.E.; Patanwala, I.; et al. British Society of Gastroenterology and UK-PSC guidelines for the diagnosis and management of primary sclerosing cholangitis. *Gut* **2019**, *68*, 1356–1378. <https://doi.org/10.1136/gutjnl-2018-317993>.
172. Lazaridis, K.N.; LaRusso, N.F. Primary Sclerosing Cholangitis. *N. Engl. J. Med.* **2016**, *375*, 2501–2502. <https://doi.org/10.1056/NEJMc1613273>.
173. Karlsen, T.H.; Folseraas, T.; Thorburn, D.; Vesterhus, M. Primary sclerosing cholangitis—A comprehensive review. *J. Hepatol.* **2017**, *67*, 1298–1323. <https://doi.org/10.1016/j.jhep.2017.07.022>.
174. Sirpal, S.; Chandok, N. Primary sclerosing cholangitis: Diagnostic and management challenges. *Clin. Exp. Gastroenterol.* **2017**, *10*, 265–273. <https://doi.org/10.2147/CEG.S105872>.
175. Dyson, J.K.; Beuers, U.; Jones, D.E.J.; Lohse, A.W.; Hudson, M. Primary sclerosing cholangitis. *Lancet* **2018**, *391*, 2547–2559. [https://doi.org/10.1016/S0140-6736\(18\)30300-3](https://doi.org/10.1016/S0140-6736(18)30300-3).
176. Jiang, X.; Karlsen, T.H. Genetics of primary sclerosing cholangitis and pathophysiological implications. *Nat. Rev. Gastroenterol. Hepatol.* **2017**, *14*, 279–295. <https://doi.org/10.1038/nrgastro.2016.154>.
177. Bergquist, A.; Montgomery, S.M.; Bahmanyar, S.; Olsson, R.; Danielsson, A.; Lindgren, S.; Prytz, H.; Hultcrantz, R.; Loof, L.A.; Sandberg-Gertzén, H.; et al. Increased risk of primary sclerosing cholangitis and ulcerative colitis in first-degree relatives of patients with primary sclerosing cholangitis. *Clin. Gastroenterol. Hepatol.* **2008**, *6*, 939–943. <https://doi.org/10.1016/j.cgh.2008.03.016>.
178. Ji, S.G.; Juran, B.D.; Mucha, S.; Folseraas, T.; Jostins, L.; Melum, E.; Kumasaka, N.; Atkinson, E.J.; Schlicht, E.M.; Liu, J.Z.; et al. Genome-wide association study of primary sclerosing cholangitis identifies new risk loci and quantifies the genetic relationship with inflammatory bowel disease. *Nat. Genet.* **2017**, *49*, 269–273. <https://doi.org/10.1038/ng.3745>.
179. Ataseven, H.; Parlak, E.; Yuksel, I.; Basar, O.; Ertugrul, I.; Sasmaz, N.; Sahin, B. Primary sclerosing cholangitis in Turkish patients: Characteristic features and prognosis. *Hepatobiliary Pancreat. Dis. Int.* **2009**, *8*, 312–315.
180. Tanaka, A.; Tazuma, S.; Okazaki, K.; Tsubouchi, H.; Inui, K.; Takikawa, H. Nationwide survey for primary sclerosing cholangitis and IgG4-related sclerosing cholangitis in Japan. *J. Hepatobiliary Pancreat. Sci.* **2014**, *21*, 43–50. <https://doi.org/10.1002/jhbp.50>.
181. Saarinen, S.; Olerup, O.; Broome, U. Increased frequency of autoimmune diseases in patients with primary sclerosing cholangitis. *Am. J. Gastroenterol.* **2000**, *95*, 3195–3199. <https://doi.org/10.1111/j.1572-0241.2000.03292.x>.
182. Liu, J.Z.; Hov, J.R.; Folseraas, T.; Ellinghaus, E.; Rushbrook, S.M.; Doncheva, N.T.; Andreassen, O.A.; Weersma, R.K.; Weismuller, T.J.; Eksteen, B.; et al. Dense genotyping of immune-related disease regions identifies nine new risk loci for primary sclerosing cholangitis. *Nat. Genet.* **2013**, *45*, 670–675. <https://doi.org/10.1038/ng.2616>.
183. Hov, J.R.; Kosmoliaptis, V.; Traherne, J.A.; Olsson, M.; Boberg, K.M.; Bergquist, A.; Schrumpf, E.; Bradley, J.A.; Taylor, C.J.; Lie, B.A.; et al. Electrostatic modifications of the human leukocyte antigen-DR P9 peptide-binding pocket and susceptibility to primary sclerosing cholangitis. *Hepatology* **2011**, *53*, 1967–1976. <https://doi.org/10.1002/hep.24299>.
184. Hov, J.R.; Lleo, A.; Selmi, C.; Woldseth, B.; Fabris, L.; Strazzabosco, M.; Karlsen, T.H.; Invernizzi, P. Genetic associations in Italian primary sclerosing cholangitis: Heterogeneity across Europe defines a critical role for HLA-C. *J. Hepatol.* **2010**, *52*, 712–717. <https://doi.org/10.1016/j.jhep.2009.11.029>.
185. Eksteen, B. Advances and controversies in the pathogenesis and management of primary sclerosing cholangitis. *Br. Med. Bull.* **2014**, *110*, 89–98. <https://doi.org/10.1093/bmb/ldu008>.
186. Hofmann, A.F. Bile acids: Trying to understand their chemistry and biology with the hope of helping patients. *Hepatology* **2009**, *49*, 1403–1418. <https://doi.org/10.1002/hep.22789>.
187. Hohenester, S.; Gates, A.; Wimmer, R.; Beuers, U.; Anwer, M.S.; Rust, C.; Webster, C.R. Phosphatidylinositol-3-kinase p110gamma contributes to bile salt-induced apoptosis in primary rat hepatocytes and human hepatoma cells. *J. Hepatol.* **2010**, *53*, 918–926. <https://doi.org/10.1016/j.jhep.2010.05.015>.
188. Hang, S.; Paik, D.; Yao, L.; Kim, E.; Trinath, J.; Lu, J.; Ha, S.; Nelson, B.N.; Kelly, S.P.; Wu, L.; et al. Bile acid metabolites control TH17 and Treg cell differentiation. *Nature* **2019**, *576*, 143–148. <https://doi.org/10.1038/s41586-019-1785-z>.
189. Bell, L.N.; Wulff, J.; Comerford, M.; Vuppalanchi, R.; Chalasani, N. Serum metabolic signatures of primary biliary cirrhosis and primary sclerosing cholangitis. *Liver Int.* **2015**, *35*, 263–274. <https://doi.org/10.1111/liv.12680>.
190. Deutschmann, K.; Reich, M.; Klindt, C.; Droge, C.; Spomer, L.; Haussinger, D.; Keitel, V. Bile acid receptors in the biliary tree: TGR5 in physiology and disease. *Biochim. Biophys. Acta Mol. Basis Dis.* **2018**, *1864*, 1319–1325. <https://doi.org/10.1016/j.bbadis.2017.08.021>.
191. Zhang, J.H.; Nolan, J.D.; Kennie, S.L.; Johnston, I.M.; Dew, T.; Dixon, P.H.; Williamson, C.; Walters, J.R. Potent stimulation of fibroblast growth factor 19 expression in the human ileum by bile acids. *Am. J. Physiol. Gastrointest. Liver Physiol.* **2013**, *304*, G940–G948. <https://doi.org/10.1152/ajpgi.00398.2012>.
192. Milkiewicz, M.; Klak, M.; Kempinska-Podhorodecka, A.; Wiechowska-Kozłowska, A.; Urasinska, E.; Blatkiwicz, M.; Wunsch, E.; Elias, E.; Milkiewicz, P. Impaired Hepatic Adaptation to Chronic Cholestasis induced by Primary Sclerosing Cholangitis. *Sci. Rep.* **2016**, *6*, 39573. <https://doi.org/10.1038/srep39573>.

193. Alvaro, D.; Gigliozzi, A.; Attili, A.F. Regulation and deregulation of cholangiocyte proliferation. *J. Hepatol.* **2000**, *33*, 333–340. [https://doi.org/10.1016/s0168-8278\(00\)80377-3](https://doi.org/10.1016/s0168-8278(00)80377-3).
194. Carpino, G.; Cardinale, V.; Renzi, A.; Hov, J.R.; Berloco, P.B.; Rossi, M.; Karlsen, T.H.; Alvaro, D.; Gaudio, E. Activation of biliary tree stem cells within peribiliary glands in primary sclerosing cholangitis. *J. Hepatol.* **2015**, *63*, 1220–1228. <https://doi.org/10.1016/j.jhep.2015.06.018>.
195. Karlsen, T.H. Primary sclerosing cholangitis: 50 years of a gut-liver relationship and still no love? *Gut* **2016**, *65*, 1579–1581. <https://doi.org/10.1136/gutjnl-2016-312137>.
196. Almeida, A.R.; Legrand, N.; Papiernik, M.; Freitas, A.A. Homeostasis of peripheral CD4⁺ T cells: IL-2R alpha and IL-2 shape a population of regulatory cells that controls CD4⁺ T cell numbers. *J. Immunol.* **2002**, *169*, 4850–4860. <https://doi.org/10.4049/jimmunol.169.9.4850>.
197. Mederacke, I.; Hsu, C.C.; Troeger, J.S.; Huebener, P.; Mu, X.; Dapito, D.H.; Pradere, J.P.; Schwabe, R.F. Fate tracing reveals hepatic stellate cells as dominant contributors to liver fibrosis independent of its aetiology. *Nat. Commun.* **2013**, *4*, 2823. <https://doi.org/10.1038/ncomms3823>.
198. Lemoine, S.; Cadoret, A.; Rautou, P.E.; El Mourabit, H.; Ratzu, V.; Corpechot, C.; Rey, C.; Bosselut, N.; Barbu, V.; Wendum, D.; et al. Portal myofibroblasts promote vascular remodeling underlying cirrhosis formation through the release of microparticles. *Hepatology* **2015**, *61*, 1041–1055. <https://doi.org/10.1002/hep.27318>.
199. Ferreira-Gonzalez, S.; Lu, W.Y.; Raven, A.; Dwyer, B.; Man, T.Y.; O'Duibhir, E.; Lewis, P.J.S.; Campana, L.; Kendall, T.J.; Bird, T.G.; et al. Paracrine cellular senescence exacerbates biliary injury and impairs regeneration. *Nat. Commun.* **2018**, *9*, 1020. <https://doi.org/10.1038/s41467-018-03299-5>.
200. Kyritsi, K.; Francis, H.; Zhou, T.; Ceci, L.; Wu, N.; Yang, Z.; Meng, F.; Chen, L.; Baiocchi, L.; Kundu, D.; et al. Downregulation of p16 Decreases Biliary Damage and Liver Fibrosis in the Mdr2(−) Mouse Model of Primary Sclerosing Cholangitis. *Gene Expr.* **2020**, *20*, 89–103. <https://doi.org/10.3727/105221620X15889714507961>.
201. Trussoni, C.E.; O'Hara, S.P.; LaRusso, N.F. Cellular senescence in the cholangiopathies: A driver of immunopathology and a novel therapeutic target. *Semin. Immunopathol.* **2022**. <https://doi.org/10.1007/s00281-022-00909-9>.
202. Cameron, R.G.; Blendis, L.M.; Neuman, M.G. Accumulation of macrophages in primary sclerosing cholangitis. *Clin. Biochem.* **2001**, *34*, 195–201. [https://doi.org/10.1016/s0009-9120\(01\)00215-6](https://doi.org/10.1016/s0009-9120(01)00215-6).
203. Grant, A.J.; Lalor, P.F.; Hubscher, S.G.; Briskin, M.; Adams, D.H. MAdCAM-1 expressed in chronic inflammatory liver disease supports mucosal lymphocyte adhesion to hepatic endothelium (MAdCAM-1 in chronic inflammatory liver disease). *Hepatology* **2001**, *33*, 1065–1072. <https://doi.org/10.1053/jhep.2001.24231>.
204. Eksteen, B.; Grant, A.J.; Miles, A.; Curbishley, S.M.; Lalor, P.F.; Hubscher, S.G.; Briskin, M.; Salmon, M.; Adams, D.H. Hepatic endothelial CCL25 mediates the recruitment of CCR9⁺ gut-homing lymphocytes to the liver in primary sclerosing cholangitis. *J. Exp. Med.* **2004**, *200*, 1511–1517. <https://doi.org/10.1084/jem.20041035>.
205. Trivedi, P.J.; Adams, D.H. Gut-liver immunity. *J. Hepatol.* **2016**, *64*, 1187–1189. <https://doi.org/10.1016/j.jhep.2015.12.002>.
206. Trivedi, P.J.; Adams, D.H. Mucosal immunity in liver autoimmunity: A comprehensive review. *J. Autoimmun.* **2013**, *46*, 97–111. <https://doi.org/10.1016/j.jaut.2013.06.013>.
207. Worthington, J.; Cullen, S.; Chapman, R. Immunopathogenesis of primary sclerosing cholangitis. *Clin. Rev. Allergy Immunol.* **2005**, *28*, 93–103. <https://doi.org/10.1385/CRIAI:28:2:093>.
208. Ruhlemann, M.; Liwinski, T.; Heinsen, F.A.; Bang, C.; Zenouzi, R.; Kummen, M.; Thingholm, L.; Tempel, M.; Lieb, W.; Karlsen, T.; et al. Consistent alterations in faecal microbiomes of patients with primary sclerosing cholangitis independent of associated colitis. *Aliment. Pharmacol. Ther.* **2019**, *50*, 580–589. <https://doi.org/10.1111/apt.15375>.
209. Kummen, M.; Holm, K.; Anmarkrud, J.A.; Nygard, S.; Vesterhus, M.; Hoivik, M.L.; Troseid, M.; Marschall, H.U.; Schruppf, E.; Moum, B.; et al. The gut microbial profile in patients with primary sclerosing cholangitis is distinct from patients with ulcerative colitis without biliary disease and healthy controls. *Gut* **2017**, *66*, 611–619. <https://doi.org/10.1136/gutjnl-2015-310500>.
210. Bajer, L.; Kverka, M.; Kostovcik, M.; Macinga, P.; Dvorak, J.; Stehlikova, Z.; Brezina, J.; Wohl, P.; Spicak, J.; Drastich, P. Distinct gut microbiota profiles in patients with primary sclerosing cholangitis and ulcerative colitis. *World J. Gastroenterol.* **2017**, *23*, 4548–4558. <https://doi.org/10.3748/wjg.v23.i25.4548>.
211. Sabino, J.; Vieira-Silva, S.; Machiels, K.; Joossens, M.; Falony, G.; Ballet, V.; Ferrante, M.; Van Assche, G.; Van der Merwe, S.; Vermeire, S.; et al. Primary sclerosing cholangitis is characterised by intestinal dysbiosis independent from IBD. *Gut* **2016**, *65*, 1681–1689. <https://doi.org/10.1136/gutjnl-2015-311004>.
212. Weston, C.J.; Shepherd, E.L.; Claridge, L.C.; Rantakari, P.; Curbishley, S.M.; Tomlinson, J.W.; Hubscher, S.G.; Reynolds, G.M.; Aalto, K.; Anstee, Q.M.; et al. Vascular adhesion protein-1 promotes liver inflammation and drives hepatic fibrosis. *J. Clin. Invest.* **2015**, *125*, 501–520. <https://doi.org/10.1172/JCI73722>.
213. Shah, A.; Crawford, D.; Burger, D.; Martin, N.; Walker, M.; Talley, N.J.; Tallis, C.; Jones, M.; Stuart, K.; Keely, S.; et al. Effects of Antibiotic Therapy in Primary Sclerosing Cholangitis with and without Inflammatory Bowel Disease: A Systematic Review and Meta-Analysis. *Semin. Liver Dis.* **2019**, *39*, 432–441. <https://doi.org/10.1055/s-0039-1688501>.
214. Yatsunenko, T.; Rey, F.E.; Manary, M.J.; Trehan, I.; Dominguez-Bello, M.G.; Contreras, M.; Magris, M.; Hidalgo, G.; Baldassano, R.N.; Anokhin, A.P.; et al. Human gut microbiome viewed across age and geography. *Nature* **2012**, *486*, 222–227. <https://doi.org/10.1038/nature11053>.

215. Vieira-Silva, S.; Sabino, J.; Valles-Colomer, M.; Falony, G.; Kathagen, G.; Caenepeel, C.; Cleynen, I.; van der Merwe, S.; Vermeire, S.; Raes, J. Quantitative microbiome profiling disentangles inflammation- and bile duct obstruction-associated microbiota alterations across PSC/IBD diagnoses. *Nat. Microbiol.* **2019**, *4*, 1826–1831. <https://doi.org/10.1038/s41564-019-0483-9>.
216. Quraishi, M.N.; Acharjee, A.; Beggs, A.D.; Horniblow, R.; Tselepis, C.; Gkoutos, G.; Ghosh, S.; Rossiter, A.E.; Loman, N.; van Schaik, W.; et al. 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. *J. Crohns Colitis* **2020**, *14*, 935–947. <https://doi.org/10.1093/ecco-jcc/jjaa021>.
217. Jiang, B.; Yuan, G.; Wu, J.; Wu, Q.; Li, L.; Jiang, P. *Prevotella copri* ameliorates cholestasis and liver fibrosis in primary sclerosing cholangitis by enhancing the FXR signalling pathway. *Biochim. Biophys. Acta Mol. Basis Dis.* **2022**, *1868*, 166320. <https://doi.org/10.1016/j.bbadis.2021.166320>.
218. Little, R.; Wine, E.; Kamath, B.M.; Griffiths, A.M.; Ricciuto, A. Gut microbiome in primary sclerosing cholangitis: A review. *World J. Gastroenterol.* **2020**, *26*, 2768–2780. <https://doi.org/10.3748/wjg.v26.i21.2768>.
219. Lemoine, S.; Kemgang, A.; Ben Belkacem, K.; Straube, M.; Jegou, S.; Corpechot, C.; Saint-Antoine IBD Network; Chazouilleres, O.; Housset, C.; Sokol, H. Fungi participate in the dysbiosis of gut microbiota in patients with primary sclerosing cholangitis. *Gut* **2020**, *69*, 92–102. <https://doi.org/10.1136/gutjnl-2018-317791>.
220. Hong, K.H.; Kim, J.W.; Jang, S.J.; Yu, E.; Kim, E.C. Liver cirrhosis caused by *Exophiala dermatitidis*. *J. Med. Microbiol.* **2009**, *58*, 674–677. <https://doi.org/10.1099/jmm.0.002188-0>.
221. Kremer, A.E.; Namer, B.; Bolier, R.; Fischer, M.J.; Oude Elferink, R.P.; Beuers, U. Pathogenesis and Management of Pruritus in PBC and PSC. *Dig. Dis.* **2015**, *33* (Suppl. S2), 164–175. <https://doi.org/10.1159/000440829>.
222. de Vries, E.; Bolier, R.; Goet, J.; Pares, A.; Verbeek, J.; de Vree, M.; Drenth, J.; van Erpecum, K.; van Nieuwkerk, K.; van der Heide, F.; et al. Fibrates for Itch (FITCH) in Fibrosing Cholangiopathies: A Double-Blind, Randomized, Placebo-Controlled Trial. *Gastroenterology* **2021**, *160*, 734–743.e6. <https://doi.org/10.1053/j.gastro.2020.10.001>.
223. Kumada, H.; Miyakawa, H.; Muramatsu, T.; Ando, N.; Oh, T.; Takamori, K.; Nakamoto, H. Efficacy of nalfurafine hydrochloride in patients with chronic liver disease with refractory pruritus: A randomized, double-blind trial. *Hepatol. Res.* **2017**, *47*, 972–982. <https://doi.org/10.1111/hepr.12830>.
224. Dull, M.M.; Wolf, K.; Vetter, M.; Dietrich, P.; Neurath, M.F.; Kremer, A.E. Endogenous Opioid Levels Do Not Correlate with Itch Intensity and Therapeutic Interventions in Hepatic Pruritus. *Front. Med.* **2021**, *8*, 641163. <https://doi.org/10.3389/fmed.2021.641163>.
225. Lindor, K.D.; Kowdley, K.V.; Harrison, M.E.; American College of Gastroenterology. ACG Clinical Guideline: Primary Sclerosing Cholangitis. *Am. J. Gastroenterol.* **2015**, *110*, 646–659. <https://doi.org/10.1038/ajg.2015.112>.
226. Rahimpour, S.; Nasiri-Toosi, M.; Khalili, H.; Ebrahimi-Daryani, N.; Nouri-Taromlou, M.K.; Azizi, Z. A Triple Blinded, Randomized, Placebo-Controlled Clinical Trial to Evaluate the Efficacy and Safety of Oral Vancomycin in Primary Sclerosing Cholangitis: A Pilot Study. *J. Gastrointest. Liver Dis.* **2016**, *25*, 457–464. <https://doi.org/10.15403/jgld.2014.1121.254.rah>.
227. Hilscher, M.B.; Tabibian, J.H.; Carey, E.J.; Gostout, C.J.; Lindor, K.D. Dominant strictures in primary sclerosing cholangitis: A multicenter survey of clinical definitions and practices. *Hepatol. Commun.* **2018**, *2*, 836–844. <https://doi.org/10.1002/hep4.1194>.
228. Stiehl, A.; Rudolph, G.; Kloters-Plachky, P.; Sauer, P.; Walker, S. Development of dominant bile duct stenoses in patients with primary sclerosing cholangitis treated with ursodeoxycholic acid: Outcome after endoscopic treatment. *J. Hepatol.* **2002**, *36*, 151–156. [https://doi.org/10.1016/s0168-8278\(01\)00251-3](https://doi.org/10.1016/s0168-8278(01)00251-3).
229. Bjornsson, E.; Lindqvist-Ottosson, J.; Asztely, M.; Olsson, R. Dominant strictures in patients with primary sclerosing cholangitis. *Am. J. Gastroenterol.* **2004**, *99*, 502–508. <https://doi.org/10.1111/j.1572-0241.2004.04106.x>.
230. European Society of Gastrointestinal Endoscopy; European Association for the Study of the Liver. 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. *J. Hepatol.* **2017**, *66*, 1265–1281. <https://doi.org/10.1016/j.jhep.2017.02.013>.
231. Rupp, C.; Hippchen, T.; Bruckner, T.; Kloters-Plachky, P.; Schaible, A.; Koschny, R.; Stiehl, A.; Gotthardt, D.N.; Sauer, P. Effect of scheduled endoscopic dilatation of dominant strictures on outcome in patients with primary sclerosing cholangitis. *Gut* **2019**, *68*, 2170–2178. <https://doi.org/10.1136/gutjnl-2018-316801>.
232. Ferreira, M.; Ribeiro, I.B.; de Moura, D.T.H.; McCarty, T.R.; da Ponte Neto, A.M.; Farias, G.F.A.; de Miranda Neto, A.A.; de Oliveira, P.; Bernardo, W.M.; de Moura, E.G.H. Stent versus Balloon Dilation for the Treatment of Dominant Strictures in Primary Sclerosing Cholangitis: A Systematic Review and Meta-Analysis. *Clin. Endosc.* **2021**, *54*, 833–842. <https://doi.org/10.5946/ce.2021.052>.
233. Song, J.; Li, Y.; Bowlus, C.L.; Yang, G.; Leung, P.S.C.; Gershwin, M.E. Cholangiocarcinoma in Patients with Primary Sclerosing Cholangitis (PSC): A Comprehensive Review. *Clin. Rev. Allergy Immunol.* **2020**, *58*, 134–149. <https://doi.org/10.1007/s12016-019-08764-7>.
234. Said, K.; Glaumann, H.; Bergquist, A. Gallbladder disease in patients with primary sclerosing cholangitis. *J. Hepatol.* **2008**, *48*, 598–605. <https://doi.org/10.1016/j.jhep.2007.11.019>.
235. Mainprize, K.S.; Gould, S.W.; Gilbert, J.M. Surgical management of polypoid lesions of the gallbladder. *Br. J. Surg.* **2000**, *87*, 414–417. <https://doi.org/10.1046/j.1365-2168.2000.01363.x>.
236. Torabi Sagvand, B.; Edwards, K.; Shen, B. Frequency, Risk Factors, and Outcome of Gallbladder Polyps in Patients with Primary Sclerosing Cholangitis: A Case-Control Study. *Hepatol. Commun.* **2018**, *2*, 1440–1445. <https://doi.org/10.1002/hep4.1276>.

237. Hildebrand, T.; Pannicke, N.; Dechene, A.; Gotthardt, D.N.; Kirchner, G.; Reiter, F.P.; Sterneck, M.; Herzer, K.; Lenzen, H.; Rupp, C.; et al. Biliary strictures and recurrence after liver transplantation for primary sclerosing cholangitis: A retrospective multicenter analysis. *Liver Transpl.* **2016**, *22*, 42–52. <https://doi.org/10.1002/lt.24350>.
238. Trivedi, P.J.; Crothers, H.; Mytton, J.; Bosch, S.; Iqbal, T.; Ferguson, J.; Hirschfield, G.M. Effects of Primary Sclerosing Cholangitis on Risks of Cancer and Death in People with Inflammatory Bowel Disease, Based on Sex, Race, and Age. *Gastroenterology* **2020**, *159*, 915–928. <https://doi.org/10.1053/j.gastro.2020.05.049>.
239. Harms, M.H.; van Buuren, H.R.; Corpechot, C.; Thorburn, D.; Janssen, H.L.A.; Lindor, K.D.; Hirschfield, G.M.; Pares, A.; Floreani, A.; Mayo, M.J.; et al. Ursodeoxycholic acid therapy and liver transplant-free survival in patients with primary biliary cholangitis. *J. Hepatol.* **2019**, *71*, 357–365. <https://doi.org/10.1016/j.jhep.2019.04.001>.
240. Wunsch, E.; Trotter, J.; Milkiewicz, M.; Raszeja-Wyszomirska, J.; Hirschfield, G.M.; Barbier, O.; Milkiewicz, P. Prospective evaluation of ursodeoxycholic acid withdrawal in patients with primary sclerosing cholangitis. *Hepatology* **2014**, *60*, 931–940. <https://doi.org/10.1002/hep.27074>.
241. Stokkeland, K.; Hoijer, J.; Bottai, M.; Soderberg-Lofdal, K.; Bergquist, A. Statin Use Is Associated with Improved Outcomes of Patients with Primary Sclerosing Cholangitis. *Clin. Gastroenterol. Hepatol.* **2019**, *17*, 1860–1866.e1. <https://doi.org/10.1016/j.cgh.2018.11.002>.
242. Zhou, M.; Luo, J.; Chen, M.; Yang, H.; Learned, R.M.; DePaoli, A.M.; Tian, H.; Ling, L. Mouse species-specific control of hepatocarcinogenesis and metabolism by FGF19/FGF15. *J. Hepatol.* **2017**, *66*, 1182–1192. <https://doi.org/10.1016/j.jhep.2017.01.027>.
243. Mizuno, S.; Hirano, K.; Isayama, H.; Watanabe, T.; Yamamoto, N.; Nakai, Y.; Sasahira, N.; Tada, M.; Omata, M.; Koike, K. Prospective study of bezafibrate for the treatment of primary sclerosing cholangitis. *J. Hepatobiliary Pancreat. Sci.* **2015**, *22*, 766–770. <https://doi.org/10.1002/jhbp.281>.
244. Hirano, F.; Kobayashi, A.; Makino, I. Inhibition of TNF-alpha-induced RANTES expression in human hepatocyte-derived cells by fibrates, the hypolipidemic drugs. *Int. Immunopharmacol.* **2003**, *3*, 225–232. [https://doi.org/10.1016/S1567-5769\(02\)00275-8](https://doi.org/10.1016/S1567-5769(02)00275-8).
245. Schierwagen, R.; Uschner, F.E.; Magdaleno, F.; Klein, S.; Trebicka, J. Rationale for the use of statins in liver disease. *Am. J. Physiol. Gastrointest. Liver Physiol.* **2017**, *312*, G407–G412. <https://doi.org/10.1152/ajpgi.00441.2016>.
246. Chung, B.K.; Hirschfield, G.M. Immunogenetics in primary sclerosing cholangitis. *Curr. Opin. Gastroenterol.* **2017**, *33*, 93–98. <https://doi.org/10.1097/MOG.0000000000000336>.
247. Ali, A.H.; Damman, J.; Shah, S.B.; Davies, Y.; Hurwitz, M.; Stephen, M.; Lemos, L.M.; Carey, E.J.; Lindor, K.D.; Buness, C.W.; et al. Open-label prospective therapeutic clinical trials: Oral vancomycin in children and adults with primary sclerosing cholangitis. *Scand. J. Gastroenterol.* **2020**, *55*, 941–950. <https://doi.org/10.1080/00365521.2020.1787501>.
248. Fickert, P.; Hirschfield, G.M.; Denk, G.; Marschall, H.U.; Altorjay, I.; Farkkila, M.; Schramm, C.; Spengler, U.; Chapman, R.; Bergquist, A.; et al. norUrsodeoxycholic acid improves cholestasis in primary sclerosing cholangitis. *J. Hepatol.* **2017**, *67*, 549–558. <https://doi.org/10.1016/j.jhep.2017.05.009>.
249. Kowdley, K.V.; Vuppalanchi, R.; Levy, C.; Floreani, A.; Andreone, P.; LaRusso, N.F.; Shrestha, R.; Trotter, J.; Goldberg, D.; Rushbrook, S.; et al. A randomized, placebo-controlled, phase II study of obeticholic acid for primary sclerosing cholangitis. *J. Hepatol.* **2020**, *73*, 94–101. <https://doi.org/10.1016/j.jhep.2020.02.033>.
250. Carey, E.J.; Eaton, J.; Clayton, M.; Gossard, A.; Iqbal, S.; Ullah, H.; Zhang, N.; Butterfield, R.; Lindor, K.D. A pilot study of vidofludimus calcium for treatment of primary sclerosing cholangitis. *Hepatol. Commun.* **2022**. <https://doi.org/10.1002/hep4.1926>.
251. Eksteen, B.; Bowlus, C.L.; Montano-Loza, A.J.; Lefebvre, E.; Fischer, L.; Vig, P.; Martins, E.B.; Ahmad, J.; Yimam, K.K.; Pockros, P.J.; et al. Efficacy and Safety of Cenicriviroc in Patients with Primary Sclerosing Cholangitis: PERSEUS Study. *Hepatol. Commun.* **2021**, *5*, 478–490. <https://doi.org/10.1002/hep4.1619>.
252. Hirschfield, G.M.; Chazouilleres, O.; Drenth, J.P.; Thorburn, D.; Harrison, S.A.; Landis, C.S.; Mayo, M.J.; Muir, A.J.; Trotter, J.F.; Leeming, D.J.; et al. Effect of NGM282, an FGF19 analogue, in primary sclerosing cholangitis: A multicenter, randomized, double-blind, placebo-controlled phase II trial. *J. Hepatol.* **2019**, *70*, 483–493. <https://doi.org/10.1016/j.jhep.2018.10.035>.
253. Allegretti, J.R.; Kassam, Z.; Carrellas, M.; Mullish, B.H.; Marchesi, J.R.; Pechlivanis, A.; Smith, M.; Gerardin, Y.; Timberlake, S.; Pratt, D.S.; et al. Fecal Microbiota Transplantation in Patients with Primary Sclerosing Cholangitis: A Pilot Clinical Trial. *Am. J. Gastroenterol.* **2019**, *114*, 1071–1079. <https://doi.org/10.14309/ajg.0000000000000115>.
254. Gidwaney, N.G.; Pawa, S.; Das, K.M. Pathogenesis and clinical spectrum of primary sclerosing cholangitis. *World J Gastroenterol.* **2017**, *23*, 2459–2469. <https://doi.org/10.3748/wjg.v23.i14.2459>.