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A Review: Pathogenesis of Cholestatic Liver Diseases



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In this review we develop the argument that cholestatic liver diseases, particularly primary biliary cholangitis and primary sclerosing cholangitis (PSC), evolve over time with anatomically an ascending course of the disease process. The first and early lesions are in "downstream" bile ducts.

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Cited by: 229 Author: Gideon M. Hirschfield, Gideon M. Hirschfi...

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Patients with cholestatic liver diseases such as primary **biliary cirrhosis**, primary **sclerosing cholangitis** and **intrahepatic cholestasis** of pregnancy commonly complain of **pruritus**. The underlying **pathogenesis remains obscure with several mediators possibly playing an important role; these include**

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**Name of Journal:** *World Journal of Hepatology*  
**Manuscript NO:** 56548  
**Manuscript Type:** REVIEW

**Review:** Pathogenesis of cholestatic liver diseases

Yokoda RT *et al.* Pathogenesis of CLD

Raquel T Yokoda, Eduardo A Rodriguez

**Abstract**

Cholestatic liver diseases (CLD) begin to develop after an impairment of bile flow start to affect the biliary tree. Cholangiocytes actively participate in the liver response to injury and repair and the intensity of this reaction is a determinant factor for the development of CLD. Progressive cholangiopathies may ultimately lead to end-stage liver disease requiring at the end orthotopic liver transplantation. This narrative review will discuss cholangiocyte biology and pathogenesis mechanisms involved in four intrahepatic CLD: Primary biliary cholangitis, primary sclerosing cholangitis, cystic

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### Cholestatic liver diseases: An era of emerging therapies