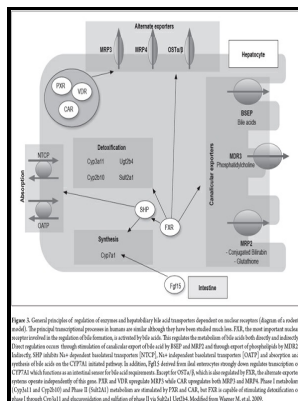


Molecular pathogenesis of cholestasis

Landes Bioscience/Eurekah.com - CiteSeerX — Review Intrahepatic cholestasis of pregnancy: molecular pathogenesis, diagnosis and management



Description: -

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Molecular Biology -- methods.
Cholestasis -- physiopathology.
Cholestasis -- metabolism.
Cholestasis -- genetics.
Molecular biology.
Cholestasis -- Pathogenesis.
Cholestasis -- Molecular aspects.
Cholestasis.Molecular pathogenesis of cholestasis
-Molecular pathogenesis of cholestasis
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Molecular Pathology of Cholestasis

In addition, localization of transporters to the cell membrane may be disrupted due to changes in microtubules. In some cases, the actual P value is presented. Black arrowheads indicate hepatic steatosis; white arrowheads indicate cytoplasmic clearing and severe hepatocyte swelling.

The molecular pathogenesis of cholestasis in sepsis

Intrahepatic cholestasis of pregnancy ICP is a reversible form of cholestasis in late pregnancy, persisting until delivery.

Molecular Pathogenesis of Cholestasis

BA profiling using ultra-HPLC—mass spectrometry For BA profiling, 20 μ L of serum was mixed with 60 μ L of acetonitrile 1% NH₄OH, including lithocholic acid-D 5 as the internal standard, and supernatant was separated by centrifugation. Additional types of PFIC with TJP5 and NR1H4 mutations have recently been designated PFIC4 and PFIC5, respectively. Cholestasis, defined as a hereditary or an acquired disruption of bile flow, is a liver disease caused by reduced bile acid BA efflux from hepatocytes intrahepatic or bile duct obstruction extrahepatic, resulting in BA accumulation and serious liver injury.

Intrahepatic cholestasis of pregnancy

New cases and follow-up from infancy to adult age. Pruritus itching is a common symptom of pregnancy, affecting around 23% of women.

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