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May 01, 2014 · Disease name. Polycystic liver disease (PLD) is a collection of rare human disorders that result from structural changes in the biliary tree development [1,2].Genetic mechanisms and/or signaling defects are the root cause of ductal structures to become separated from the biliary tree finally resulting in cyst formation [2,3].Typically, these disconnected biliary structures are present in a ...

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## Polycystic liver disease

Polycystic liver disease usually describes the presence of multiple cysts scattered throughout normal liver tissue. PLD is commonly seen in association with autosomal-dominant polycystic kidney disease, with a prevalence of 1 in 400 to 1000, and accounts for 8–10% of all cases of end stage renal disease. The much rarer autosomal-dominant polycystic liver disease will progress without any kidney involvement.

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## Polycystic Liver Disease: Symptoms, Causes, Treatments

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You may be diagnosed with polycystic liver disease if: You have a family member with **PLD**, are under the age of 40, and have more than one cyst. You have a family member with **PLD**, are older than 40, and have more than three cysts. You have no family members with **PLD**, are over the age of 40, and have more than 20 cysts.

**Author:** Hedy Marks

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<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC6924961>

Dec 20, 2019 · Establish the **Diagnosis**. **Polycystic liver disease** (PLD) is characterized by more than 20 fluid-filled biliary epithelial-lined cysts in the **liver**. 1 The majority of PLD cases occur as an extrarenal manifestation of autosomal dominant **polycystic** kidney **disease** (ADPKD) caused by mutations in PKD1 and PKD2. 2 A separate condition, autosomal dominant **polycystic liver disease** (ADPLD), is caused ...

**Author:** Armani Patel, Arlene B. Chapman, Ad... **Publish Year:** 2019

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