

Name of Journal: *World Journal of Clinical Cases*

Manuscript NO: 38709

Manuscript Type: Case report

Polycystic kidney and hepatic disease 1 gene mutations in von Meyenburg complexes: Case report

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Abstract

Von Meyenburg complexes (VMCs) are a rare type of ductal plate malformations. We herein report two Chinese families with VMCs and the suspicious gene mutation of this disease. Proband A was a 62-year-old woman with abnormal echographic presentation of the liver. She received

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Von Meyenburg complexes, or biliary hamartomas, are frequently incidentally detected. ... imaging (MRI) is the reference investigation tool for complex **cases**. ... **polycystic liver** and **kidney diseases**, Caroli **disease** and Caroli syndrome, **ARPKD** is caused by **mutations** of PKHD1, a **gene** located on chromosome 6, coding ...

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Polycystic **liver disease-1** is an autosomal dominant condition characterized by ... See also PCLD2 (617004), caused by **mutation** in the SEC63 **gene** (608648) on ... normal but contained microcysts and typical **von Meyenburg complexes** from ... was found only with adult **PKD** and was not observed in **cases** of only PCLD.

Von Meyenburg complex: case report and literature review - Scielo.br

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The present **study reports** a case of **von Meyenburg complex** and discusses the ... the association with