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Clinical characteristics and ABCC2 genotypic in Dubin-Johnson syndrome: A case report and review of literature

Huan Wu, Xue-Ke Zhao, Juan-Juan Zhu

Abstract

BACKGROUND

Dubin-Johnson syndrome (DJS) is a benign autosomal recessive liver disease by the mutation of *ABCC2* gene that is characterized by chronic or intermittent conjugated hyperbilirubinemia, with chronic idiopathic jaundice as the main clinical manifestation. Genetic alterations of the *ABCC2* gene are commonly used for diagnosing DJS; however,

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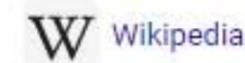
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