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A molecular overview of Progressive Familial Intrahepatic Cholestasis

PFIC on a molecular overview

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Progressive familial intrahepatic cholestasis 2 ...

https://www.ncbi.nlm.nih.gov/gtr/conditions/C3489789

Mild ATP8B1 deficiency is characterized by **intermittent episodes of cholestasis manifest** as **severe pruritus** and **jaundice**; **chronic liver damage** does not typically develop.

Expanding etiology of progressive familial intrahepatic ...

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