Progressive familial intrahepatic cholestasis type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Progressive familial intrahepatic cholestasis type 1. ORPHA:79306

PFIC1, a type of progressive familial intrahepathic cholestasis (PFIC, see this term), is an infantile hereditary disorder in bile formation that is hepatocellular in origin and associated with extrahepatic features.