Belgian Genetic Tests database

DISEASE:
Caroli disease

NAME:	Caroli disease
DESCRIPTION:	Caroli disease (CD) is a rare congenital liver disease characterized by non-obstructive cystic dilatations of the intra-hepatic and rarely extra-hepatic bile ducts.
ORPHACODE:	53035
XREF(S):	Orphanet MedDRA MeSH MeSH OMIM ICD-10
ANALYTE(S):	PKHD1
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/2994

RELATED CONTENT

Related Genetic Tests

- Caroli Disease
- Cholestasis (gene panel)
- Ciliopathy (gene panel)
- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)

Related Laboratories

- Centre de Génétique Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Gent

Related Analytes

- PKHD1 ciliary IPT domain containing fibrocystin/polyductin

Related Gene Panels

- Cholestasis (40 genes) - UCL
- Ciliopathy (120 genes) - UGent
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG

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