DISEASE:
Familial colorectal cancer Type X

NAME:	Familial colorectal cancer Type X
DESCRIPTION:	A rare inherited cancer-predisposing syndrome characterized by the fulfilment of the Amsterdam criteria for hereditary nonpolyposis colorectal cancer (HNPCC) but without alterations, either somatic or germline, in the DNA mismatch repair (MMR) genes.
ORPHACODE:	440437
SYNONYMS:	FCCTX
XREF(S):	Orphanet ICD-10 ICD-10 ICD-10 ICD-10 ICD-10 ICD-10 ICD-10 ICD-10
ANALYTE(S):	BMPR1A SEMA4A RPS20

CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Colon carcinoma (hereditary/familial) (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- bone morphogenetic protein receptor type 1A
- ribosomal protein S20
- semaphorin 4A

Related Gene Panels

- Colon carcinoma (hereditary/familial) (gene panel) (12 genes) - VUB