Rodzinny rak jelita grubego typu X
Kod Orpha: 440437 Kod OMIM:

Opis choroby *

Definicja
A rare, hereditary nonpolyposis colon cancer defined in individuals meeting the Amsterdam criteria for Lynch syndrome, but lacking germline mutations in the mismatch repair genes. It is characterized by a later onset, preferential involvement of distal colon and rectum, lower risk of developing extracolonic cancer, a higher adenoma/carcinoma ratio, a higher differentiation of tumor cells, a more heterogeneous tumor architecture and an infiltrative growth pattern, when compared to Lynch syndrome cases.
Dane

Klasyfikacja
Choroba

Synonimy
FCCTX
FCCTX

Kod ORPHA
440437

Kod OMIM
-

Kod ICD10
C18.7

Kod ICD11
-

---

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.