

Rodziny 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.