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
-

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*Źródło

orphanet