

Opis choroby *

Definicja

A rare inherited cancer-predisposing syndrome characterized by predisposition to a wide variety of cancers, including neoplasms of the digestive tract, urinary tract, kidney, endometrium, ovary, brain, and prostate, as well as sebaceous skin tumors, depending on the gene involved. Tumors may occur at any age but often arise in young people. Factors influencing individual tumor risk include sex, age, affected gene, and personal history of cancer.

Dane

Klasyfikacja

Choroba

Kod ORPHA

144

Kod OMIM

614385

Kod ICD10

D48.9

Kod ICD11

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*Źródło

orphanet