

Zespół Muira i Torrego

Kod Orpha: 587 Kod OMIM: 158320

Opis choroby *

Definicja

A form of hereditary nonpolyposis colon cancer characterized by the development of cutaneous sebaceous neoplasia and at least one visceral malignancy, most frequently gastrointestinal carcinoma. The malignancies are usually multiple, occur at an early age, but tend to be of low-grade and have a relatively low incidence of metastases. Sebaceous tumors are usually multiple, with sebaceous adenomas being the commonest. Multiple keratoacanthomas, usually located on the face or the trunk, have been reported as a feature. Cutaneous tumors may precede or follow the first presentation of internal malignancy, which usually involves the gastrointestinal tract, the breast or the genitourinary tract.

Dane

Klasyfikacja

Choroba

Synonimy

Multiple keratoacanthoma, Muir-Torre type
Rogowiak kolczystokomórkowy typu Muira i Torrego

Kod ORPHA

587

Kod OMIM

158320

Kod ICD10

L72.8

Kod ICD11

2C31.1

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

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