

Opis choroby *

Definicja

A genodermatosis characterized by the presence of multiple hamartomas in various tissues and an increased risk for malignancies of the breast, thyroid, endometrium, kidney and colorectum. When CS is accompanied by germline *PTEN* mutations, it belongs to the PTEN hamartoma tumor syndrome (PHTS) group.

Dane

Klasyfikacja

Choroba

Synonimy

Cowden disease

Choroba Cowdena

Zespół mnogich zmian typu hamartoma

Multiple hamartoma syndrome

Kod ORPHA

201

Kod OMIM

616858

Kod ICD10

Q85.8

Kod ICD11

LD2D.Y

*Źródło

orphanet