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

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*Źródło

orphanet