Opis choroby *

Definicja

A rare subtype of mixed epithelial-mesenchymal tumor, often presenting as a large, exophytic polypoid lesion, which may extend through the cervix, composed of benign or atypical epithelium and low-grade malignant stroma. It usually presents with dysfunctional bleeding or vaginal discharge and less often abdominal pain. Association with long-term unopposed estrogen therapy, tamoxifen therapy and a history of pelvic radiation has been reported.

Dane

Klasyfikacja
Choroba

Kod ORPHA
213600

Kod OMIM
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Kod ICD10
C54.2

Kod ICD11
2B5D.1

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*Źródło

orphanet