Opis choroby *

Definicja

A rare genetic, malignant breast tumor characterized by early onset breast cancer in association with a germline mutation. Tumors arising in carriers of BRCA1 and BRCA2 mutations differ morphologically and genetically from each other, as well as from sporadic breast cancers. Most BRCA1-associated tumors are invasive ductal adenocarcinomas of no special type, typically of higher grade than sporadic tumors, and more often negative for hormone receptors. In addition, more cases with features of typical or atypical medullary carcinoma are seen in these patients. Likewise, BRCA2-associated tumors tend to be of higher grade than sporadic ones, although their phenotype is similar. They show a low frequency of HER-2 expression.

Dane

Klasyfikacja
Choroba

Synonimy
Familial breast cancer
Rodzinny rak piersi
Familial breast carcinoma
Hereditary breast carcinoma

Kod ORPHA
227535

Kod OMIM
613399

Kod ICD10
C50.0

Kod ICD11
2C6Y

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*Źródło

orphanet