Opis choroby *

Definicja

A rare multiple endocrine neoplasia (MEN) syndrome that is principally characterized by the association of medullary thyroid carcinoma (MTC) with other endocrine tumors. The variant MEN 2A is defined by MTC associated with pheochromocytoma and/or primary hyperparathyroidism (MEN2A); the variant MEN 2B is defined as an aggressive form of MTC in association with pheochromocytoma but without primary hyperparathyroidism.

Dane

Klasyfikacja Choroba Synonimy MEN2 MEN2

Kod ORPHA

653

Kod OMIM 155240

Kod ICD10 D44.8

Kod ICD11 2F7A.0

<u>*Źródło</u>

orphanet