

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

*Źródło

orphanet