Multiple endocrine neoplasia type 2

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Multiple endocrine neoplasia type 2. ORPHA:653

Multiple endocrine neoplasia type 2 (MEN2) is a multiple endocrine neoplasia (see this term), a polyglandular cancer syndrome characterized by the occurrence of medullary thyroid carcinoma (MTC), pheochromocytoma (PCC; see these terms), in one variant, primary hyperparathyroidism (PHPT). There are three forms: MEN2A, MEN2B, and familial medullary thyroid carcinoma (FMTC) (see these terms).