Multiple endocrine neoplasia type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Multiple endocrine neoplasia type 1. ORPHA:652

Multiple endocrine neoplasia Type 1 (MEN1) is a frequent form of MEN (see this term), a rare inherited cancer syndrome, characterized by the development of neuroendocrine tumors of the parathyroid, pancreas, and anterior pituitary gland, and less commonly the adrenal cortical gland, with other non-endocrine tumors in some patients.