Multiple endocrine neoplasia type 4

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Multiple endocrine neoplasia type 4. ORPHA:276152

Multiple endocrine neoplasia type 4 (MEN4) is a very rare form of MEN (see this term), an inherited cancer syndrome, characterized by parathyroid and anterior pituitary tumors, possibly associated with adrenal, renal, and reproductive organ tumors.