VIPoma

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. VIPoma. ORPHA:97282

VIPoma is an extremely rare type of pancreatic neuroendocrine tumor (see this term) that secretes vasoactive intestinal polypeptide (VIP) leading to the manifestations of watery diarrhea, hypokalemia and achlorhydia or hypochhlorhydia (known as WDHA syndrome).