Hereditary pheochromocytoma-paraganglioma

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Hereditary pheochromocytoma-paraganglioma. ORPHA:29072

Hereditary paraganglioma-pheochromocytomas (PGL/PCC) are rare neuroendocrine tumors represented by paragangliomas (occurring in any paraganglia from the skull base to the pelvic floor) and pheochromocytomas (adrenal medullary paragangliomas; see this term).