Opis choroby *

Definicja

A rare, isolated, non-familial pheochromocytoma/paraganglioma tumor arising from neuroendocrine chromaffin cells of the adrenal medulla (pheochromocytoma) or from extra-adrenal chromaffin tissue (paraganglioma). The majority of these tumors are benign and the presenting symptoms are typically caused by the increased catecholamine production of the tumor, including hypertension (often paroxysmal), tachycardia, anxiety and/or excessive sweating.

Dane

Klasyfikacja
Choroba

Kod ORPHA
276621

Kod OMIM
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Kod ICD10
C74.1

Kod ICD11
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*Źródło

orphanet