

Opis choroby *

Definicja

A rare, hereditary, pheochromocytoma/paraganglioma tumor arising from neuroendocrine chromaffin cells of the adrenal medulla (pheochromocytoma) or from any paraganglia from the skull base to the pelvic floor (paraganglioma). Clinical manifestations are often linked to excess catecholamines production causing sustained or paroxysmal elevations in blood pressure, headache, episodic profuse sweating, palpitations, pallor and apprehension or anxiety. Hereditary pheochromocytoma/paraganglioma tumors tend to present at younger ages, to be multi-focal, bilateral, and recurrent, or to have multiple synchronous neoplasms.

Dane

Klasyfikacja

Choroba

Synonimy

Familial pheochromocytoma-paraganglioma
Rodzinny guz chromochłonny - przyzwojak

Kod ORPHA

29072

Kod OMIM

614165

Kod ICD10

D35.6

Kod ICD11

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*Źródło

orphanet