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