Opis choroby *

Definicja

A rare neuroendocrine tumor arising from chromaffin cells of the adrenal medulla (pheochromocytoma) or from sympathetic and parasympathetic ganglia (paraganglioma). These tumors are most often benign and may produce catecholamines in excess causing hypertension and sometimes severe acute cardiovascular complications.

Dane

Klasyfikacja
Grupa fenomenów

Kod ORPHA
573163

Kod OMIM
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Kod ICD10
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Kod ICD11
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*Źródło

orphanet