Opis choroby *

Definicja

A rare pituitary tumor originating from normally hormone-producing cells of the adenohypophysis, characterized by a sellar or extrasellar mass manifesting with clinical signs secondary to mass effect, but without evidence for hormonal hypersecretion. Typical manifestations are visual disturbances, headaches, cranial nerve dysfunction, and hypopituitarism but the mass may also be discovered incidentally.

Dane

Klasyfikacja
Choroba

Synonimy
NFPA
NFPA

Kod ORPHA
91349

Kod OMIM
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Kod ICD10
D35.2

Kod ICD11
2F37.0

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*Źródło

orphanet