

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

*Źródło

orphanet