Opis choroby *

Definicja

A rare pituitary disease characterized by hemorrhagic or non-hemorrhagic necrosis of the pituitary gland. Clinical manifestations typically comprise sudden and severe headache (often with nausea and vomiting), visual disturbances (visual-field defects, loss of visual acuity), oculomotor palsies, and variable degrees of altered consciousness, ranging from lethargy to coma. Acute endocrine dysfunction may also be present, most commonly corticotropic deficiency with severe hypotension and hyponatremia as well as secondary adrenal failure, but also thyrotropic and gonadotropic deficiency.

Dane

Klasyfikacja
Choroba

Synonimy
Pituitary tumor apoplexy
Pituitary tumor apoplexy

Kod ORPHA
95613

Kod OMIM
-

Kod ICD10
E23.6

Kod ICD11
5A61.0

---

*Źródło

orphanet