Panhypophysitis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Panhypophysitis. ORPHA:95513

Panhypophysitis is a rare, acquired pituitary hormone deficiency, a type of primary hypophysitis characterized by an inflammation of the entire pituitary gland. Common clinical presentation is diabetes insipidus with polyuria and polydipsia and partial or panhypopituitarism. Other symptoms may include headaches, nausea/vomiting, visual disturbances and fatigue.