Opis choroby *

Definicja

A rare, acquired, endocrine disease characterized by the triad of diffuse skin and mucosa hyperpigmentation, markedly elevated serum adrenocorticotropin (ACTH) levels and an enlarging corticotroph adenoma, which manifest following total bilateral adrenalectomy performed for the treatment of Cushing's disease. Additionally, patients may present with headaches, visual field defects, cranial nerve palsy, pituitary apoplexy, diabetes insipidus, panhypopituitarism, and, occasionally, paraovarian or paratesticular tumors.

Dane

Klasyfikacja
Zespół kliniczny

Kod ORPHA
199244

Kod OMIM
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Kod ICD10
E24.1

Kod ICD11
5A70.3

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*Źródło

orphanet