Opis choroby *

Definicja

A rare cranial malformation characterized by hyperostosis frontalis interna, variably associated with metabolic and endocrine disorders (such as obesity, diabetes mellitus, and hirsutism, among others). Compression by calvarial thickening may lead to cerebral atrophy and present with cognitive impairment, neuropsychiatric symptoms, headaches, and epilepsy. The condition predominantly affects women.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Hyperostosis frontalis interna
Hiperostoza czołowa wewnętrzna

Kod ORPHA
77296

Kod OMIM
144800

Kod ICD10
M85.2

Kod ICD11
FB80.3

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*Źródło

orphanet