

## **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

---

#### \*Źródło

orphanet