

## Opis choroby \*

### Definicja

A rare endocrine disease characterized by neonatal hypoglycemia, prolonged cholestatic jaundice, and seizures. Typical are low plasma ACTH and cortisol levels in the absence of structural pituitary defects, and sometimes low partial growth hormone deficiency is associated.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

199296

#### Kod OMIM

201400

#### Kod ICD10

E23.6

#### Kod ICD11

5A74.Y

---

[\\*Źródło](#)

orphanet