

## Opis choroby \*

### Definicja

A rare inborn error of metabolism characterized by congenital hypertryptophanemia and hyperserotonemia. Patients are typically asymptomatic, although developmental delay, intellectual disability, and behavioral abnormalities, among others, have been reported in association.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

2224

#### Kod OMIM

600627

#### Kod ICD10

E70.8

#### Kod ICD11

5C50.3

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### \*Źródło

orphanet