

## Opis choroby \*

### Definicja

Congenital atransferrinemia is a very rare hematologic disease caused by a transferrin (TF) deficiency and characterized by microcytic, hypochromic anemia (manifesting with pallor, fatigue and growth retardation) and iron overload, and that can be fatal if left untreated.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Congenital hypotransferrinemia

Wrodzona hipotransferynemia

#### Kod ORPHA

1195

#### Kod OMIM

209300

#### Kod ICD10

E88.0

#### Kod ICD11

5D0Y

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#### [\\*Źródło](#)

orphanet