

## Opis choroby \*

### Definicja

A rare inborn error of metabolism characterized by a deficiency in erythrocyte catalase, an enzyme responsible for the breakdown of hydrogen peroxide. The disorder is usually asymptomatic but may be associated with oral ulcerations and gangrene, or diabetes mellitus and atherosclerosis in certain populations.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Catalase deficiency

Niedobór katalazy

#### Kod ORPHA

926

#### Kod OMIM

614097

#### Kod ICD10

E80.3

#### Kod ICD11

5C57.1

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#### \*Źródło

orphanet