

## Opis choroby \*

### Definicja

A disorder of glyoxylate metabolism characterized by an excess of oxalate resulting in kidney stones, nephrocalcinosis and ultimately renal failure and systemic oxalosis. There are 3 types of PH, types 1-3, all caused by liver-specific enzyme defects.

### Dane

### Klasyfikacja

Choroba

**Kod ORPHA**

416

**Kod OMIM**

613616

**Kod ICD10**

E74.8

**Kod ICD11**

5C51.2Y

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[\\*Źródło](#)

orphanet