

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

*Źródło

orphanet