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