DISEASE:
Primary hyperoxaluria type 3

NAME:	Primary hyperoxaluria type 3
DESCRIPTION:	Primary hyperoxaluria type 3 (PH3) is a disorder of glyoxylate metabolism that can be asymptomatic or characterized by oxalate nephrolithiasis.
ORPHACODE:	93600
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	HOGA1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Hyperoxaluria
- Tubulopathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- 4-hydroxy-2-oxoglutarate aldolase 1

Related Gene Panels

- Tubulopathy/Nephrolithiasis (106 genes) - IPG