DISEASE:
Dent disease type 2

NAME:	Dent disease type 2
DESCRIPTION:	A rare genetic renal tubular disease, characterized by manifestations of proximal tubule dysfunction with low-molecular-weight (LMW) proteinuria, hypercalciuria, nephrolithiasis, nephrocalcinosis, and progressive renal failure. Extra-renal involvement is frequent, but may be mild and not recognized.
ORPHACODE:	93623
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	OCRL
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Cataract (gene panel)
- Genetic disorders of Calcium and Phosphate metabolism (gene panel)

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- Centrum Medische Genetica - UZ Gent
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Related Analytes

- OCRL inositol polyphosphate-5-phosphatase

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