Belgian Genetic Tests database

DISEASE:
Gitelman syndrome

NAME:	Gitelman syndrome
DESCRIPTION:	A rare syndrome characterized by hypokalemic metabolic alkalosis in combination with significant hypomagnesemia and low urinary calcium excretion.
ORPHACODE:	358
XREF(S):	Orphanet MeSH MedDRA OMIM ICD-10
ANALYTE(S):	SLC12A3 CLCNKB
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/549

RELATED CONTENT

Related Genetic Tests

- Tubulopathy (gene panel)

Related Laboratories

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

Related Analytes

- chloride voltage-gated channel Kb
- solute carrier family 12 member 3

Related Gene Panels

- Tubulopathy/Nephrolithiasis (106 genes) - IPG
- essai