

DISEASE:
Bartter syndrome type 3

NAME:	Bartter syndrome type 3
DESCRIPTION:	A form of Bartter syndrome characterized by a later age at onset than the other types of Bartter syndrome, typically presenting beyond the first year of life with failure to thrive, hypokalemic and hypochloremic metabolic alkalosis, increased levels of plasma renin and aldosterone and low to normal blood pressure.
ORPHACODE:	93605
SYNOMYS:	Bartter syndrome type III
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	CLCNKB
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/3527>

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](#)

Related Gene Panels

- [Tubulopathy/Nephrolithiasis \(106 genes\) - IPG](#)

Source URL: <http://gentest.healthdata.be/disease/3527>