DISEASE:
Liddle syndrome

NAME:	Liddle syndrome
DESCRIPTION:	A rare genetic form of low-renin hypertension characterized by hypertension associated with decreased plasma levels of potassium and aldosterone.
ORPHACODE:	526
SYNONYMS:	Pseudoaldosteronism Pseudohyperaldosteronism type 1
XREF(S):	Orphanet MeSH MedDRA MedDRA OMIM ICD-10 OMIM OMIM
ANALYTE(S):	SCNN1B SCNN1G SCNN1A
CREATED:	13 May 2019 - 01:02

CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/510

RELATED CONTENT

Related Genetic Tests

- Bronchiectasies with or without elevated sweat chloride panel (5 genes)

Related Laboratories

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

Related Analytes

- sodium channel epithelial 1 subunit alpha
- sodium channel epithelial 1 subunit beta
- sodium channel epithelial 1 subunit gamma

Related Gene Panels

- Cystic Fibrosis / Liddle syndrome / Pseudohypoaldosteronism type 1 (3 genes) - ULG
- Pulmonary/Bronchiectasies (5 genes) - IPG