

ANALYTE:
SCNN1A

NAME:	sodium channel epithelial 1 subunit alpha
SYMBOL:	SCNN1A
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ENaCalpha amiloride-sensitive sodium channel subunit alpha
XREF(S):	Orphanet Genatlas HGNC OMIM Reactome SwissProt Ensembl
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/analyte/1115>

RELATED CONTENT

Related Genetic Tests

- [Bronchiectasies with or without elevated sweat chloride panel \(5 genes\)](#)
- [Bronchiectasis \(4 genes\)](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Nephropathies, hereditary \(gene panel\)](#)
- [Primary cardiac arrhythmias \(Atrial fibrillation / Brugada syndrome / Catech. polymorphic ventricular tachycardia / Early repolarisation syndrome / Idiopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome\) \(gene panel\)](#)
- [Respiratory disorders \(gene panel\): non-CF bronchiectasis; pulmonary hypertension; interstitial lung disease](#)
- [Tubulopathy \(gene panel\)](#)

Related Diseases

- [Brugada syndrome](#)
- [Generalized pseudohypoaldosteronism type 1](#)
- [Idiopathic bronchiectasis](#)
- [Liddle syndrome](#)

Related Gene Panels

- [Bronchiectasis \(4 genes\) - UCL](#)
- [Bronchiectasis with or without elevated sweat chloride \(4 genes\)](#)
- [Cystic Fibrosis / Liddle syndrome / Pseudohypoaldosteronism type 1 \(3 genes\) - ULG](#)
- [Nephropathies, hereditary \(219 genes\) - KUL](#)
- [Nephropathy panel - UGent](#)

- Panel Nephro-ULG-V1
- Primary cardiac arrhythmias (113 genes) - VUB
- Pulmonary/Bronchiectasies (5 genes) - IPG
- Respiratory Disorders panel (137 genes) - Ugent
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

Source URL: <http://gentest.healthdata.be/analyte/1115>