GENETIC TEST:
Cystic Fibrosis / Congenital absence of the vas deferens / CFTR-related disorders (88 hot spot mutations)

FULL NAME:	Cystic Fibrosis / Congenital absence of the vas deferens / CFTR-related disorders (88 hot spot mutations)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Dried blood spot card, Peripheral (whole) blood on EDTA, Chorionic villi, Amniotic fluid
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	PCR based technique
RIZIV CODE:	565353-565364
ACCREDITATION (ISO 15189):	2021-03-25 / 2024-09-09

EQA:	• Cystic Fibrosis, • Cystic Fibrosis, • Cystic Fibrosis, • Cystic Fibrosis, • Cystic Fibrosis, • Cystic fibrosis, • Cystic fibrosis, • Cystic fibrosis
TURNAROUND TIME (MAXIMUM):	3 - weeks
CREATED:	18 Jul 2019 - 15:04
CHANGED:	19 Jan 2024 - 09:10
URL:	https://ulbgenetics.be/compendium-des-analyses/#GM56

RELATED CONTENT

Related Diseases

- Congenital bilateral absence of vas deferens
- Cystic fibrosis
- Hereditary chronic pancreatitis
- Idiopathic bronchiectasis

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB

Related Analytes

- CF transmembrane conductance regulator