GENETIC TEST:
Cystic Fibrosis / Congenital absence of the vas deferens / related disorders (50 hot spot mutations)

FULL NAME:	Cystic Fibrosis / Congenital absence of the vas deferens / related disorders (50 hot spot mutations)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	PCR based technique
RIZIV CODE:	565353-565364
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02

EQA:	• Cystic Fibrosis, • Cystic fibrosis
TURNAROUND TIME (MAXIMUM):	2-4 week
CREATED:	18 Jul 2019 - 15:04
CHANGED:	01 Mar 2023 - 14:56
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/13903

Source URL: http://gentest.healthdata.be/genetic_test/75

RELATED CONTENT

Related Diseases

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

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- CF transmembrane conductance regulator

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