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

FULL NAME:	Cystic Fibrosis / Congenital absence of the vas deferens / CFTR-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, Chorionic villi, Amniotic fluid
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	PCR based technique
RIZIV CODE:	565353-565364
ACCREDITATION (ISO 15189):	2023-11-09 / 2024-05-08

EQA:	• Cystic Fibrosis, • Cystic Fibrosis, • Cystic Fibrosis, • Cystic Fibrosis, • Cystic Fibrosis, • Cystic fibrosis, • Cystic fibrosis, • Cystic fibrosis, • Cystic Fibrosis (CF) and CFTR-related disorders
TURNAROUND TIME (MAXIMUM):	1 month
CREATED:	18 Jul 2019 - 15:04
CHANGED:	22 Jan 2024 - 13:40
URL:	https://labogidsmedgen.uza.be/analyses/mucoviscidose

RELATED CONTENT

Related Diseases

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

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- CF transmembrane conductance regulator