GENETIC TEST:
Cystic Fibrosis / Congenital absence of the vas deferens / CFTR-related disorders (Sequencing CFTR gene)

FULL NAME:	Cystic Fibrosis / Congenital absence of the vas deferens / CFTR-related disorders (Sequencing CFTR gene)
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:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565471-565482
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
TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	18 Jul 2019 - 14:57
CHANGED:	22 Jan 2024 - 13:39
URL:	https://labogidsmedgen.uza.be/analyses/mucoviscidose

Source URL: http://gentest.healthdata.be/genetic_test/518

RELATED CONTENT

Related Diseases

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

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen

Related Analytes

- CF transmembrane conductance regulator