GENETIC TEST:
Cystic Fibrosis / Congenital bilateral absence of vas deferens (CBAVD) / Idiopathic pancreatitis (50 recurrent mutations)

FULL NAME:	Cystic Fibrosis / Congenital bilateral absence of vas deferens (CBAVD) / Idiopathic pancreatitis (50 recurrent 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:	Next Generation Sequencing (NGS)
RIZIV CODE:	565353-565364
ACCREDITATION (ISO 15189):	2021-09-11 / 2026-09-10

EQA:	• Cystic Fibrosis, • Cystic Fibrosis, • Cystic Fibrosis, • Cystic fibrosis
TURNAROUND TIME (MAXIMUM):	4 weeks
CREATED:	18 Jul 2019 - 15:04
CHANGED:	16 May 2022 - 09:39

RELATED CONTENT

Related Diseases

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

Related Laboratories

- Centrum Medische Genetica - UZ Gent

Related Analytes

- CF transmembrane conductance regulator