# GENETIC TEST:
# Cystic Fibrosis / related disorders (50 hot spot mutations)

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| **FULL NAME:** | Cystic Fibrosis / related disorders (50 hot spot mutations) |
| **TEST TYPE:** | Clinical |
| **TEST SPECIALTY:** | Molecular Genetics |
| **TEST PURPOSE:** | Carrier diagnosis,<br>Newborn screening,<br>Post-natal Diagnosis,<br>Prenatal diagnosis |
| **SPECIMEN:** | Dried blood spot card,<br>Peripheral (whole) blood on EDTA,<br>Chorionic villi,<br>Amniotic fluid |
| **METHOD CATEGORY:** | Targeted variant analysis |
| **METHOD TECHNIQUE:** | PCR based technique |
| **RIZIV CODE:** | 565353-565364 |
| **ACCREDITATION (ISO 15189):** | 2022-12-22 / 2027-12-21 |

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| **EQA:** | • Cystic Fibrosis,<br>• Cystic Fibrosis,<br>• Cystic Fibrosis,<br>• Cystic Fibrosis,<br>• Cystic Fibrosis,<br>• Cystic fibrosis,<br>• Cystic fibrosis,<br>• Cystic fibrosis,<br>• Cystic fibrosis |
| **TURNAROUND TIME (MAXIMUM):** | 1 month |
| **CREATED:** | 18 Jul 2019 - 15:04 |
| **CHANGED:** | 19 Jan 2024 - 11:54 |

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# RELATED CONTENT

## Related Diseases

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

## Related Laboratories

- Centre de Génétique Médicale UCL

## Related Analytes

- CF transmembrane conductance regulator