

LASTNAME, FIRSTNAME

DOB: mm/dd/yyyy

Account Number:00000000



Patient ID:

Age: 00

Ordering Physician:

Specimen ID: 000-000-0000-0

Sex: Female

Date Collected: mm/dd/yyyy

Date Received: mm/dd/yyyy

Date Reported: mm/dd/yyyy

Date Entered: mm/dd/yyyy

Specimen Type: Whole Blood

Ethnicity: Not Provided

Indication: Carrier Test / Screening

## Cystic Fibrosis (CF) Full-gene Carrier Screen

**Summary: ■ POSITIVE**

## SAMPLE REPORT

### Variants Detected

Disorder (Gene)	Result	Interpretation
Cystic fibrosis (CFTR) NM_000492.3	POSITIVE: CARRIER Heterozygous for c.1521_1523delCTT (p.Phe508del), pathogenic, CHR7:117199645-117199648.	Predicted to be a carrier. <b>Risk:</b> AT INCREASED RISK FOR AFFECTED PREGNANCY. If this individual's reproductive partner is also a carrier of a pathogenic variant in the same gene, the risk for an affected fetus is 25%. Genetic counseling and reproductive partner carrier screening is recommended.

### Recommendations

Genetic counseling is recommended to discuss the potential clinical and/or reproductive implications of positive results, as well as recommendations for testing family members and, when applicable, this individual's partner. Genetic counseling services are available. To access Labcorp Genetic Counselors please visit <https://womenshealth.labcorp.com/genetic-counseling> or call (855) GC-CALLS (855-422-2557).

### Additional Clinical Information

Cystic fibrosis (CF) is an autosomal recessive disorder with variable severity and age at onset. Signs and symptoms of classic CF may include elevated sweat chloride levels, progressive lung disease, pancreatic insufficiency, and male infertility. Symptoms of mild CF may include pancreatic sufficiency. Symptoms of CFTR-related disorders may include pancreatitis, bronchiectasis, and isolated male infertility due to congenital absence of the vas deferens (CBAVD). Treatment is dietary and supportive. Genotype-targeted therapies may be available for some individuals. In severely affected individuals, lung transplantation may be indicated. (Ong, PMID:20301428).

### Comments

This interpretation is based on the clinical information provided and the current understanding of the molecular genetics of the disorder(s) tested. Information about the disorder(s) tested is available at <https://womenshealth.labcorp.com>.

### Methods/Limitations

**Next-generation Sequencing:** Genomic regions of interest are selected using the Twist Biosciences® hybridization capture method and sequenced via the Illumina® next generation sequencing platform. Sequencing reads are aligned with the human genome reference GRCh37/hg19 build. Regions of interest include coding exons, intron/exon junctions (+/- 20 nucleotides) and additional genomic regions with known significant pathogenic variants. Analytical sensitivity at 30X coverage is estimated to be >99% for single nucleotide variants, >99% for insertions/deletions less than six base pairs and >96% for insertions/deletions between six and forty-five base pairs. Variant detection is performed by QIAGEN CLC Genomics and in-house algorithms. Expected minimum size resolution for CNVs in CFTR is 60 bp of coding sequence. Precise breakpoints are not reported. Single-exon deletions or duplications are not detected in some cases due to CNV size limitations, or due to isolated data quality variation or intrinsic sequence properties. Confirmatory testing by orthogonal technologies includes Sanger sequencing, MLPA analysis.

**Reported variants:** Pathogenic and likely pathogenic variants are reported for all tests. Benign and likely benign variants are typically not reported. Variants of uncertain significance are reported when included in the test specification. Variants are specified using the numbering and nomenclature recommended by the Human Genome Variation Society (HGVS, <http://www.hgvs.org/>). Variant classification and confirmation are consistent with ACMG standards and guidelines (Richards, PMID:25741868; Rehm, PMID:23887774). Detailed variant classification information and reevaluation are available upon request.

**Limitations:** Technologies used do not detect germline mosaicism and do not rule out the presence of large chromosomal aberrations including rearrangements and gene fusions, or variants in regions or genes not included in this test, or possible inter/intragenic interactions between variants, or repeat expansions. Variant classification and/or interpretation may change over time if more information becomes available. False positive or false negative results may occur for reasons that include: rare genetic variants, sex chromosome abnormalities, pseudogene interference, blood transfusions, bone marrow transplantation, somatic or tissue-specific mosaicism, mislabeled samples, or erroneous representation of family relationships.

### References

Deignan JL, Astbury C, Cutting GR *et al*. CFTR variant testing: a technical standard of the American College of Medical Genetics and Genomics (ACMG). *Genet Med* 22, 1288 (2020). PMID: 32404922

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Specimen ID: 000-000-0000-0

Sex: Female

# Cystic Fibrosis (CF) Full-gene Carrier Screen

## References (Cont.)

Ong T, Marshall SG, Karczeski BA, et al. Cystic Fibrosis and Congenital Absence of the Vas Deferens. 2001 Mar26 [Updated 2017 Feb 2]. In: Adam MP, Ardinger HH, Pagon RA, et al., editors. GeneReviews® [Internet]. PMID: 20301428

Rehder C, Bean LJH, Bick D. et al. Next-generation sequencing for constitutional variants in the clinical laboratory, 2021 revision: a technical standard of the American College of Medical Genetics and Genomics (ACMG). *Genet Med* 23, 1399 (2021). PMID: 33927380

Richards S, Aziz N, Bale S et al. Standards and guidelines for the interpretation of sequence variants: a joint consensus recommendation of the American College of Medical Genetics and Genomics and the Association for Molecular Pathology. *Genet Med* 17, 405 (2015). PMID: 25741868

## Performing Labs

Component Type	Performed at	Laboratory Director
Technical component, processing	Laboratory Corporation of America, 1912 TW Alexander Drive, RTP, NC 27709-0150	Laboratory Director
Technical component, analysis	Laboratory Corporation of America, 1912 TW Alexander Drive, RTP, NC 27709-0150	Laboratory Director
Professional component	Laboratory Corporation of America, 1912 TW Alexander Drive, RTP, NC 27709-0150	Laboratory Director

For inquiries, the physician may contact the lab at 800-345-4363

This test was developed and its performance characteristics determined by Labcorp. It has not been cleared or approved by the Food and Drug Administration.

### Patient Details

LASTNAME, FIRSTNAME

Phone:

Date of Birth: mm/dd/yyyy

Age: 00

Sex: Female

Patient ID:

Alternate Patient ID:

### Physician Details

CLIENT NAME

CLIENT ADDRESS

Phone: 00000000

Account Number: 00000000

Physician ID:

NPI:

### Specimen Details

Specimen ID: 000-000-0000-0

Control ID:

Alternate Control Number:

Date Collected: mm/dd/yyyy 0000 Local

Date Received: mm/dd/yyyy 0825 ET

Date Entered: mm/dd/yyyy 1508 ET

Date Reported: mm/dd/yyyy 1411 ET

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