

THIS IS NOT A TEST REQUEST FORM. Please complete and submit with the test request form or electronic packing list.

PATIENT HISTORY FOR CYSTIC FIBROSIS (CF) TESTING

Patient Name: _____ **Date of Birth:** _____ **Sex:** Female Male
Ordering Provider: _____ **Provider's Phone:** _____
Practice Specialty: _____ **Provider's Fax:** _____
Genetic Counselor: _____ **Counselor Phone:** _____

Patient's Ethnicity (check all that apply)

- Black or African American Asian Hispanic or Latino Native American or Other Pacific Islander
 Ashkenazi Jewish White Middle Eastern Other: _____

Is the patient pregnant? No Yes N/A

Does the patient have symptoms? No Yes (check all that apply)

- Azoospermia COPD Pancreatitis
 Bilateral absence of the vas deferens Failure to thrive Pneumonia
 Bronchiectasis Fetal echogenic bowel Positive newborn screen
 Chronic Cough Meconium ileus Pseudomonas
 Other Symptoms: _____

Has sweat chloride testing been performed? No Yes Unknown

If yes, what was the result? normal (<30) borderline (30-60) elevated (>60) QNS Unknown

Has the patient undergone previous DNA testing for CF? No Yes Unknown

If yes, describe the test(s) and results: _____

Does the patient have a family history of CF? No Yes Unknown

If yes, specify the relationship of the family member to the patient _____

Indicate if the relative is: a healthy carrier affected with CF List CF variant(s): _____

Is the patient's reproductive partner a CF carrier? No Yes If yes, list the variant: _____

Does the patient's reproductive partner have a family history of CF? No Yes Unknown

If yes, specify the relationship of family member(s) to the partner: _____

Is the partner's relative a healthy carrier or affected?

Check the test you intend to order.

- 2013661 Cystic Fibrosis (CFTR) 165 Pathogenic Variants:** Recommended for carrier screening and as first line diagnostic test in symptomatic patients. Clinical sensitivity for carrier detection is 78% in African American/Black, 96% in Ashkenazi Jewish, 55% in Asian American, 92% in Caucasian, and 80% in Hispanic patients.
 2013664 Cystic Fibrosis (CFTR) 165 Pathogenic Variants w/Reflex to Sequencing w/Reflex to Deletion/Duplication: Test begins with 165 variants and reflexes until two pathogenic variants are identified. Clinical sensitivity is 99%.
 0051640 Cystic Fibrosis (CFTR) Sequencing with Reflex to Deletion/Duplication: *CFTR* gene sequencing; if two pathogenic variants are not identified, deletion/duplication testing is performed. Clinical sensitivity is 99%.
 2001961 Familial Mutation, Targeted Sequencing: Tests for a mutation previously identified in a family member; a copy of relative's lab result is REQUIRED.
 3003144 Deletion/Duplication Analysis by MLPA: Tests large *CFTR* deletion/duplication previously identified in a family member; copy of relative's lab report is REQUIRED.

Master Label

For questions, contact an ARUP genetic counselor at 800-242-2787 ext. 2141.