

ATTACHMENT 3

21 December 2020

Dr. Anthony S. Fauci, Director
National Institute of Allergy and Infectious Diseases
5601 Fishers Lane
Rockville, MD 20892
301- 496 - 5717

**Subject : I Hereby Accuse You of 'Gross Criminal Negligence'
Connectable to the Death of Mr. Spencer William Smith**

2 Pages

Mission:

To protect, promote & improve the health of all people in Florida through integrated state, county & community efforts.



Ron DeSantis
Governor

Scott A. Rivkees, MD
State Surgeon General

Vision: To be the **Healthiest State** in the Nation

Mandatory Reporting of COVID-19 Laboratory Test Results: Reporting of Cycle Threshold Values

December 3, 2020

Laboratories are subject to mandatory reporting to the Florida Department of Health (FDOH) under section 381.0031, Florida Statutes, and Florida Administrative Code, Chapter 64D-3.

- All positive, negative and indeterminate COVID-19 laboratory results must be reported to FDOH via electronic laboratory reporting or by fax immediately. This includes all COVID-19 test types—polymerase chain reaction (PCR), other RNA, antigen and antibody results. For a list of county health departments and their reporting contact information, please visit www.FLhealth.gov/chdepcontact.
- Cycle threshold (CT) values and their reference ranges, as applicable, must be reported by laboratories to FDOH via electronic laboratory reporting or by fax immediately.

As per Florida Administrative Code, rule 64D-3.031, laboratories must report all of the following:

- The patient's:
 - First and last name, including middle initial
 - Address (including street, city, state and ZIP code)
 - Telephone number (including area code)
 - Date of birth
 - Sex
 - Race
 - Ethnicity (Hispanic or non-Hispanic)
 - Pregnancy status, if applicable
 - Social Security number
- The laboratory:
 - Name, address and telephone number of laboratory performing test
 - Type of specimen (e.g., stool, urine, blood, mucus, etc.)
 - Date of specimen collection
 - Specimen collection site (e.g., cervix, eye) if applicable
 - Date of report
 - Type of test performed and results, including reference range, titer when quantitative procedures are performed and all available results on speciation, grouping or typing of organisms
- The submitting provider's:
 - Name
 - Address (including street, city, state and ZIP code)
 - Telephone number (including area code)
 - National provider number (NPI)

If your laboratory is not currently reporting CT values and their reference ranges, the lab should begin reporting this information to FDOH within seven days of the date of this memorandum. If your laboratory is unable to report CT values and their reference ranges, please fill out the [brief questionnaire attached](#) to this memorandum and submit by facsimile to the FDOH's Bureau of Epidemiology confidential fax line at 850-414-6894, within seven days of the date of this memorandum

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Attachment

Name of person completing questionnaire	
Name of laboratory	
Street address	
City, state and ZIP code	

1. Is your laboratory a CLIA-certified laboratory performing diagnostic molecular testing for the detection of SARS-CoV-2?
- Yes
 No

2. Does your laboratory perform multiple assays for the molecular detection of SARS-CoV-2?
- Yes
 No

3. Please list all the platforms/assays that your laboratory uses.

4. Do the molecular assays your laboratory performs include real-time PCR with the test result being based on a CT value?
- Yes
 No (Your survey is complete, please fax to 850-414-6894)

5. Please select all the reason(s) why your laboratory is not able to report the CT value to FDOH.
- Although the qualitative result is generated based on a CT value, the assay/instrument does not provide the user with the actual CT value—it only provides the qualitative result
 The laboratory does not have a separate mechanism to report the CT value to FDOH since the CT value does not get reported to the submitting provider
 Other (please list the reasons)

Fax to 850-414-6894