

Coronavirus Disease 2019 (COVID-19)

Interim Guidelines for Collecting, Handling, and Testing Clinical Specimens from Persons for Coronavirus Disease 2019 (COVID-19)

Summary of Recent Changes

Revisions were made on March 21, 2020 to reflect the following:

A new document, Completing a CRF and Specimen Guidance [2 pages], has been posted that contains
instructions for completing a supplemental guidance regarding specimen collection, storage, and shipping to Cl
laboratories.

Revisions were made on March 19, 2020. to reflect the following:

- Allowance for OP as an acceptable specimen type if NP swabs are not available.
- Standard operating procedure for public health labs to create their own viral transport media [5 pages] in accordance with CDC's protocol.

Revisions were made on March 17, 2020. to reflect the following:

- Recommendation for collection for testing of lower respiratory tract specimens.
- Updated description of collecting a Nasopharyngeal swab.

Revisions were made on March 13, 2020. to reflect the following:

• Recommendation to collect and test a single upper respiratory nasopharyngeal swab (NP).

March 19, 2020

Health care providers should contact their local/state health department immediately to notify them of patients with fever and lower respiratory illness who they suspect may have COVID-19. Local and state public health staff will determine if the patient meets the criteria for testing for COVID-19. The state and local health department will assist clinicians to collect, store, and ship specimens appropriately, including during afterhours or on weekends/holidays. Clinical specimens should be collected for routine testing of respiratory pathogens at either clinical or public health land Note that clinical laboratories should NOT attempt viral isolation from specimens collected from persons suspected thave COVID-19 unless this is performed in a BSL3 laboratory. Testing for other pathogens by the provider should be done as part of the initial evaluation but should not delay testing for COVID-19.

Specimen Type and Priority

For initial diagnostic testing for COVID-19, CDC recommends collecting and testing an upper respiratory nasopharyng swab (NP). Collection of oropharyngeal swabs (OP) is a lower priority and if collected should be combined in the san tube as the NP. Collection of only OP swab is acceptable if other swabs are not available.. Collection of sputum should only be done for those patients with productive coughs. Induction of sputum is not recommended. Specimens should collected as soon as possible once a PUI is identified, regardless of the time of symptom onset. Maintain proper infection of the control when collecting specimens.

CDC also recommends testing lower respiratory tract specimens, if available. For patients who develop a productive cough, sputum should be collected and tested for SARS-CoV-2. The induction of sputum is not recommended. For patients for whom it is clinically indicated (e.g., those receiving invasive mechanical ventilation), a lower respiratory tract specimen.

Specimens should be collected as soon as possible once a PUI is identified, regardless of the time of symptom onset. Maintain proper infection control when collecting specimens. See Biosafety FAQs for handling and processing specimens uspected cases and PUIs.

General Guidelines

Store specimens at 2-8°C and ship overnight to CDC on ice pack. Label each specimen container with the patient's ID number (e.g., medical record number), unique specimen ID (e.g., laboratory requisition number), specimen type (e.g. serum) and the date the sample was collected. Complete a CDC Form 50.34 for each specimen submitted. In the upp left box of the form, 1) for *test requested* select "Respiratory virus molecular detection (non-influenza) CDC-10401" are for *At CDC*, *bring to the attention of* enter "Stephen Lindstrom: 2019-nCoV PUI".

 Please refer to our instruction guidance for submitting CDC Form 50.34 found here: Completing a CRF and Spec Guidance [2 pages]

I. Respiratory Specimens

A. Lower respiratory tract

Bronchoalveolar lavage, tracheal aspirate

Collect 2-3 mL into a sterile, leak-proof, screw-cap sputum collection cup or sterile dry container.

Sputum

Have the patient rinse the mouth with water and then expectorate deep cough sputum directly into a sterile, leak-proscrew-cap sputum collection cup or sterile dry container.

B. Upper respiratory tract

Nasopharyngeal swab (NP) /oropharyngeal swab (OP)

Use only synthetic fiber swabs with plastic shafts. Do not use calcium alginate swabs or swabs with wooden shafts, at they may contain substances that inactivate some viruses and inhibit PCR testing. Place swabs immediately into steri tubes containing 2-3 ml of viral transport media. In general CDC is now recommending collecting only the NP swab. both swabs are used, NP and OP specimens should be combined at collection into a single vial. OP swabs remain an acceptable specimen type.

Nasopharyngeal swab: Insert a swab into nostril parallel to the palate. Swab should reach depth equal to distance from nostrils to outer opening of the ear. Leave swab in place for several seconds to absorb secretions. Slowly remove swab while rotating it.

Oropharyngeal swab (e.g., throat swab): Swab the posterior pharynx, avoiding the tongue.

Nasopharyngeal wash/aspirate or nasal aspirate

Collect 2-3 mL into a sterile, leak-proof, screw-cap sputum collection cup or sterile dry container.

II. Storage

Store specimens at 2-8°C for up to 72 hours after collection. If a delay in testing or shipping is expected, store specin at -70°C or below.

III. Shipping

Specimens PUI's must be packaged, shipped, and transported according to the current edition of the International Ai Transport Association (IATA) Dangerous Goods Regulationsexternal icon . Store specimens at 2-8°C and ship overr to CDC on ice pack. If a specimen is frozen at -70°C ship overnight to CDC on dry ice. Additional useful and detailed information on packing, shipping, and transporting specimens can be found at Interim Laboratory Biosafety Guidelin for Handling and Processing Specimens Associated with Coronavirus Disease 2019 (COVID-19).

For additional information, consultation, or the CDC shipping address, contact the CDC Emergency Operations Cente (EOC) at 770-488-7100.

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