



Please, PRINT; *indicates required fields

Patient Information

Name* (last) _____ (first) _____ (initial) _____

DOB* ____/____/____ Phone #* () -

Address* _____

City* _____ State* _____ Zip* _____ Patient Medical Record Number*: _____

Sex:* M F

(mark all applicable)

- Ethnicity*: Hispanic/Latino Non-Hispanic/Non-Latino Unknown
 Race*: White Black/African American Asian
 American Indian/Alaska Native Native Hawaiian/Other Pacific Islander Other

Submitter Information

Practitioner Name* (last) _____ (first) _____ (initial) _____

NPI _____ Facility Name* _____

Phone # () - Fax #* () - *Include the fax number in which you would like patient results sent.

Address* _____

City* _____ State* _____ Zip* _____

Email: _____

Testing Priority* (MDHHS COVID-19 Testing Prioritization Criteria)

- Priority 1: Hospitalized patients, healthcare workers with symptoms
- Priority 2: Patients in long-term care facilities with symptoms, >65 years old with symptoms, underlying conditions with symptoms, first responders with symptoms
- Priority 3: Critical infrastructure workers with symptoms, individuals with mild symptoms
- Priority 4: Asymptomatic individuals
- Non-symptomatic preoperative testing

Clinical Information

Diagnosis _____

Onset (mm-dd-yyyy) ____/____/____

Antibiotics (list and start dates) _____

Specimen Information

Collection Date (mm-dd-yyyy)* ____/____/____ Time (hour:minute) _____ AM / PM By: _____

Source/Type* (check one only)

- Nasopharynx
- Other (specify): _____

**Specimens must be kept between 2-8° C from collection until delivery to testing facility. If the sample will not reach our facility within 72 hours, please freeze. ** Minimum specimen volume of 2ml.

For initial diagnostic testing for COVID-19, CDC recommends collecting and testing an upper respiratory specimen. Nasopharyngeal specimen is the preferred choice for swab-based SARS-CoV-2 testing. When collection of a nasopharyngeal swab is not possible, the following are acceptable alternatives:

- An oropharyngeal (OP) specimen collected by a healthcare professional, or
- A nasal mid-turbinate (NMT) swab collected by a healthcare professional or by onsite self-collection (using a flocked tapered swab), or
- An anterior nares (nasal swab; NS) specimen collected by a healthcare professional or by onsite self-collection (using a flocked or spun polyester swab).

After collection, specimen

was** (check one only)

- Frozen
- Refrigerated

Test Request

x COVID-19

*FDA recommends that test reports include a general statement that the test has been validated but the FDA's independent review of this validation is pending.