



**LABORATORY ALLIANCE**

of Central New York, LLC

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## SARS-CoV-2 and Flu A/Flu B Molecular Assay

Effective February 7, 2022, Laboratory Alliance's Microbiology Department will offer a new multiplex gene amplification assay to detect SARS-CoV-2, influenza A and influenza B in provider-collected nasopharyngeal (N/P) and anterior nasal (nasal) swab specimens as well as self-collected patient swab specimens. These specimens should be collected from individuals suspected of respiratory viral infection consistent with COVID-19 and/or influenza.

**Background:** SARS-CoV-2, influenza A and influenza B are common causes of respiratory illness. Clinical signs and symptoms of these respiratory infections are often similar requiring the use of a laboratory test for proper diagnosis. The SARS-CoV-2 and Flu A/Flu B test is a multiplex assay based on the use of transcription-mediated amplification, similar to PCR, that reliably detects the presence of these viruses in N/P and nasal specimens. Specimens are collected in the usual manner and placed in either Universal Transport Medium (UTM), Viral Transport Medium (VTM), or MedSchenker STM.

**Test Name: COVID/FLU AB AMPLIFI (SARS-CoV-2, Flu A, Flu B by RT-TMA)**

**Test Code: CFNAAT**

**Method:** Nucleic Acid Amplified Test (NAAT): Real-Time Transcription Mediated Amplification (RT-TMA)

**Specimen Collection:** Collect N/P or nasal swab

**Specimen Transport:** Transport N/P or nasal specimens at room temperature. Specimens placed in transport medium following collection can be stored for 24 hours at 15 to 30°C or store specimens refrigerated at 2 to 8°C for up to 7 days.

**Unacceptable Specimens:** The use of any swab other than Dacron or flocked swabs and any transport medium other than UTM, VTM or MedSchenker.

**Schedule:** Daily

**CPT Code:** 87636

**Billing Code:** 3010469

## Interpretation of Results:

**Negative result** indicates that the virus was not detected but does not rule out the presence of disease

**Positive result** indicates that the virus was detected and may be responsible for the respiratory infection

## Preferred Specimen Collection Device(s):

**UTM - Universal Transport Media (Viral, Chlamydia, Mycoplasma, Ureaplasma Culture, Bordetella pertussis)**



**Viral Transport Medium**



**MedSchenker STM 3 ml tube and NP swab**

