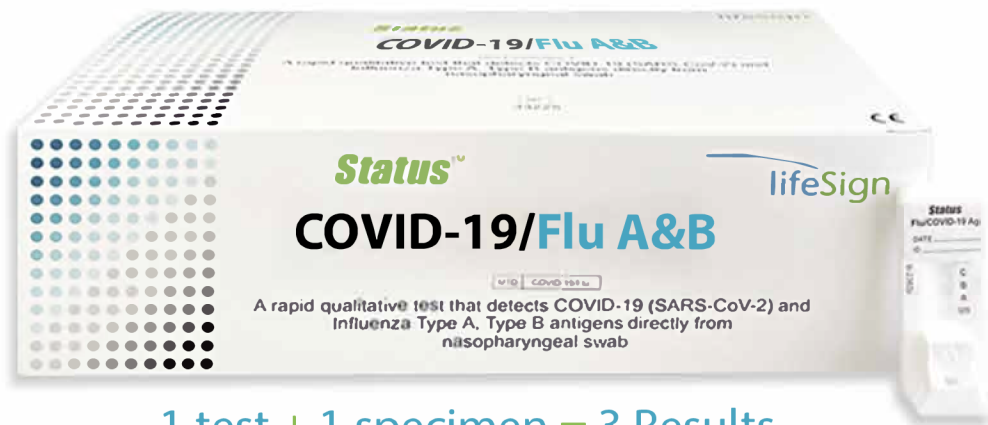
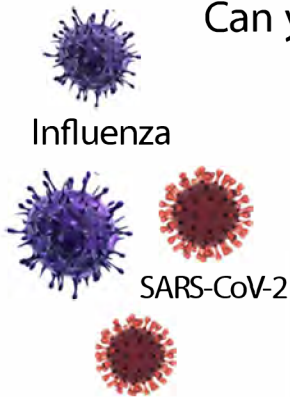


# Status™

# COVID-19/Flu A&B

A Rapid Immunoassay for the Simultaneous Direct Detection and Differential Diagnosis of SARS-CoV-2, Influenza Type A and Type B Antigen from anterior nasal and nasopharyngeal swab specimens

Infections with these viruses may present similar symptoms. Can you tell them apart? **We Can.**



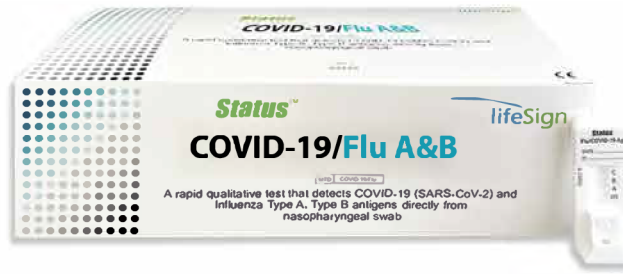
Learn More

1 test + 1 specimen = 3 Results



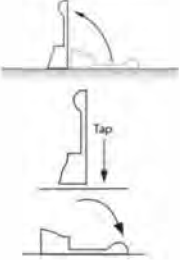
- **COVID-19 - Anterior Nasal - Sensitivity 93.8%, Specificity 100%**  
**Nasopharyngeal - Sensitivity 93.1%, Specificity 100%**
- **Flu A - Sensitivity 91.4%, Specificity 95.7%**
- **Flu B - Sensitivity 87.6%, Specificity 95.9%**
- **FDA Emergency Use Authorization (EUA)**
- **Visually read in 15 minutes**
- **Flocked swabs for superior specimen collection and patient comfort**

#### ORDERING INFORMATION

| PRODUCT                  | # OF TESTS | ITEM # |
|--------------------------|------------|--------|
| Status™ Covid-19/Flu A&B | 25         | 33225  |



## Easy 3 Step Procedure

|   |  |  |
|---|--|--|
| 1 |  <p>squeeze</p>   | <p>Tear the tab off the Extraction Reagent Capsule and dispense entire contents into the Extraction Well.</p>  |
| 2 |   | <p>Insert the specimen swab in the Swab Stand.</p> <ul style="list-style-type: none"> <li>• <b>Rotate</b> swab 3 times to mix the specimen.</li> <li>• <b>Let stand</b> 1 minute.</li> <li>• <b>Rotate</b> swab 3 times again and discard the swab.</li> </ul> |
| 3 |  <p>Tap</p>  | <p>Raise the device upright and <b>let stand</b> 1–2 seconds.</p> <p>Gently <b>tap</b> the device to ensure the liquid flows into the hole.</p> <p>Lay the device back down.</p> <p style="text-align: center;"><b>Set a timer for 15 minutes.</b></p>         |
| 4 | <p><b>Read test results at 15 minutes.</b></p> <p><b>NOTE: False positive or false negative results can occur if the test is not read between 15 and 20 minutes.</b></p> |  |

- Status™ COVID-19/FLU A&B is an antigen test for use with anterior nasal and nasopharyngeal swab specimens.
- Status™ COVID-19/FLU A&B test has been granted emergency use authorization (EUA) by the FDA for use at the point-of-care setting. During the duration of the emergency this test can be performed in a patient care setting that is operating under a CLIA Waiver, Certificate of Compliance or Certificate of Accreditation.



Proudly made in the USA.

For more information or to place an order, contact



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