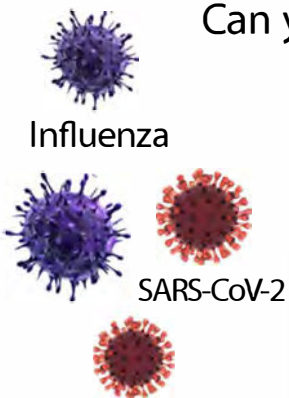


Status™

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**

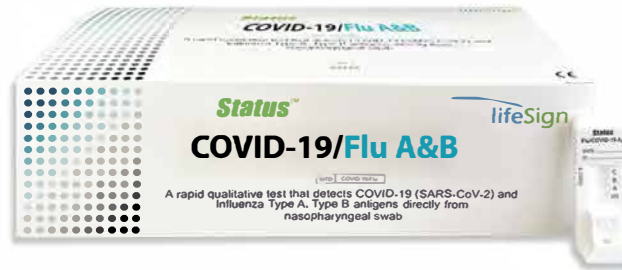
ORDERING INFORMATION		
PRODUCT	# OF TESTS	ITEM #
Status™ Covid-19/Flu A&B	25	33225

Proudly made in the USA



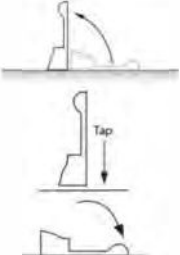
For more information or to place an order, contact:

A PBM Group Company

85 Orchard Road • Skillman • New Jersey • 08558
800.526.2125 • 732.246.3366 • Fax: 732.246.0570 • www.lifesignmed.com



Easy 3 Step Procedure

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

- Refer to our website at www.lifesignmed.com for complete test procedure details, training video and package insert.
- 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.

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