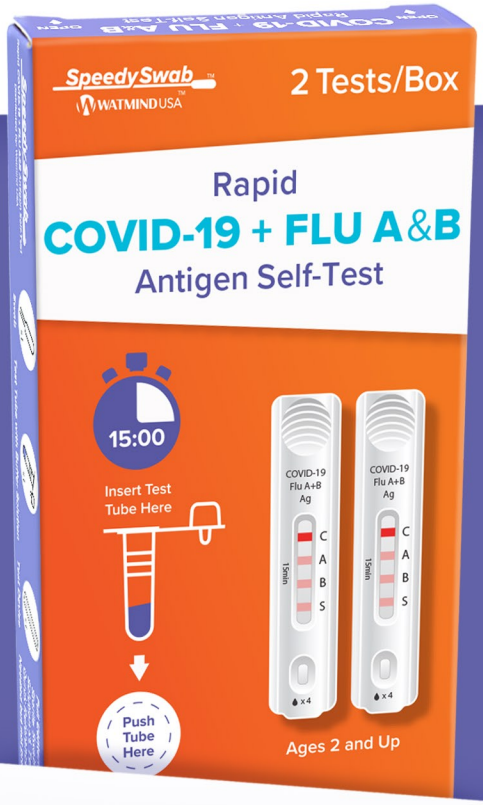
Distributed by:
WATMIND USA™

FDA EUA Approval

SpeedySwab™

Rapid COVID-19 + FLU A&B Antigen Self-Test



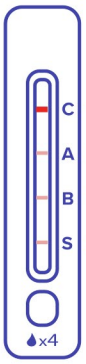
SpeedySwab™ is an FDA EUA authorized Rapid COVID-19, influenza A and Influenza B Antigen Self-Test designed for nasal application and is highly effective in detecting all COVID-19 variants and sub-variants with a class leading COVID-19 Clinical Sensitivity of 92.5%.

The test uses the long-proven Lateral Flow Assay (LFA) technology. In three easy steps, simply swirl the soft tip of the nasal swab ½" – ¾" into each nostril for 15 seconds, insert the swab into a test tube with a buffer solution, and apply 4 drops onto the test card. Results will be available in 15 minutes.

Available with a 18-Month Shelf Life.

Positive Test Result

Flu A+B+ COVID-19



C = Control Line
A = Influenza A Line
B = Influenza B Line
S = COVID-19 (SARS-CoV 2)

Product Benefits

- **Reliable** design supporting highly sensitive detection of Influenza A and B, and COVID-19 genetic variants and sub-variants
- **Simple** format is designed for self-administration on adults, or by adults on children ages 2 and up for ease of use and provides results in 15 minutes
- Our **production capabilities** give us the ability to quickly produce large quantities of tests, allowing us to provide for the market as demand increases

Available Products

1 Test/Box

2 Tests/Box

4 Tests/Box

25 Tests/Box

*In the USA, this product has not been FDA cleared or approved; but has been authorized by FDA under an Emergency Use Authorization. This product has been authorized only for the detection of proteins from SARS-CoV-2, influenza A and influenza B, not for any other viruses or pathogens. The emergency use of this product is only authorized for the duration of the declaration that circumstances exist justifying the authorization of emergency use of *in vitro* diagnostics for detection and/or diagnosis of COVID-19 under Section 564(b)(1) of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C § 360bbb-3(b)(1), unless the declaration is terminated or authorization is revoked sooner.