QUIDEL

October 1, 2021 to December 31, 2021

Buy 6 kits for the price of 5

Sofia® Influenza A+B FIA kits*

For use with Sofia 2 and Sofia
Quidel Catalog #20218

Sofia 2 Flu + SARS Antigen FIA kits**

For use with Sofia 2
Quidel Catalog #20377

FDA Emergency Use Authorization (EUA)

Current and new Sofia customers

ORDER NOW!

Contact your Quidel Account Manager to order your promotional test kits.
Complete the form and send it along with your invoice(s) to Quidel at

1.858.431.3513 or customerservice@quidel.com.

The promotional test kits will be sent directly to your office from Quidel.

Name_____

Contact_____ Telephone_____ E-Mail _____

Address_____

City_____ State_____ ZIP Code_____

***Minimum of five (5) Sofia Influenza A+B FIA kits must be invoiced during the promotional period by new or current Sofia customer in order to qualify for the discounted price. Promotion valid from October 1, 2021 through December 31, 2021. Form must be submitted within 90 days of promotional period end date to qualify for the promotion.**

****Minimum of five (5) Sofia 2 Flu + SARS Antigen FIA kits must be invoiced during the promotional period by new or current Sofia customer in order to qualify for the discounted price. Promotion valid from October 1, 2021 through December 31, 2021. Form must be submitted within 90 days of promotional period end date to qualify for the promotion.**

Please note that the value of the special offer(s) the Customer may receive from the manufacturer under this program is a "discount or other reduction in price" to Customer under Section 1128B(b)(3)(A) of the Social Security Act [42 U.S.C. 1320a-7b(b)(3)(a)]. Accordingly, Customer shall disclose this and any other discounts or other reductions in price received under this program under any state or federal program which provides cost or charge-based reimbursement to the Customer for the products and services purchased under this program. Quidel reserves the right to cancel this promotion at any time.

The Sofia 2 Flu + SARS Antigen FIA has not been FDA cleared or approved but has been authorized by the FDA under an EUA for use by authorized laboratories for the detection of proteins from SARS-CoV-2, and influenza, not for any other viruses or pathogens. This assay 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 authorization is terminated or revoked sooner.

FX1018001EN00 (10/21)