Sofia® 2

AMS | ATLANTIC MEDICAL SOLUTIONS

Buy 6 kits for the price of 5

Sofia SARS Antigen FIA*

For use with Sofia 2 and Sofia platforms
QuidelOrtho Catalog #20374
FDA Emergency Use Authorization (EUA)

Sofia 2 SARS Antigen+ FIA**

For use with Sofia 2 platform
QuidelOrtho Catalog #20405
CLIA waived

Contact your QuidelOrtho Account Manager to order your promotional test kits. Complete the form and send it along with your invoice(s) to QuidelOrtho at **858.455.4960** or **customerservice@quidelortho.com**.

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

Name

Telephone

Email

Address

City

State

ZIP code

Order now!

July 1, 2024 to September 30, 2024

*Minimum of five (5) Sofia SARS Antigen FIA kits must be invoiced during the promotional period by new or current Sofia customers in order to qualify for the discounted price. Promotion valid from July 1, 2024 to September 30, 2024. **Form must be submitted within 90 days of promotional period and date to qualify for the promotion**.

Minimum of five (5) Sofia 2 SARS Antigen+ FIA kits must be invoiced during the promotional period by new or current Sofia customers in order to qualify for the discounted price. Promotion valid from July 1, 2024 to September 30, 2024. **Form must be submitted within 90 days of promotional period and 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 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.

New QuidelOrtho branding may not be available in all markets, subject to country-specificregulatory approval. Please confirm with your local commercial team.

QuidelOrtho™

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