The BD Veritor™ Plus System sales event is on.

Save up to 20% on pediatric care value packs

1 kit = 30 tests • Only valid for assay combination advertised below through December 31, 2024

BD item number	Value pack includes:
256132	**Two** - BD Veritor™ System for Rapid Detection of Respiratory Syncytial Virus CLIA-waived kits
	Two - BD Veritor™ System for Rapid Detection of Group A Strep CLIA-waived kits

Learn more about the BD respiratory season offer at: **https://go.bd.com/pediatricvaluepackLP**

The right outcome begins with the right test...

Trust the rapid, digitally displayed results of the BD Veritor™ Plus System to guide your patients to the appropriate path to care.

To learn more, contact a sales representative to take advantage of this limited time offer today!
We're here to help. 1.844.823.5433

Offer valid through December 31, 2024. Offers cannot be combined. Promotion may not offer additional discounts for BD Veritor™ Plus System users under existing pricing contract/agreement. Offer is applicable to new and current BD Veritor™ Plus System users. The value of any rebates, discounts, or incentives provided may constitute a "discount or other reduction in price" under Section 1128B(b)(3)(A) of the Social Security Act (42 U.S.C. Sec.1320a-7b(b)(3)(A)). Customer shall satisfy any and all requirements imposed on buyers relating to discounts or reductions in price, including, when required by law, to disclose all discounts or other reductions in price received from BD and to accurately report under any state of federal healthcare program the net cost actually paid by customer.

* In the USA, the BD Veritor™ System for Rapid Detection of SARS-CoV-2 & Flu A+B has not been FDA cleared or approved but has been authorized by the FDA under an Emergency Use Authorization for use by authorized laboratories; use by laboratories certified under the CLIA, 42 U.S.C. §263a, that meet requirements to perform moderate, high, or waived complexity tests. The product is authorized for use at the Point of Care (POC), i.e., in patient care settings operating under a CLIA Certificate of Waiver, Certificate of Compliance, or Certificate of Accreditation.

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; and, in the USA, 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-1 9 under Section 564(b)(1) of the Federal Food, Drug and Cosmetic Act, 21 U.S.C. é360bbb-3(b)(1), unless the declaration is terminated or authorization is revoked sooner.

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