VACUETTE®



CE VACUETTE® Virus Stabilization Tube

For In Vitro Diagnostic Use



Intended Use

VACUETTE® Virus Stabilization Tubes are intended for the transport and storage of nasopharyngeal and oropharyngeal swab specimens. The product is to be used by healthcare professionals for SARS-CoV-2, influenza A and influenza B testing only.

Product Description

VACUETTE® Virus Stabilization Tubes are made of PET with a pre-defined volume of a phosphate buffered saline solution at a pH of 7.4±0.2 to allow for the storage of the swab specimens for up to 72h at 4°C. The tubes are fitted with **VACUETTE®** Safety Caps. The tube interior is sterile. The product is single use only and can be used on a single patient only.

Precautions/Cautions

- 1. Do not use tubes if foreign matter is present.
- 2. Do not use tubes after their expiration date.
- 3. Keep sterile tubes closed until use.
- Do not pool the additive of different VACUETTE® Virus Stabilization Tubes.
- 5. Do not use the additive from **VACUETTE**[®] Virus Stabilization Tubes for premoistening or prewetting the applicator swab before collecting the sample or for rinsing or irrigating the sampling sites.
- 6. Do not ingest the additive in the tube.
- 7. Specimens must be collected and handled using personal protective equipment against biological risk.
- 8. Handle all biological samples according to the policies and procedures of your facility.
- 9. Obtain appropriate medical attention in the case of any exposure to biological samples.
- 10. Ensure that after placing the specimen into the tube, that the cap is securely closed.
- 11. The tubes have a round bottom and are not self standing. Use of a sample rack is recommended.
- 12. The additive is transparent. Functionality of the product is not indicated by a color change.
- 13. Refer to the instructions for use of diagnostic assays for information on the correct sample material, correct storage and stability.
- 14. The user must validate the product when combining it with swabs, diagnostic kits or instruments before its use.

Storage

Storage guidelines for tubes before use

Store tubes at 4-25°C (40-77°F).

NOTE: Avoid exposure to direct sunlight. Exceeding the maximum recommended storage temperature may lead to impairment of the tube quality (i.e. drying out of liquid additives, coloring, etc.)

Limitations

- There is limited data available on test performance with specimens which have been frozen in any transport media; therefore, specimen stability should be validated if freezing is necessary. Please follow your facility protocol for this purpose.
- 2. The additive of the VACUETTE® Virus Stabilization Tube does not contain any RNase inhibitors.
- The VACUETTE® Virus Stabilization Tube has a dimension of 13x100mm. The length of the inserted swab should not exceed 90mm.

Specimen Collection and Handling

Please follow your institution's policy regarding the correct sample collection and processing.

Remove the cap from the tube by twisting in an anti-clockwise direction. Insert swab and close the tubes by twisting in a clockwise direction until firmly closed.

Please ensure a correct transport of the sample in the tube according to international regulations, such as UN3373.

Disposal

- The general hygiene guidelines and legal regulations for the proper disposal of infectious material should be considered and followed.
- 2. Disposable gloves and other personal protective equipment prevent the risk of infection.
- Contaminated or filled tubes must be disposed of in suitable biohazard disposable containers, which can then be autoclaved and incinerated afterwards. Follow your facility's protocol.
- 4. Disposal should take place in an appropriate incineration facility or through autoclaving (steam sterilization).

980360_Rev01_01-2021 Page 1 / 2





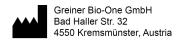
Label Information				
		Manufacturer	1	Temperature limit
		Use-by date	8	Do not re-use
	LOT	Batch code	[]i	Consult instructions for use
	REF	Catalogue number	IVD	In vitro diagnostic medical device
	STERILE R	Sterilized using irradiation		

References:

ISO / EN / ANSI/AAMI Standards

ISO 11137 "Sterilization of health care products – Requirements for validation and routine control – Radiation sterilization"

U.S. Food and Drug Administration. COVID-19 Testing Supplies: FAQs on Testing for SARS-CoV-2. Available at: https://www.fda.gov/medical-devices/coronavirus-covid-19-and-medical-devices/covid-19-testing-supplies-faqs-testing-sars-cov-2. Retrieved on November 25th, 2020.



www.gbo.com/preanalytics office@at.gbo.com Phone +43 7583 6791

980360_Rev01_01-2021 Page 2 / 2