



EN – Instruction for use

VWR Specimen Containers, PP, sterile

APPLICABLE PRODUCTS

					
CAT. NO.	VOLUME	SHAPE	CAP	SPECS	UOM
216-1853	60ML	CYLINDRICAL	RED	INDIV. WRAPPED	500/PAK
216-0857	120ML	CONICAL	BLUE	BULK	300/PAK
216-0862	120ML	CONICAL	GREEN	BULK	300/PAK
216-0856	120ML	CONICAL	RED	INDIV. WRAPPED	250/PAK
216-0861	120ML	CONICAL	RED	BULK	300/PAK
216-0860	120ML	CONICAL	WHITE	BULK	300/PAK
216-0869	120ML	CONICAL	WHITE	INDIV. WRAPPED	250/PAK
216-0892	120ML	CONICAL	RED WITH SPATULA	BULK	300/PAK
216-1161	120ML	CONICAL	YELLOW/VACUUM TUBES	INDIV. WRAPPED	250/PAK
216-2200	120ML	CONICAL	YELLOW/VACUUM TUBES	BULK	300/PAK

INTENDED PURPOSE

Collection of human specimens, other than blood.

SAMPLE MATERIAL

Sputum, urine and faeces.

DIAGNOSTICS

Diagnoses are only to be made by authorized and trained persons.

STORAGE CONDITIONS AND SHELF LIFE

Store in a dry place, away from sources of heat and direct sunlight.
The containers must be used by the expiry date stated.



















PRECAUTIONARY MEASURES ON HEALTH HAZARDS

Effective measures must be taken to protect against infection in line with laboratory guidelines.

INSTRUCTIONS FOR ENVIRONMENTAL DISPOSAL

Dispose of contents/container to an appropriate treatment and disposal facility in accordance with applicable laws and regulations, and product characteristics at time of disposal.

SYMBOLS USED

Symbol	Symbol title and	Symbol	Symbol title and description	Symbol	Symbol title and description
	CE marking of conformity		In vitro diagnostic medical device		Made in Italy
	Catalogue number		Product Quantity in the box		Keep dry
	Fragile, handle with care		Manufacturer		Keep away from sunlight
	Do not use if the packaging is		Unique Device Identification		Caution
	Consult instructions for use		Date of manufacture		Temperature limit
	Use-by date		Batch code		Sterilized using irradiation

DISCLAIMER

We pay the utmost attention to the quality of our products. In case of any malfunction or dissatisfaction please use the complaint form on the contact page of your local vwr.com website and indicate the reference and batch number of the product concerned. The attention of the users is drawn to the possible risks when the product is used for purposes other than what it is intended for. This IFU does not exempt the user from knowing and implementing all the information regulating their activity. The user will take under their responsibility the relevant precautions needed in the use that they make of this product.

CHANGE LOG

This instruction for use was established in January 2025 and cancels all previous instructions. The information is based on our knowledge and our experience on this day.

VERSION 04

Changes regarding latest IFU version:

- General update of the layout
- Minor corrections and clarifications

CONTACT



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Any serious incident that has occurred in relation to this product must be reported to the manufacturer and the Competent Authority of the Member State in which the user and/or the patient is established. Please contact Avantor's customer service department.