

Certificate of Conformity

VWR Catalogue Number	See List
Description	Conical Specimen Containers with Screw Caps, Capacity 120 mL
Country of Origin	Manufactured in Italy
Date of Issue (yyyy-mm-dd)	2025-07-11

Catalogue Number	Cap colour	Version	Type
216-0855	Red	Separate cap	Non-Sterile
216-0856	Red	Assembled cap	Sterile
216-0857	Blue	Assembled cap	Sterile
216-0858	Blue	Assembled cap	Non-Sterile
216-0859	Red	Separate cap	Non-Sterile
216-0860	White	Assembled cap	Sterile
216-0861	Red	Assembled cap	Sterile
216-0862	Green	Assembled cap	Sterile
216-0863	White	Assembled cap	Non-Sterile
216-0864	Red	Assembled cap	Non-Sterile
216-0865	Green	Assembled cap	Non-Sterile
216-0868	Red	Assembled cap	Non-Sterile
216-0869	White	Assembled cap	Sterile
216-1185	—	Without cap	Non-Sterile
216-3949	Green	Separate cap	Non-Sterile

Separate caps

Catalogue Number	For	Cap colour
216-1288	Urine container, 120 mL	Red

Quality System Compliance

This certificate confirms that the product has been sourced from a supplier or manufacturer who has declared their Quality Management System (QMS) complies with **ISO 9001:2015 & ISO 13485:2016**.

QC Testing

Representative products samples are collected and inspected in accordance with current applicable QC requirements.

Product Specifications

The product is in conformance with the requirements of the following standards and regulations:

Material	Container Cap	Polypropylene Polyethylene
FDA Statement	Raw material used to produce the products are listed in List of Authorised Substances of Regulation EU/10/2011. The material grades conform to EU Directive 2020/1245 on materials and articles intended to come into contact with food.	
Sterilization	Product labelled as sterile are sterilized using Beta rays as per ISO 11137-1 and meets SAL 10 ⁻⁶ with a dosage range of 17-20 kGy.	
ATP Assay	Not applicable.	
DNase & RNase Free	Not applicable.	
BSE/TSE	No use of any raw material produced from or substances derived from animal origin. The manufacturing process of the product does not use any ingredient of animal origin and no material derived from or exposed to animals affected by or under quarantine for transmitting Animal Bovine Spongiform Encephalopathy/Spongiform Encephalopathy (BSE)/(TSE).	
Cytotoxicity	Not applicable.	
Non-Pyrogenic Statement	Not applicable.	
Latex Statement	The products are not made from natural rubber latex.	
BPA Statement	Bisphenol A (BPA) has not been intentionally incorporated in the manufacturing of the products.	
DEHP Statement	Not applicable.	

RoHS	Not applicable.
REACH Statement	No substances as listed on >Substance of Very High Concern< candidate list are used in manufacturing of the raw materials used to produce this product. No routinely analysis is performed.
Storage Conditions	Store in a dry place, away from sources of heat and direct sunlight.
Shelf Life	60 months.

Disclaimer: VWR's Quality Management System (QMS) is ISO 9001:2015 certified as a wholesale distributor of finished products, ensuring effective oversight of finished product handling, traceability, and distribution processes. VWR is not a manufacturer and does not engage in manufacturing, co-manufacturing or processing activities such as design, packaging, labeling, testing or quality release. Consequently, the supplier or manufacturer's QMS certification may differ from VWR's.

VWR states that this declaration will not discharge the user from their obligation to ensure the product is suitable for the intended use. This product is intended for research use only. It is not for use in diagnostic procedures, therapeutic applications, or any human or clinical use.

This document has been produced electronically and is valid without a signature.

Version History

Sl.no	Version number	Description	Month & Year
1	1.0	Created General CoC using new layout	09-2021
2	1.1	FDA statement added	11-2023
3	1.2	Sterilization statement updated	04-2025
4	2.0	Modified Quality system compliance and disclaimer	07-2025