Certificate of Conformity

VWR Catalogue Number	331-0384
Description	HDPE Water sampling Bottle, with Tamper Evident Red Colour Screw Cap, 500 mL Capacity, 20mg/L Sodium Thiosulphate Dossage, Square shape, 33 mm Neck Size, 149.5 mm Height without cap, 71.5 mm length and width, Sterile
Lot Number	2205230058
Date of Sterilization (yyyy-mm-dd)	2022-05
Expiry Date (yyyy-mm-dd)	2024-04
Country of Origin	Manufactured in India
Date of Issue (yyyy-mm-dd)	2023-01-16

Quality System Compliance

Products are manufactured under the **ISO 9001:2015 & ISO 15378:2017** standard. Products are inspected and controlled through whole production processing in accordance with current applicable product specifications and QC SOP.

QC Testing

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

Test Name	Test Reference	Accuracy	Result
Leak Test	SOP/QA/07A	General Insp. Level-1, AQL -1.5	Leak Proof
Ethylene Oxide Dose	SOP/PRO/21	NA	PASS
Dimensional	SOP/QA/01	General Insp. Level-1, AQL -1.5	Within Tolerance

Product Specifications

Material	**Bottle**: High Density Polyethylene, 100% Virgin, FDA 21 CFR Compliance, USP Class VI Passed, Colour: Natural. **Closure with EP Liner:** Polypropylene, 100% Virgin, FDA 21 CFR Compliance, USP Class VI Passed. Colour: Red. Monomers and additives used to produce the products are listed in List of Authorised Substances of Regulation EU/10/2011. The material grades conform to EU Directive 1935/2004 on materials and articles intended to come into contact with food.
Sterilization	Product is ETO (Ethylene Oxide) irradiated as per ISO 11135:2014 and meets SAL 10^{-3}.
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	The product meets the requirement for polyolefin resins intended for food packaging application as described in FDA olefin polymer regulation 21 CFR 177.1520(c) 3.2a.
Non-Pyrogenic Statement	Not Applicable.
Latex Statement	This product is not made from natural rubber latex.
BPA Statement	Bisphenol are not used in the manufacture of the raw material and are not expected to be present.
DEHP Statement	Please see RoHS information below.
RoHS	No substances (Lead, Cadmium, Mercury, Hexavalent Chromium (Cr6+), Poly Brominated Biphenyls (PBB), Poly Brominated Diphenyl ethers (PBDE), Bis(2-Ethyhexyl) Phthalate (DEHP), Benzyl Butyl Phthalate (BBP), Dibutyl Phthalate (DBP), Diisobutyl Phthalate (DIBP)) are used in manufacturing the raw materials and final product. No routinely analyse is performed.

REACH Statement	No substances as listed on >Substance of Very High Concern< candidate list published 2022-06-10 are used in manufacturing of the raw materials used to produce this product. No routinely analyse is performed.
Storage Conditions	Store at 25 + 3 / - 8 °C, Relative Humidity 50 ± 10 % RH.

Disclaimer: VWR states that this declaration will not discharge the user from their obligation to ensure the product is suitable for the intended use. The purpose of the product is for use in laboratory only.
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