Certificate of Conformity

VWR Catalogue Number	**See List**
Description	**Dressing forceps, Purple Colour**
Country of Origin	**Manufactured in China**
Date of Issue (yyyy-mm-dd)	**2024-09-17**

Catalogue Number	Type	Length [mm]	Thichness [mm]	Width [mm]
232-0218	Non Sterile	110	1.5	9.9
232-0219	Sterile	110	1.5	9.9
232-0220	Non Sterile	130	1.6	13
232-0221	Sterile	130	1.6	13

Quality System Compliance

Products are manufactured under the **ISO 13485:2012** 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.

Inspected according to AQL level S-2 1.5 sampling plan per ISO 2859-1:1999.

Product Specifications

Material Acrylonitrile Butadiene Styrene.

Sterilization Product labeled as sterile is Ethylene Oxide (EO) irradiated according to AAMI/ISO 11135-1:2007 recommended practices in effect at the time of validation. Products labeled sterile meets a minimum requirement of 10^{-6} SAL (which is not applicable for non-sterile products).

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	Not Applicable.
BPA Statement	Not Applicable.
DEHP Statement	Not Applicable.
RoHS	Not Applicable.
REACH Statement	Not Applicable.
Storage Conditions	Store at room temperature.
Shelf Life	Product labelled as sterile has shelf life of 3 years and 5 years for non-sterile products.

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.
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 CoQ using new layout**	09-2021
2	2.0	**Removed** "Class I category device" information under product specification	09-2024