Certificate of Conformity

VWR Catalogue Number	734-2978
Description	Cell Lifters, Sterile, 234 mm Total length, 19 mm flat blade with 9 mm J-hook end
Lot Number	231209589A
Date of Manufacture (yyyy-mm-dd)	2023-12-09
Date of Sterilization (yyyy-mm-dd)	2023-12-11
Expiry Date (yyyy-mm-dd)	2026-12-09
Country of Origin	Manufactured in China
Date of Issue (yyyy-mm-dd)	2024-02-21

Quality System Compliance

Products are manufactured under the **ISO 9001:2015 & ISO 13485:2016** 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.

Product Specifications

The product is Class I category device as defined by FDA in 21CFR Parts 862-892.

Material	High Density Polyethylene (HDPE),Compliance with USP Class VI The composition of the product complies with FDA regulation 21 CFR 177.1520 (Olefin polymers), paragraphs (c) 2.1 and (c) 2.2.
Sterilization	Product labelled as sterile is EB (Electron Beam) irradiated and dose released upon ISO 11137 recommended practices at the time of validation and meets a minimum requirement of 10^{-6} SAL with a specified dose range of 15-30 kGy.
ATP Assay	Not Applicable.

DNase & RNase Free	This product is free of any detectable DNase/RNase contamination.
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). according to EMEA 410/01 revision 3.
Cytotoxicity	Testing is conducted to quality all material resins using ISO 10993 standards for cytotoxicity and have been shown to be non-toxic.
Non-Pyrogenic Statement	The acceptance level for product is 0.5 EU/ml or less than 20 EU/device. (TAL Gel Clot Method).
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	Not Applicable.
Storage Conditions	Room temperature

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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