Certificate of Conformity

VWR Catalogue Number	See List
Description	VWR® Cell Culture Dishes, Sterile
Country of Origin	Manufactured in China
Date of Issue (yyyy-mm-dd)	2025-07-23

Increased cell attachment treated surface

Catalogue Number	Type	Diameter [mL]	Height [mm]	Growth area [cm^2]
734-2814	Cell culture dish, increased cell attachment surface	35	12.3	8.5
734-2815	Cell culture dish, increased cell attachment surface, with gripping ring	60	18	21.2
734-2816	Cell culture dish, increased cell attachment surface, with gripping ring	90	17	55.0
734-2817	Cell culture dish, increased cell attachment surface, with gripping ring	100	22	60.8
734-2818	Cell culture dish, increased cell attachment surface	150	22	143.0

TC-treated surface

Catalogue Number	Type	Diameter [mL]	Height [mm]	Growth area [cm^2]
734-2317	Cell culture dish, TC-treated	35	12.3	8.5
734-2318	Cell culture dish, TC-treated, with gripping ring	60	18	21.2
734-2319	Cell culture dish, TC-treated, with gripping ring	70	15	36.3
734-2320	Cell culture dish, TC-treated, with gripping ring	90	17	55.0
734-2321	Cell culture dish, TC-treated, with gripping ring	100	22	60.8
734-2322	Cell culture dish, TC-treated	150	22	143.0
734-2754	Cell culture dish, TC-treated	150	22	143.0

Non treated surface

Catalogue Number	Type	Diameter [mL]	Height [mm]	Growth area [cm^2]
734-2793	Cell culture dish, non treated	35	12,3	8,5
734-2794	Cell culture dish, non treated, with gripping ring	60	18	21,2
734-2795	Cell culture dish, non treated, with gripping ring	90	17	55,0
734-2796	Cell culture dish, non treated, with gripping ring	100	22	60,8
734-2797	Cell culture dish, non treated	150	22	143,0

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

Material	Polystyrene (PS).
Sterilization	Products are EB (Electron beam) irradiated as per ISO 11137 and meets SAL 10^{-6} 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).
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). Test passed.

Latex Statement	This product is not made from natural rubber latex.
BPA Statement	Bisphenols are not used in the manufacture of the raw material and are not expected to be present.
DEHP Statement	Not Applicable.
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.
Shelf Life	3 years.

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 manufacture 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 CoQ using new layout	09-2021
2	2.0	Modified Quality system compliance and disclaimer	07-2025