

Certificate of Conformity

VWR Catalogue Number	See list
Description	VWR® Cell Culture Flasks, Non-Treated or Treated
Country of Origin	Manufactured in China
Date of Issue (yyyy-mm-dd)	2026-06-30

Catalogue Number	Description
734-3692	600 ml, 175cm ² , TC-treated, w/ Plug Seal Cap, Sterilized(5/pkg,50/cs)
734-3696	600 ml, 175cm ² , Non-treated, w/Plug Seal Cap, Sterilized(5/pkg,50/cs)
734-3693	600 ml, 175cm ² , TC-treated, w/Vent Cap, Sterilized, (5/pkg,50/cs)
734-3694	600 ml, 175cm ² , TC-treated, w/Plug Seal Cap, Sterilized, (5/pkg/triple bags,50/cs)
734-3695	600 ml, 175cm ² , TC-treated, w/Vent Cap, Sterilized, (5/pkg,50/cs)
734-3697	600 ml, 175cm ² , Non-treated,w/Vent Cap, Sterilized (5/pkg/triple bags,50/cs)

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

Material	Flask - Polystyrene (PS), Compliance with USP Class VI. Cap - High Density Polyethylene (HDPE), Compliance with USP Class VI.
Sterilization	Product labelled as sterile is 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) 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) Test passed.
Latex Statement	The product is Latex free.
BPA Statement	Bisphenol-A (BPA) and Bisphenol-F (BPF) are not used in the manufacture or the formulation of the raw materials and are not expected to be present. However, tests have not been performed for these chemical materials.
DEHP Statement	Please refer RoHS.
RoHS	Raw materials comply with the requirements of the Directive 2011/65/EU, as amended, and (EU) 2015/863 concerning the limits of cadmium, lead, mercury, hexavalent chromium, dibutyl phthalate(DBP), Diisobutyl phthalates(DIBP), Butyl benzyl phthalate(BBP), Bis(2-ethylhexyl)phthalate(DEHP), Polybrominated Biphenyls (PBB) and Polybrominated Diphenyl Ethers (PBDE).
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 CoC using new layout	12-2025
2	1.0	More products added	03-2026
3	1.0	Table information corrected	06-2026