

Certificate of Quality

VWR Catalogue Number	See List	
Description	VWR® Tissue Culture Plate Inserts, Polycarbonate (PC) Membrane, Sterilized, Standard Line	
Country of Origin	Manufactured in China	
Date of Issue (yyyy-mm-dd)	2022-01-12	

VWR Catalogue Number	Number of wells	Cell growth area	Insert Size	Membrane	Pore size
10769-176	6	4.67 cm ²	24 mm	Translucent	0.1 µm
10769-178	6	4.67 cm ²	24 mm	Translucent	1.0 µm
10769-180	6	4.67 cm ²	24 mm	Transparent	12.0 µm
10769-186	6	4.67 cm ²	24 mm	Translucent	0.4 μm
10769-188	6	4.67 cm ²	24 mm	Transparent	3.0 µm
10769-190	6	4.67 cm ²	24 mm	Transparent	8.0 µm
10769-198	12	1.12 cm²	12 mm	Translucent	0.1 µm
10769-200	12	1.12 cm²	12 mm	Translucent	1.0 µm
10769-202	12	1.12 cm²	12 mm	Transparent	12.0 µm
10769-210	12	1.12 cm²	12 mm	Transparent	3.0 µm
10769-212	12	1.12 cm²	12 mm	Transparent	8.0 µm
10769-208	12	1.12 cm²	12 mm	Translucent	0.4 μm
10769-220	24	0.33 cm ²	6.5 mm	Translucent	0.1 µm
10769-222	24	0.33 cm ²	6.5 mm	Translucent	1.0 µm
10769-224	24	0.33 cm ²	6.5 mm	Transparent	12.0 µm
10769-230	24	0.33 cm ²	6.5 mm	Translucent	0.4 μm
10769-232	24	0.33 cm ²	6.5 mm	Transparent	3.0 µm
10769-234	24	0.33 cm ²	6.5 mm	Transparent	8.0 µm
10769-236	24	0.33 cm ²	6.5 mm	Transparent	5.0 μm



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 product specifications.

Membrane Thickness 0.1 mm

Product Specifications

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

Plate Material Polystyrene (PS),

> Insert Polystyrene (PS), Insert Membrane Polycarbonate (PC)

Sterilization Product is EB (Electron beam) irradiated, and dose released upon

> ISO11137 recommended practices in effect at the time of validation. Products 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).

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 The product is Latex free.

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), 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 Store at room temperature.

Shelf-Life 3 years.

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.