Certificate of Quality

VWR Catalogue Number	See List
Description	VWR® Crystallizing Dishes
Country of Origin	Manufactured in Czech Republic
Date of Issue (yyyy-mm-dd)	2026-01-29

VWR Catalogue Number	Capacity	Ø	Height	Unit
10754-764	50 ml	50 mm	35 mm	Pack of 6
				Case of 24
10754-766	80 ml	60 mm	35 mm	Pack of 6
				Case of 24
10754-768	160 ml	70 mm	50 mm	Pack of 6
				Case of 24
10754-770	170 ml	80 mm	40 mm	Pack of 6
				Case of 24
10754-772	270 ml	90 mm	50 mm	Pack of 6
				Case of 18
10754-774	340 ml	100 mm	50 mm	Pack of 6
				Case of 18
10754-776	700 ml	125 mm	65 mm	Pack of 4
				Case of 12
10754-778	1200 ml	150 mm	75 mm	Pack of 4
				Case of 8
10754-780	1800 ml	170 mm	90 mm	Pack of 4
				Case of 8
10754-782	2600 ml	190 mm	100 mm	Pack of 3
				Case of 6

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 and ISO 28000 standard.

QC Testing

Representative products samples are collected and inspected in accordance with current applicable product specifications.

Test 1: Hydrolytic resistance of glass grains at 98°C in according to ISO 719:1985

Test reference:	ISO 719:2023
Acceptance Criteria:	HGB1: < 0.10 ml/g glass
Test Result:	0.02 ml/g glass
Compliance:	YES

Test 2: Volume and Dimensional verification

Test reference:	Technical drawings
Acceptance Criteria:	Technical drawings
Test Result:	OK
Compliance:	YES

The product is in conformance with the requirements of the following standards and regulations:

- **Glass characteristics:** ISO 3585 Borosilicate glass 3.3 –Properties.
- **Technical standards for products:** ISO 4796 Laboratory glassware. Bottles.
- **Glass containers for pharmaceutical use:** European Pharmacopoeia 10th, Art. 3.2.1. US Pharmacopoeia - Glass Type I

Product Specifications

Material	SIMAX® Borosilicate glass, in fired-on, chemically resistant ceramic enamel (print). The FDA classifies glass in general as GRAS (Generally Recognized as Safe).
Sterilization	The products are non-sterile.
ATP Assay	The products do not contain ATP.
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	No substances (Lead, Cadmium, Mercury, Hexavalent Chromium (Cr^{6+}), 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 routine analysis is performed.
REACH Statement	No substances as listed on >Substance of Very High Concern< candidate list are used in manufacturing of the raw materials used to produce this product. No routine analysis is performed.
Storage Conditions	Store at room temperature in normal warehouse conditions; Protect from any possible contamination.
Shelf-Life	Indefinite.

Version History

Sl.no	Version number	Description	Month & Year
1	1.0	Created General CoQ using new layout	January 2026