Certificate of Quality

VWR Catalogue Number	See List
Description	VWR® Disposable Serological Pipets, Premium Line
Country of Origin	Manufactured in China
Date of Issue (yyyy-mm-dd)	2025-09-02

Individually Wrapped in Paper/Plastic

VWR Catalogue Number	Capacity [mL]	Graduations [mL]	Sterility	Colour Code
75816-102	1	0.01	Sterile	Yellow
75816-104	2	0.01	Sterile	Green
75816-094	5	0.1	Sterile	Blue
76201-710	5	0.1	Sterile	Blue
75816-100	10	0.1	Sterile	Orange
76204-736	10	0.2	Sterile	Orange
75816-090	25	0.2	Sterile	Red
76204-738	25	0.5	Sterile	Red
75816-088	50	0.5	Sterile	Purple
75816-084	100	1.0	Sterile	Pink
76184-872	1	0.01	Sterile	Yellow
76184-742	2	0.01	Sterile	Green
76184-744	10	0.1	Sterile	Blue
76184-746	10	0.1	Sterile	Orange
76184-748	25	0.2	Sterile	Red
76184-750	50	0.5	Sterile	Purple
76184-752	100	1.0	Sterile	Pink

Packed in Polyethylene Bag with Zip Lock

VWR Catalogue Number	Capacity [mL]	Graduations [mL]	Sterility	Colour Code
75816-112	1	0.01	Sterile	Yellow
75816-114	2	0.01	Sterile	Green
75816-106	5	0.01	Sterile	Blue
75816-098	10	0.01	Sterile	Orange
76184-754	10	0.1	Sterile	Orange
75816-092	25	0.2	Sterile	Red
76184-756	50	0.5	Sterile	Purple
75816-086	100	1.0	Sterile	Pink

Bulk Packed in Polyethylene Bags

VWR Catalogue Number	Capacity [mL]	Graduations [mL]	Sterility	Colour Code
75816-116	1	0.01	Non-Sterile	Yellow
75816-110	2	0.01	Non-Sterile	Green
75816-108	5	0.1	Non-Sterile	Blue
75816-096	10	0.1	Non-Sterile	Orange

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 the ISO 9001:2015 & ISO 13485:2016 standard.

QC Testing

Representative products samples are collected and inspected in accordance with current applicable QC requirements.

Test/Procedure	Results
Visual Attributes	Pass
Volumetric	Pass
Ink fastness	Pass

Volumetric Accuracy: Volumetric Accuracy is calibrated for accurate dispensing to within ±2% of full range.

Product Specifications

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

Material	**Pipet** GPPS (General purpose Polystyrene, Compliance with USP Class VI; Colour: Clear) **Mouthpiece Plug** Polyolefin **Food contact statement:** The product meets the requirements of the 21 CFR 177.1640 **Haemolytic statement:** The product has no effect on haemolytic properties under the test condition HR less than 5%.
Sterilization	Product labelled as sterile are EB (Electron Beam) irradiated and dose released upon ISO 11137 recommended practices in effect at the time of validation and they meet a minimum requirement of 10^{-6} SAL with a specified dose range of 15 - 30 kGy.
ATP Assay	Not applicable.
DNase & RNase Free	The products are 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 the product is 0.06 EU/mL. (TAL Gel Clot Method).
Latex Statement	The products are latex free.
BPA Statement	Bisphenol-A (BPA) and Bisphenol-F (BPF) are not used in the manufacture of the raw material and are not expected to be present.
DEHP Statement	Diethylhexyl phthalate (DEHP) is not included in the manufacture or the formulation of the raw material and are not expected to be present.
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 routinely analyse is performed.

REACH Statement	Not applicable.
Storage Conditions	Store at normal temperature.
Shelf Life	Products labelled as sterile has shelf life period of 3 years which is not applicable for non-sterile products.

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 manufacturer and does not engage in manufacturing, co- manufacturing, or processing activities such as design, packaging, labelling, 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	March 2021
2	2.0	Updated Quality System Compliance and Disclaimer statement.	September 2025