Certificate of Quality

VWR Catalogue Number	**See List**
Description	**VWR® Spec-Wipe® 4 Wiper**
Country of Origin	**Manufactured in China**
Date of Issue (yyyy-mm-dd)	**2026-05-26**

VWR Catalogue number	Sterility	Width x Length	Unit
21912-046	Non-sterile	22.9×22.9 cm (9×9")	Pack of 150
			Case of 1200
76377-480	Sterile	30.5×30.5 cm (12×12")	Case of 300

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** standard.

QC Testing

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

Product Specifications

Material 100% polyester.

Sterilization The product labelled as sterile has been dosimetrically released as sterile. All aspects of the manufacturing process are within specifications and the specified minimum dose has been delivered to the product. This is in accordance with procedures outlined in ANSI/AAMI/ISO 11137-2 Method VDmax: Sterilization of healthcare products – Requirements for validation and routine control – Radiation Sterilization, to a Sterility Assurance Level of 10^{-6}.

Min Specified Dose: 25.0 kGy Max Specified Dose: 50.0 kGy

ATP Assay No testing performed.

DNase & RNase Free No testing performed.

BSE/TSE No testing performed.

Cytotoxicity	No testing performed.
Non-Pyrogenic Statement	No testing performed.
Latex Statement	No testing performed.
BPA Statement	No testing performed.
DEHP Statement	The products are free of DEHP.
RoHS	Not Applicable.
REACH Statement	Not Applicable.
Storage Conditions	Store at standard warehouse conditions within an environment of ambient temperature and humidity.
Shelf Life	There is no definite shelf life.

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	May 2026