

Certificate of Conformity

VWR Catalogue Number	See List
Description	VWR® Spec-Wipe® 7, Cleanroom Wipes
Country of Origin	Manufactured in China
Date of Issue (yyyy-mm-dd)	2026-02-23

Catalogue Number	L X W [mm]	Packed
115-0043	229 × 229	100 wipes/bag
115-0044	305 × 305	100 wipes/bag

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**.

QC Testing

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

Product Specifications

Material	100% polyester.
Sterilization	Non-Sterile.
ATP Assay	Not Applicable.
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	Latex or latex derivatives are not used in the production of this product.
BPA Statement	Not Applicable.
DEHP Statement	Not Applicable.
RoHS	Not Applicable.
REACH Statement	Not Applicable.
Storage Conditions	Stored in standard warehouse conditions within an environment of ambient temperature and humidity.
Shelf Life	No 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, 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.

Annex

Attribute; (units)	Value **	Test Method
Basis weight; (g/m ²)	136	
Absorbency in water		
Intrinsic; (mL/g)	2.3	IENT-RP-CC004.2, Sec. 7.1
Extrinsic; (mL/m ²)	312	IENT-RP-CC004.2, Sec. 7.1
Sorptive rate; (seconds)	<1.5	
Extractables		IENT-RP-CC004.2, Sec. 6.1.2
In deionized water; (g/m ²)	0.015	
In isopropyl alcohol; (g/m ²)	0.015	
Specific ions		IENT-RP-CC004.2, Sec. 6.2.2
Sodium; (ppm)	0.11	
Chloride (ppm)	0.13	
Particle generation		IENT-RP-CC004.2, Sec. 5.1
P ₀ ≥ 0.5µm; (x10 ⁶ /m ²)	4.8	
Fibers > 100µm; (x10 ³ /m ²)	0.6	IENT-RP-CC004.2, Sec. 5.2

** ND = None detected; levels are below detection limit of test equipment

Version History

Sl.no	Version number	Description	Month & Year
1	1.0	Created General CoQ using new layout	02-2026