

CERTIFICATE OF ANALYSIS

Thermo Fisher Scientific's Quality System has been found to conform to Quality Management System Standard ISO9001:2015 by TUV SUD America Inc.
 Certificate Number. 951001163

Catalogue Number	022928
Lot Number	T17M705
Description	Acetone, HPLC Grade, 99.5+%
CAS Number	67-64-1
Quality Test/Release Date	04/Jun/2026
Suggested retest date	04/Jun/2031
Country of Origin	United States
Declaration of Origin	Synthetic
BSE/TSE	Not directly derived from or manufactured with any animal byproducts in any way (including but not limited to fermentation or nutrient broth, catalysts, enzymes).

Result Name	Units	Specifications	Test Value
Color (APHA)		≤10 (U.S. specification)	<5
Assay (GC)	%	≥99.5%	99.96
Appearance (Color)		Clear colorless	Clear colorless
Form		Liquid	Liquid
Impurity content	ppb	Fluorescence at Emission maximum for impurities ≤1ppb (as Quinine sulfate) (U.S. specification)	<1
Impurity content	ppt	Heptachlor epoxide: ≤10 ppt by GC-ECD (U.S. specification)	4
Water Content (Karl Fischer Titration)	%	≤0.50%	0.27
Optical Absorbance at	AU	330nm: ≤1.00AU (U.S. specification)	0.71
Optical Absorbance at	AU	340nm: ≤0.10AU (U.S. specification)	0.05
Optical Absorbance at	AU	350nm: ≤0.01AU (U.S. specification)	0.001
Optical Absorbance at	AU	375nm: ≤0.005AU (U.S. specification)	<0.001
Optical Absorbance at	AU	400nm: ≤0.005AU (U.S. specification)	<0.001
Residue on Evaporation	ppm	≤1 ppm	<0.1



Derek Roy
 Quality Manager, LCD

Note: The data listed is valid for all package sizes of this lot of this product, expressed as an extension of the catalogue number listed above.



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Products are processed under ISO 9001:2015 quality management systems and samples are tested for conformance to the noted specifications. Certain data may have been supplied by third parties. We disclaim the implied warranties of merchantability and fitness for a particular purpose, and the accuracy of third-party data or information associated with the product. Products are for research and development use only. Products are not for direct administration to humans or animals. It is the responsibility of the final formulator or end user to determine suitability, and to qualify and/or validate each product for its intended use.

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