

## CERTIFICATE OF ANALYSIS

Thermo Fisher Scientific's Quality System has been found to conform to Quality Management System Standard ISO9001:2015 by Intertek Global  
Certificate Number: CERT-0120633

Catalogue Number	C185
Lot Number	261019
Description	CARBON DISULFIDE, LOW BENZENE MEETS A.C.S. SPECIFICATION
CAS Number	75-15-0
Quality Test/Release Date	16/Jun/2026
Suggested retest date	15/Jun/2031
Country of Origin	United States
Declaration of Origin	Organic - non animal
BSE/TSE	No animal products are used as starting raw material ingredients, or used in processing, including lubricants, processing aids, or any other material that might migrate to the finished product.

Result Name	Units	Specifications	Test Value
APPEARANCE		REPORT	Clear, colorless liquid free of suspended matter
ASSAY	%	$\geq 99.9$	99.99
COLOR	APHA	$\leq 10$	<5
EVAPORATION RESIDUE	%	$\leq 0.002$	0.00002
HYDROGEN SULFIDE IDENTIFICATION	PASS/FAIL	= P.T. (ABOUT 1.5 PPM)	<1.5
	PASS/FAIL	= PASS TEST	PASS TEST
OPTICAL ABS AT 382 NM	ABSORBANCE UNITS	$\leq 1.0$	0.9
OPTICAL ABS AT 388 NM	ABSORBANCE UNITS	$\leq 0.50$	0.39
OPTICAL ABS AT 403 NM	ABSORBANCE UNITS	$\leq 0.10$	0.05
OPTICAL ABS AT 410 NM	ABSORBANCE UNITS	$\leq 0.05$	0.02
OPTICAL ABS AT 500 NM	ABSORBANCE UNITS	$\leq 0.01$	0.002
SULFUR DIOXIDE	PASS/FAIL	= P.T. (ABOUT 2.5 PPM)	PASS TEST
TRACE ORGANIC IMPURITY	ppm	$\leq 1$	<1
WATER (H <sub>2</sub> O)	%	$\leq 0.05$	0.002



Ash Frey  
Supervisor, Quality Control

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.

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 use or further manufacturing. 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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