

## CERTIFICATE OF ANALYSIS

Thermo Fisher Scientific's Quality System has been found to conform to Quality Management System Standard ISO9001:2015 by Intertek. Certificate Number. 2317548

Catalogue Number	13266
Lot Number	A0481787
Description	L-Serine,99%
CAS Number	56-45-1
Quality Test/Release Date	01/Apr/2026
Suggested retest date	01/Apr/2031
Country of Origin	CHINA
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
Appearance (Color)		White	White
Appearance (Form)		Crystalline powder or crystals or needles	Crystalline powder
Infrared spectrum		Conforms	Conforms
Titration with HClO <sub>4</sub>	%	98.5 to 101.5	99.6
Other amino acids	%	=<0.5 (TLC)	=<0.5 (TLC)
Loss on drying	%	=<0.2 (105°C, 3 hrs)	0.06 (105°C, 3 hrs)
Heavy metals (as Pb)	ppm	=<10	=<10
Sulfated ash	%	=<0.1	=<0.001
Specific optical rotation		+14.4° to +16.0° (20°C, 589 nm) (c=10, 2 N HCl)	+14.8° (20°C, 589 nm) (c=10, 2 N HCl)
Chloride (Cl)	ppm	=<200	=<200
Sulfate (SO <sub>4</sub> )	ppm	=<200	=<200
Ammonium (NH <sub>4</sub> )	ppm	=<200	=<200
Arsenic (As)	ppm	=<1	=<1
Iron (Fe)	ppm	=<10	=<10
UV transmittance	%	>=95 (c=10, H <sub>2</sub> O, 430 nm) 1 cm cell	99.3 (c=10, H <sub>2</sub> O, 430 nm) 1 cm cell
pH		5.0 to 6.5 (2.5 % at 25°C)	5.9 (2.5 % at 25°C)

Geert Torfs  
Supervisor, QC

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

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.

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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