Fisher Scientific UK Ltd.
Bishop Meadow Road
Loughborough,
LE11 5RG, United Kingdom

CERTIFICATE OF ANALYSIS

Thermo Fisher Scientific's Quality System has been found to conform to Quality Management System Standard ISO9001:2015 by INTERTEK SAI Global Certificate Number. QMS42099

Catalogue Number	F/1501
Lot Number	2195359
Description	Formaldehyde solution SLR Specified reagent for general lab work
Quality Test/Release Date	13/Sep/2021
Expiry Phrase	Use within 2 yrs of opening
Country of Origin	United Kingdom
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
Acidity/alkalinity (meq/g)	meq/g	<= 0.05	0.008
Assay	%	>= 37 and <= 41	37.2
Calcium (Ca)	ppm	<= 50	<5
Chemical form		Mobile liquid	Mobile liquid
Colour	APHA	<= 20	<20
Copper (Cu)	ppm	<= 2	<0.5
Iron (Fe)	ppm	<= 5	<0.5
Lead (Pb)	ppm	<= 2	<0.5
Magnesium (Mg)	ppm	<= 20	<0.5
Methanol	%	>= 9 and <= 14	10
Potassium (K)	ppm	<= 5	<0.5
Sulfated ash	%	<= 0.1	0.004
Total chloride (Cl)	%	<= 0.005	<0.001
Total silicon (Si)	ppm	<= 10	<0.5
Total sulfur (S)	ppm	<= 100	<5
Visual colour		Clear colourless	Clear colourless
Wt/ml at 20C	g	>= 1.075 and <= 1.09	1.082
Zinc (Zn)	ppm	<= 10	<0.5

Additional Information Please note: The assay stated on the COA is by w/v.

A. Ganatra

Ashok Ganatra
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.

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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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