

BIOSPECTRA

100 Majestic Way, Bangor, PA 18013 / www.biospectra.us

Effective Date:	23-Mar-2021	23-Mar-2024	: Date of Next Review
Prepared By:	Jared L Lobb	19-002973 v.8.0	: Supersedes
QA/QC Approval:	Carissa McCollan	Wendy Santay	: Management Approval
Reason for Revision:	See Revision History in ensur.		

CERTIFICATE OF ANALYSIS

TRIS

BIO EXCIPIENT GRADE / NEW CODE TRIS-3255-05

(HISTORICAL CODE TR3255-K005)

LOT: TRIS-0122-00047

NH₂C(CH₂OH)₃ ▲ F.W. 121.14 g/mol. ▲ CAS# 77-86-1
 Manufacturing Date: 11/23/21 Expiration Date: 11/30/24
 Manufacturing Site: 1474 Rockdale Lane, Stroudsburg, PA 18360
 Packaging Date: 2/20/22 Packaging Site: 100 Majestic Way, Bangor PA, 18013
 Meets or Exceeds USP, EP and JPC Specifications

USP COMPENDIA

ANALYSIS	SPECIFICATION	TEST RESULT
Assay (Dried Basis)	99.0-101.0%	99.7%
Identification A	Passes Test	Passes Test
Identification B	Passes Test	Passes Test
Identification C	Passes Test	Passes Test
Loss on Drying	1.0% max.	0.2%
Melting Range	168-172°C	170 - 172 °C
pH (1 in 20)	10.0 – 11.5	10.7
Residue on Ignition	0.1% max.	<0.1%

EP COMPENDIA

ANALYSIS	SPECIFICATION	TEST RESULT
Appearance of Solution	Passes Test	Passes Test
Assay (Dried Basis)	99.0-100.5%	99.7%
Chloride (Cl)	≤ 100 ppm	<100 ppm
Identification A	Passes Test	Passes Test
Identification B (Melting Range)	168-172°C	170 - 172 °C
Identification C	Passes Test	Passes Test
Identification D	Passes Test	Passes Test
Iron (Fe)	10 ppm max.	<10 ppm
Loss on Drying @105°C	0.5% max.	0.2%
pH (5%)	10.0-11.5	10.7
Related Substances	≤ 1.0%	<1.0%

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ANALYSIS	SPECIFICATION	TEST RESULT
Sulfated Ash	0.1% max.	<0.1%

JPC ANALYSIS

ANALYSIS	SPECIFICATION	TEST RESULT
Arsenic (As)	1.6 ppm max.	≤ 1.6 ppm
Assay (Dried Basis)	99.0-101.0%	99.7%
Clarity and Color of Solution	Passes Test	Passes Test
Heavy Metals	8 ppm max.	≤ 8 ppm
Identification A	Passes Test	Passes Test
Identification B	Passes Test	Passes Test
Loss on Drying	0.5% max.	0.2%
Melting Point	168-172°C	170 - 172 °C
pH	10.3 – 10.7	10.5
Residue on Ignition	0.1% max.	<0.1%

ADDITIONAL ANALYSES

ANALYSIS	SPECIFICATION	TEST RESULT
Appearance and Color	White, crystalline powder to needle-like crystals	White, crystalline powder to needle-like crystals
Absorbance (1M)	0.06 a.u. max	0.01 a.u.
	0.01 a.u. max	<0.01 a.u.
Absorbance (10%)	0.03 a.u. max.	0.01 a.u.
	0.02 a.u. max.	0.01 a.u.
Absorbance (40%)	0.2 a.u. max.	<0.2 a.u.
APHA Color, 20% Solution	20 APHA max.	<20 APHA
Assay (Ultrapure, Dried Basis)	99.9% min	99.9%
Endotoxins	≤ 2.5 EU/g	1.7 EU/g
Enzymes	None	None
	None	None
	None	None
Heavy Metals (As Pb)	1 ppm max.	≤ 1 ppm
Insoluble Matter	0.005% max.	<0.005%
Karl Fischer Water	1.0% max.	0.1%
Loss on Drying	0.3% max.	0.2%
Microbial Content	≤ 100 CFU/g	<100 CFU/g
	≤ 100 CFU/g	<100 CFU/g
Related Substances	0.1% max.	<0.1%

ANALYSIS	SPECIFICATION	TEST RESULT
Residue on Ignition	0.05% max.	0.04%
Arsenic (As)	≤ 1.6 ppm	≤ 1.6 ppm
Calcium (Ca)	≤ 1 ppm	≤ 1 ppm
Copper (Cu)	≤ 1 ppm	≤ 1 ppm
Trace Metals		
Iron (Fe)	≤ 1 ppm	≤ 1 ppm
Lead (Pb)	≤ 1 ppm	≤ 1 ppm
Magnesium (Mg)	≤ 5 ppm	≤ 5 ppm
Manganese (Mn)	≤ 1 ppm	≤ 1 ppm
Zinc (Zn)	≤ 1 ppm	≤ 1 ppm

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-000496

INTENDED USE: Material represented by this Certificate of Analysis is suitable for use as an excipient. It is manufactured in accordance with the ICH Q7 Good Manufacturing Practice Guide. The material represented by this Certificate of Analysis is not suitable to be used as an Active Pharmaceutical Ingredient, Drug Product or Household Item.

RESIDUAL SOLVENTS: Based on the manufacturing process and the controlled handling, storage and analysis of this product, this product complies with the requirements and specifications listed in the current USP method <467> Tables 1, 2, 3, or 4.

Prepared by: John Hughes Date: 3/1/22 Job Title: QA Specialist

Reviewed by: Car Date: 3/1/22 Job Title: QA Manager