

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 McCollian	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-0123-00267

NH₂C(CH₂OH)₃ * F.W. 121.14 g/mol. * CAS# 77-86-1

Manufacturing Date: 11/03/23 Expiration Date: 11/30/26

Manufacturing Site: 1474 Rockdale Lane, Stroudsburg, PA 18360

Packaging Date: 11/18/23 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%	100.3%
Identification A	Passes Test	Passes Test
Identification B	Passes Test	Passes Test
Identification C	Passes Test	Passes Test
Loss on Drying	1.0% max.	0.1%
Melting Range	168 - 172 °C	171 - 172 °C
pH (1 in 20)	10.0 - 11.5	10.9
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%	100.3%
Chloride (Cl)	≤ 100 ppm	< 100 ppm
Identification A	Passes Test	Passes Test
Identification B (Melting Range)	168 - 172 °C	171 - 172 °C
Identification C	Passes Test	Passes Test
Identification D	Passes Test	Passes Test
Iron (Fe)	10 ppm max.	< 0.30 ppm
Loss on Drying @ 105°C	0.5% max.	0.1%
pH (5%)	10.0 - 11.5	10.9
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%	100.3%
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.	0.1%
Melting Point	168 - 172 °C	171 - 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
	260nm	0.06 a.u. max
Absorbance (1M)	280nm	0.01 a.u.
	430nm	0.01 a.u.
	260nm	0.03 a.u. max.
Absorbance (10%)	280nm	0.01 a.u.
	430nm	< 0.003 a.u.
Absorbance (40%)	290nm	< 0.2 a.u.
APHA Color, 20% Solution	20 APHA max.	< 20 APHA
Assay (Ultrapure, Dried Basis)	99.9% min	100.2%
Endotoxins	≤ 2.5 EU/g	< 1.0 EU/g
	DNase	None
Enzymes	Protease	None
	RNase	None
Heavy Metals (As Pb)	1 ppm max.	≤ 1 ppm
Insoluble Matter	0.005% max.	< 0.001%
Karl Fischer Water	1.0% max.	0.1%
Loss on Drying	0.3% max.	0.1%
Microbial Content	TAMC	≤ 100 CFU/g
	TYMC	≤ 100 CFU/g
Related Substances	0.1% max.	< 0.1%

ANALYSIS	SPECIFICATION	TEST RESULT
Residue on Ignition	0.05% max.	< 0.01%
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
	Lead (Pb)	≤ 1 ppm
	Magnesium (Mg)	≤ 5 ppm
	Manganese (Mn)	≤ 1 ppm
	Zinc (Zn)	≤ 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 **m**anufacturing 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: M. Mhafer Date: 12/06/23 Job Title: QA Mater. Disp. Tech. III

Reviewed by: John Singh Date: 12/6/23 Job Title: QA Mater. Disp. Supervisor

