DCN: 16-001164 v.3.0

## **BI**SPECTRA

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

Effective Date:	28-Mar-2018	28-Mar-2021	: Date of Next Review
Prepared By:	Jamie Storm	16-001164 v.2.0	: Supersedes
QA/QC Approval:	Crystal Hamelburg	Dora Meissner	: Management Approval
Reason for Revision:	See Revision History in ensur.		

## **TRIS**

## **CERTIFICATE OF ANALYSIS**

## BIO EXCIPIENT GRADE / TR3220-G100

LOT: TR3220-010-0219

NH<sub>2</sub>C(CH<sub>2</sub>OH)<sub>3</sub> ^ F.W. 121.14 g/mol. ^ CAS# 77-86-1 Manufacture Date: 3/20/2018 Retest Date: 3/31/2020 Manufacturing Site: 1474 Rockdale Lane, Stroudsburg, PA 18360

Packaging Date: 2/27/2019

Packaging Site: 100 Majestic Way, Bangor PA, 18013

Meets or Exceeds USP Specifications

Analysis		SPECIFICATION	TEST RESULT	
Absorbance (40%)	290nm	0.2 a.u. max.	0. 0751 a.u.	
Appearance and Color		White / Crystals	White / Crystals	
Assay		99.0 - 101.0%	100.31%	
	RNase	None Detected	None Detected	
Enzymes	DNase	None Detected	None Detected	
	Protease	None Detected	None Detected	
Heavy Metals		5 ppm max.	< 5 ppm	
Identification B		Passes Test	Passes Test	
Identification C		Passes Test	Passes Test	
Identification (IR)		Passes Test	Passes Test	
Insoluble Matter		0.005% max	<0.0010%	
Karl Fischer Water		2.0% max.	0.32%	
Loss on Drying	Loss on Drying		0.1543%	
Melting Range		168-172°C	170.3 − 171.4 °C	
pH (5%)		10.0 - 11.5	10.892 @ 21.80 °C	
Residue On Ignition		0.1% max.	<0.0200%	
	Arsenic (As)	5 ppm max.	< 5 ppm	
	Calcium (Ca)	5 ppm max.	< 5 ppm	
Torres Madel	Copper (Cu)	5 ppm max.	< 5 ppm	
Trace Metals	Iron (Fe)	5 ppm max.	< 5 ppm	
	Lead (Pb)	5 ppm max.	< 5 ppm	
N	Magnesium (Mg)	5 ppm max.	< 5 ppm	

DCN: 16-001164 v.3.0

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-000496

INTENDED USE: Material represented by this Certificate of Analysis is suitable to be used only as the following: ICH Q7 Compliant cGMP Manufactured Excipient for use in further Manufacturing. 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 STATEMENT: Based on the manufacturing process and the controlled handling and storage of this product, there is no potential or any of the residual solvents listed in the current USP method <467> Tables 1, 2, 3, or 4 to be present at the specified limits; furthermore, if tested this product would comply with USP/NF requirements.

Prepared by:	Date: _	2128/19
Reviewed by: # Berne	Date: _	2128/19