DCN: 19-002851 v.2.0



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

ECC .: D .	7 4 2020	7 4 2022	D . CN . D .
Effective Date:	/-Aug-2020	/-Aug-2023	: Date of Next Review
Prepared By:	Amy Hosein	19-002851 v.1.0	: 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 / TR3254-K001

LOT: TR3254-008-0820

 $NH_2C(CH_2OH)_3 \land F.W. 121.14 g/mol. \land CAS# 77-86-1$

Manufacture Date: 7/14/20

Retest Date: 7/31/22

Manufacturing Site: 1474 Rockdale Lane, Stroudsburg, PA 18360

Packaging Date: 8/28/20 Packaging Site: 100 Majestic Way, Bangor PA, 18013

USP COMPENDIA				
ANALYSIS		SPECIFICATION	TEST RESULT	
Appearance and Color		White/Crystals	White/Crystals	
Assay		99.0 - 101.0%	99.7%	
Endotoxin		\leq 2.5 EU/g	<1.0EU/g	
Identification A		Passes Test	Passes Test	
Identification B		Passes Test	Passes Test	
Identification C		Passes Test	Passes Test	
Loss on Drying		≤ 1.0%	0.1%	
Melting Range		168-172°C	170-172°C	
pH (1 in 20)		10.0 - 11.5	10.7 @ 24.8°C	
Residue on Ignition		≤ 0.1%	<0.1%	
MissalialContant	TAMC	$\leq 500 \text{ CFU/g}$	<10CFU/g	
MicrobialContent	TYMC	≤ 200 CFU/g	<10CFU/g	

EP COMPENDIA				
ANALYSIS	SPECIFICATION	TEST RESULT		
Appearance of Solution	Passes Test	Passes Test		
Assay	99.0 - 100.5%	99.7%		
Chloride (Cl)	≤ 100 ppm	<100ppm		
Identification A	Passes Test	Passes Test		
Identification B	168-174°C	170-172°C		
Identification C	Passes Test	Passes Test		
Iron (Fe)	< 10ppm	<10ppm		

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		DC11. 17-002031 1.2.0		
EP COMPENDIA				
Analysis	SPECIFICATION	TEST RESULT		
Loss on Drying at 105°C	≤ 0.5%	0.1%		
pН	10.0 - 11.5	10.7 @ 24.8°C		
Related Substances	≤ 1.0%	<1.0%		
Sulfated Ash	≤ 0.1%	<0.1%		

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-000496

<u>RESIDUAL SOLVENTS STATEMENT</u>: 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.

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.

Prepared by:	_ Date: _	911/20	_ Job Title: _	DA Supervisor
Reviewed by: Muy Jaut	_ Date: _	09/01/20	_ Job Title: _	Of Manager