BIOSPECTRA

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

Effective Date:	1-Aug-2022	1-Aug-2025	: Date of Next Review
Prepared By:	Wendy Santay	BSI-COA-0097 v.8.0	: Supersedes
QA/QC Approval:	Carissa McCollian	Amy Yencho	: Management Approval
Reason for Revision:	See Revision History in MasterControl.		

CERTIFICATE OF ANALYSIS

TREHALOSE, DIHYDRATE

BIO EXCIPIENT GRADE / NEW CODE TRED-3250-10

(HISTORICAL CODE TE3250-K010)

LOT: TRED-0122-00037

$C_{12}H_{22}O_{11} \cdot 2H_2O$ ▲ F.W. 378.33 g/mol. ▲ CAS# 6138-23-4

Manufacturing Date: 6/22/22 Retest Date: 6/30/24

Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 12/5/22 Packaging Site: 100 Majestic Way, Bangor PA, 18013

Meets or Exceeds USP/NF, EP and JP Specifications

NF COMPENDIA

ANALYSIS		SPECIFICATION	TEST RESULT
[1]Assay		[3]98.0% - 101.0%	100.4%
Chloride and Sulfate, *Chloride*		≤ 0.0125%	≤ 0.0125%
Color and Clarity of Solution	A720	≤ 0.050	<0.003
	A420 – A720	≤ 0.100	0.016
[2]Endotoxins		[3]≤ 2.4 EU/g	≤0.2 EU/g
[2]Identification A		Conforms to Standard	Conforms to standard
[2]Identification B		Passes Test	Passes Test
[2]Identification C		Passes Test	Passes Test
[2]Microbial Content	*Escherichia coli*	Absent/g	Absent/g
	Salmonella species	Absent/10g	Absent/10g
	TAMC	≤ 100 CFU/g	10 CFU/g
	TYMC	≤ 100 CFU/g	≤ 10 CFU/g
[2]Nitrogen Determination		≤ 0.005%	≤ 0.001 %
[2]Optical Rotation, Specific Rotation @ 20°C		+197° to +201°	+199°
[2]pH @ 25°C		4.5 – 6.5	5.6
[1]Related Substances	Total Impurities with RRT <1.0	≤ 0.5%	≤ 0.5%
	Total Impurities with RRT >1.0	≤ 0.5%	≤ 0.5%
[2]Residue on Ignition		≤ 0.1%	≤ 0.1%

ANALYSIS	SPECIFICATION	TEST RESULT
[2]Soluble Starch	Passes Test	Passes Test
Chloride and Sulfate, *Sulfate*	≤ 0.0200%	≤ 0.0200%
[2]Water Determination	9.0% to 11.0%	10.2%

EP COMPENDIA

ANALYSIS		SPECIFICATION	TEST RESULT
[1]Assay		[3]98.0 - 101.0%	100.4%
Appearance of Solution		Clear, colorless	Clear, colorless
Chlorides		≤ 0.0125%	≤ 0.0125%
[2]Endotoxins		[3]≤ 2.4 EU/g	<0.2 EU/g
[2]Identification A		Conforms to Standard	Conforms to standard
[2]Identification B		Passes Test	Passes Test
[2]Identification C		Passes Test	Passes Test
[1]Related Substances	Impurity A	≤ 0.5%	≤ 0.5%
	Impurity B	≤ 0.5%	≤ 0.5%
	Unspecified Impurities	≤ 0.2%	≤ 0.2%
	Total Impurities	≤ 1.0%	≤ 1.0%
[2]Microbial Content	*Escherichia coli*	Absent/g	Absent/g
	Salmonella species	Absent/10g	Absent/10g
	TAMC	≤ 100 CFU/g	10CFU/g
	TYMC	≤ 100 CFU/g	<10CFU/g
[2]pH @ 25°C		4.5 - 6.5	5.6
[2]Soluble Starch		Passes Test	Passes Test
[2]Specific Optical Rotation @ 20°C		+197° to +201°	+199°
Sulfated Ash		≤ 0.1%	≤ 0.1%
Sulfates		≤ 0.0200%	≤ 0.0200%
[2]Water		9.0% to 11.0%	10.2%

JP COMPENDIA

ANALYSIS	SPECIFICATION	TEST RESULT
[1]Assay	98.0% - 101.0%	100.4%
Chloride	≤ 0.018%	≤ 0.018%
[2]Dextrin, Soluble Starch, Sulfite	Passes Test	Passes Test
Heavy Metals (as Pb)	≤ 5 ppm	≤ 5 ppm
[2]Identification 1	Passes Test	Passes Test

ANALYSIS		SPECIFICATION	TEST RESULT
[2]Identification 2		Passes Test	Passes Test
[2]Identification 3		Conforms to Standard	Conforms to Standard
[2]Nitrogen		≤ 0.005%	0.001%
[2]Optical Rotation @ 20°C		+197° to +201°	+199°
[2]pH @ 25°C		4.5 – 6.5	5.6
[2]Residue on Ignition		≤ 0.1%	≤ 0.1%
[1]Related Substances	Total Impurities with RRT <1.0	≤ 0.5%	≤ 0.5%
	Total Impurities with RRT >1.0	≤ 0.5%	≤ 0.5%
Sulfate		≤ 0.024%	≤ 0.024%
[2]Water		9.0% to 11.0%	10.2%

NON-COMPENDIAL ANALYSES

ANALYSIS	SPECIFICATION	TEST RESULT
Appearance and Color	White to Almost White Crystalline Powder	White to Almost White Crystalline Powder
[1]Residual Ethanol	≤ 200 ppm	<200 ppm
[1]Residual Isopropyl Alcohol	≤ 250 ppm	<250 ppm
[1]Residual Methanol	≤ 50 ppm	<50 ppm

[1]Alternate Validated Method
[2]Analyses are Harmonized
[3]Specifications is more stringent than Compendia Monograph

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: BSI-ATM-0027

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 STATEMENT: 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. Ethanol and Methanol are not used in the manufacturing process.

Prepared by: [signature] Date: 12/9/22 Job Title: QA Specialist

Reviewed by: [signature] Date: 12/9/22 Job Title: Assoc. Director of Quality