BI@SPECTRA

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-3250-05 (HISTORICAL CODE TE3250-K005) LOT: TRED-0124-00020

C₁₂H₂₂O₁₁ 2H₂O ↓ F.W. 378.33 g/mol. ↓ CAS# 6138-23-4 Manufacturing Date: 08/11/23 Retest Date: 08/31/25 Manufacturing Site: 100 Majestic Way, Bangor PA, 18013 Packaging Date: 08/07/24 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%	100.0%		
Chloride and Sulfate, Chloride		\leq 0.0125%	< 0.0125%		
Color and Clarity	A720	\leq 0.050	< 0.003		
of Solution	A420 - A720	≤ 0.100	0.013		
² Endotoxins		$^{3} \le 2.4 \text{ EU/g}$	< 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		
Escherichia coli		Absent/g	Absent/g		
² Microbial Salmonella species		Absent/10g	Absent/10g		
Content	TAMC	\leq 100 CFU/g	< 100 CFU/g		
	TYMC	$\leq 100 \text{ CFU/g}$	< 100 CFU/g		
² Nitrogen Determinati	on	\leq 0.005%	< 0.005 %		
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	$\leq 0.5\%$	0.11%		
Related Substances	Total Impurities with RRT >1.0	\leq 0.5%	< 0.01%		
² Residue on Ignition		$\leq 0.1\%$	< 0.1%		

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Analysis	Specification	TEST RESULT	
² Soluble Starch	Passes Test	Passes Test	
Chloride and Sulfate, Sulfate	$\leq 0.0200\%$	< 0.0200%	
² Water Determination	9.0% to 11.0%	9.4%	

		EP COMPENDIA		
ANALYSIS		Specification	TEST RESULT	
¹ Assay		³ 98.0 - 101.0%	100.0%	
Appearance of Solution		Clear, colorless	Clear, colorless	
Chlorides		≤ 0.0125%	< 0.0125%	
² Endotoxins		$^{3} \le 2.4 \text{ EU/g}$	< 0.2 EU/g	
² Identification	А	Conforms to Standard	Conforms to standard	
² Identification B		Passes Test	Passes Test	
² Identification C		Passes Test	Passes Test	
	Impurity A	≤ 0.5%	< 0.10%	
¹ Related	Impurity B	$\leq 0.5\%$	< 0.10%	
Substances	Unspecified Impurities	≤ 0.2%	0.11%	
	Total Impurities	≤ 1.0%	0.11%	
	Escherichia coli	Absent/g	Absent/g	
² Microbial	Salmonella species	Absent/10g	Absent/10g	
Content	TAMC	≤ 100 CFU/g	< 100 CFU/g	
	TYMC	\leq 100 CFU/g	< 100 CFU/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%	9.4%	

JP COMPENDIA					
ANALYSIS	Specification	TEST RESULT			
¹ Assay	98.0% - 101.0%	100.0%			
Chloride	$\leq 0.018\%$	< 0.018%			
² Dextrin, Soluble Starch, Sulfite	Passes Test	Passes Test			
Heavy Metals (as Pb)	\leq 5 ppm	< 5 ppm			
² Identification 1	Passes Test	Passes Test			

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			DUN: D51-CUA-0097 V.0.1	
Analysis		SPECIFICATION	TEST RESULT	
² Identification 2		Passes Test	Passes Test	
² Identification 3		Conforms to Standard	Conforms to Standard	
² Nitrogen		≤ 0.005%	< 0.005%	
² Optical Rotation @ 20°C		+197° to +201°	+199°	
² pH @ 25°C		4.5 - 6.5	5.6	
² Residue on Ig	nition	≤ 0.1%	< 0.1%	
¹ Related	Total Impurities with RRT <1.0	≤ 0.5%	0.11%	
Substances	Total Impurities with RRT >1.0	≤ 0.5%	< 0.01%	
Sulfate		$\leq 0.024\%$	< 0.024%	
² Water		9.0% to 11.0%	9.4%	

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	\leq 200 ppm	< 95 ppm			
¹ Residual Isopropyl Alcohol	\leq 250 ppm	< 135 ppm			
¹ Residual Methanol	\leq 50 ppm	< 25 ppm			

¹Alternate Validated Method

²Analyses are Harmonized

³Specifications is more stringent than Compendia Monograph

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: BSI-ATM-0027

<u>INTENDED USE:</u> 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.

<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. Ethanol and Methanol are not used in the manufacturing process.

Prepared by: Anil Mc (all	Date: _	8/13/24	Job Title: _	QA Tech I
Reviewed by: Joon Bugh	Date: _	8/13/24	Job Title: _	QA Supervisor

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