DCN: 18-002600 v.8.0

BI SPECTRA

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

	Effective Date:	1-Mar-2021	1-Mar-2024	: Date of Next Review
	Prepared By:	Jaron Hughes	18-002600 v.7.2	: Supersedes
	QA/QC Approval:	Carissa McCollian	Wendy Santay	: Management Approval
Re	eason for Revision:	See Revision History in ensur.	2	***

CERTIFICATE OF ANALYSIS

TREHALOSE, DIHYDRATE

BIO EXCIPIENT GRADE / NEW CODE TRED-3250-92

(HISTORICAL CODE TE3250-G100)

LOT: TRED-0122-00003

C₁₂H₂₂O₁₁ · 2H₂O F.W. 378.33 g/mol. CAS# 6138-23-4

Manufacturing Date: 08/01/21 Retest Date: 08/31/24

Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 01/10/22 Packaging Site: 100 Majestic Way, Bangor PA, 18013

Meets or Exceeds USP/NF, EP and JP Specifications

	Wicets of Ex	ceeds USP/NF, EP and JP Specifica	ttions	
		NF COMPENDIA		
Anal	YSIS	SPECIFICATION	TEST RESULT	
¹ Assay		³ 98.0% - 101.0%	99.4%	
Chloride and Sulfate,	Chloride	≤ 0.0125%	≤ 0.0125%	
Color and Clarity	A720	≤ 0.050	0.003	
of Solution	A420 - A720	≤ 0.100	0.017	
² Endotoxins		$^3 \le 2.4 \text{ EU/g}$	$\leq 0.2 \; EU/g$	
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	$\leq 100 \text{ CFU/g}$	≤ 10 CFU/g	
	TYMC	≤ 100 CFU/g	$\leq 10 \text{ CFU/g}$	
² Nitrogen Determinat	ion	≤ 0.005%	≤ 0.005 %	
² Optical Rotation, Spe 20°C	ecific Rotation @	+197° to +201°	+199°	
²pH @ 25°C		4.5 - 6.5	5.7	
¹ Related Substances	Total Impurities with RRT < 1.0	≤ 0.5%	≤ 0.5%	
Rotated Substances	Total Impurities with RRT >1.0	≤ 0.5%	≤ 0.5%	
² Residue on Ignition		$\leq 0.1\%$	≤ 0.1%	

DCN: 18-002600 v.8.0

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%	9.3%

		EP COMPENDIA	
Anai	LYSIS	SPECIFICATION	TEST RESULT
¹ Assay		³ 98.0 - 101.0%	99.4%
Appearance of Solut	cion	Clear, colorless	Clear, colorless
Chlorides		≤ 0.0125%	≤ 0.0125%
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
	Impurity A	≤ 0.5%	≤ 0.5%
¹ Related	Impurity B	≤ 0.5%	≤ 0.5%
	nspecified Impurities	≤ 0.2%	$\leq 0.2\%$
	Total Impurities	≤ 1.0%	≤ 1.0%
	Escherichia coli	Absent/g	Absent
² Microbial	Salmonella species	Absent/10g	Absent
Content	TAMC	≤ 100 CFU/g	< 10CFU/g
	TYMC	≤ 100 CFU/g	< 10CFU/g
²pH @ 25°C		4.5 - 6.5	5.7
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%	9.3%

	JP COMPENDIA		
Analysis	SPECIFICATION	TEST RESULT	
¹ Assay	98.0% - 101.0%	99.4%	
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	

DCN: 18-002600 v.8.0

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°
²pH @ 25°C		4.5 - 6.5	5.7
² Residue on Ignition		≤ 0.1%	≤ 0.1%
¹ Related	Total Impurities with RRT < 1.0	≤ 0.5%	≤ 0.5%
Substances	Total Impurities with RRT >1.0	≤ 0.5%	≤ 0.5%
Sulfate		≤ 0.024%	≤ 0.024%
² Water	<u> </u>	9.0% to 11.0%	9.3%

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	≤ 5000 ppm	≤ 5000 ppm					
¹ Residual Methanol	≤ 3000 ppm	≤3000 ppm					

¹Alternate Validated Method

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 18-002375

<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.

Prepared by:	M. Shaf	fire	Date: _	01/12/22	Job Title:	QA Tech.	I

Reviewed by: Jan Augh Date: 1/12/22 Job Title: QA Specialist

²Analyses are Harmonized

³Specifications is more stringent than Compendia Monograph

•