DCN: 18-002600 v.8.0

BI SPECTRA

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

Effective Date: 1-Mar-20	021	1-Mar-2024	: Date of Next Review	
Prepared By: Jaron Hu	ighes	18-002600 v.7.2	: Supersedes	
QA/QC Approval: Carissa N	McCollian	Wendy Santay	: Management Approval	
Reason for Revision: See Revi	sion History in ensur.		A	

CERTIFICATE OF ANALYSIS

TREHALOSE, DIHYDRATE

BIO EXCIPIENT GRADE / NEW CODE TRED-3250-01

(HISTORICAL CODE TE3250-K001)

LOT: TRED-0122-00018

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

Manufacturing Date: 07/22/21

Retest Date: 07/31/24

Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 4/29/22 Packaging Site: 100 Majestic Way, Bangor PA, 18013

Meets or Exceeds USP/NF, EP and JP Specifications

Buryana		NF COMPENDIA	
Anal	YSIS	SPECIFICATION	TEST RESULT
¹ Assay		³ 98.0% - 101.0%	99.6%
Chloride and Sulfate,	Chloride	≤ 0.0125%	≤ 0.0125%
Color and Clarity	A720	≤ 0.050	0.001
of Solution	A420 - A720	≤ 0.100	0.021
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	$\leq 100 \text{ CFU/g}$	$\leq 10 \text{ CFU/g}$
	TYMC	$\leq 100 \text{ CFU/g}$	$\leq 10 \text{ CFU/g}$
² Nitrogen Determination		$\leq 0.005\%$	≤ 0.005 %
² Optical Rotation, Sp 20°C	ecific Rotation @	+197° to +201°	+ 199°
²pH @ 25°C		4.5 - 6.5	5.5
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%

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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%	9.8%

		EP COMPENDIA	
A	NALYSIS	SPECIFICATION	TEST RESULT
¹ Assay		³ 98.0 - 101.0%	99.6%
Appearance of S	Solution	Clear, colorless	Clear, colorless
Chlorides		≤ 0.0125%	\leq 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%
Substances	Unspecified Impurities	≤ 0.2%	$\leq 0.2\%$
	Total Impurities	≤ 1.0%	$\leq 1.0\%$
	Escherichia coli	Absent/g	Absent/g
² Microbial	Salmonella species	Absent/10g	Absent/10g
Content	TAMC	$\leq 100 \text{ CFU/g}$	< 10CFU/g
	TYMC	$\leq 100 \text{ CFU/g}$	< 10CFU/g
²pH @ 25°C		4.5 - 6.5	5.5
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%	9.8%

JP COMPENDIA		
Analysis	SPECIFICATION	TEST RESULT
¹ Assay	98.0% - 101.0%	99.6%
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%	< 0.005%
² Optical Rotati	on @ 2 0°C	+197° to +201°	+ 199°
²pH @ 25°C		4.5 - 6.5	5.5
² Residue on Ign	nition	≤ 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		$\leq 0.024\%$	$\leq 0.024\%$
² Water		9.0% to 11.0%	9.8%

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:	Date: $\frac{5/z}{z^2}$	Job Title: QA specialist
Reviewed by:	Date: 5/2/22	Job Title: DA Manager

²Analyses are Harmonized

³Specifications is more stringent than Compendia Monograph