## BISPECTRA

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

Effective Date:	21-Jun-2022	21-Jun-2025	: Date of Next Review
Prepared By:	Crystal Hamelburg	BSI-COA-0098 v.4.0	: Supersedes
QA/QC Approval:	Wendy Santay	Dora Meissner	: Management Approval
Reason for Revision:	See Revision History in MasterControl		

## **CERTIFICATE OF ANALYSIS**

## D-GALACTOSE, PLANT DERIVED

## BIO EXCIPIENT GRADE / NEW CODE GALP-3250-10

(HISTORICAL CODE GA3250-K010)

LOT: GALP-0124-00085

C<sub>6</sub>H<sub>12</sub>O<sub>6</sub> ♣ F.W. 180.16 g/mol. ♣ CAS# 59-23-4

Manufacturing Date: 05/16/24 Retest Date: 05/31/26 Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 10/17/24 Packaging Site: 100 Majestic Way, Bangor PA, 18013

Meets or Exceeds EP and NF Specifications

EP COMPENDIA				
Analysis	SPECIFICATION	TEST RESULT		
Acidity or Alkalinity	Passes Test	Passes Test		
Appearance	White to almost white, crystalline powder	White to almost white, crystalline powder		
Appearance of Solution	Passes Test	Passes Test		
Assay	97.0 – 102.0%	99.5 %		
Barium	Passes Test	Passes Test		
Identification A	Passes Test	Passes Test		
Identification B	Passes Test	Passes Test		
Identification C	Passes Test	Passes Test		
Microbial Content TAMC	$\leq 100 \text{ CFU/g}$	< 10 CFU/g		
Proteins	$\leq 0.1 \text{ mg/mL}$	<0.1 mg/ml		
Impurities A and B	≤ 1.0%	<0.05 %		
Related Unspecified Substances Impurities	≤ 0.3% each	< 0.05 %		
Total Impurities	≤ 2.0%	0.07 %		
Sulfated Ash	≤ 0.1%	< 0.1 %		
Water	≤ 1.0%	0.3 %		

			BSI-COA-0098 v.4.
		NF COMPENDIA	
Analysis		SPECIFICATION	TEST RESULT
Acidity		Passes Test	Passes Test
Appearance of So	lution	Passes Test	Passes Test
Assay		98.0-102.0%	99.5%
Barium		Passes Test	Passes Test
Identification A		Passes Test	Passes Test
Identification B		Passes Test	Passes Test
Identification C		Passes Test	Passes Test
Limit of Lead		≤ 0.5 ppm	< 0.005 ppm
	Escherichia coli	Absent	Absent
1	Pseudomonas aeruginosa	Absent	Absent
Microbial	Salmonella species	Absent	Absent
Content	Staphylococcus aureus	Absent	Absent
	TAMC	≤ 1000 CFU/g	< 10 CFU/g
	TYMC	≤ 100 CFU/g	< 10 CFU/g
Lactose and 1,6-galactosyl- galactose		≤0.6%	< 0.05%
	Galacturonic acid	≤0.6%	< 0.05%
	Dextrose	≤0.6%	< 0.05%
Related	Tagatose	≤0.6%	< 0.05%
Substances	Dulcitol	≤0.6%	< 0.05%
	Arabinose	≤0.6%	0.07%
Any unspecified impurity  Total Impurities		≤0.2%	< 0.05%
		≤1.0%	0.07%
Residue on Ignition		≤ 0.1%	< 0.1%
Optical Rotation,	Specific Rotation	+78.0° to +81.5°	+80.5°
Water		≤ 1.0%	0.3%

ANALYSIS	SPECIFICATION	TEST RESULT
Endotoxins	≤2.5 EU/g	< 1.0 EU/g
Glucose	≤ 0.1%	< 0.05%
Lead	≤ 0.5 ppm	< 0.005 ppm
Residual Ethanol	≤ 500 ppm	< 240 ppm
Residual Isopropanol	≤ 5000 ppm	< 2520 ppm
Residual Methanol	≤ 100 ppm	< 80 ppm
Residual Methyl Isobutyl Ketone	≤ 500 ppm	< 250 ppm

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: BSI-ATM-0026

<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: Daullall	_ Date: _	10/22/24	Job Title: QATCCh 111
Reviewed by: Jan Burth	_ Date: _	10/23/24	_ Job Title: QA Supervisor

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