DCN: 19-002872 v.4.0



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

Effective Date: 06 Apr 2021		06 Apr 2024	: Date of Next Review
Prepared By: Jared L Lobb		19-002872 v.3.0	: Supersedes
QA/QC Approval: Carissa McCollian		Wendy Santay	: Management Approval
Reason for Revision: See Revision History in	ensur.		

CERTIFICATE OF ANALYSIS

D-GALACTOSE, PLANT DERIVED

BIO EXCIPIENT GRADE / GALP-3251-93

(HISTORICAL CODE GA3251-G500)

LOT: GALP-0121-00020

C₆H₁₂O₆ → F.W. 180.16 g/mol. → CAS# 59-23-4

Manufacturing Date: 10/1/21 Retest Date: 10/31/23

Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 10/21/21 Packaging Site: 100 Majestic Way, Bangor PA, 18013

EP COMPENDIA				
ANA	LYSIS	SPECIFICATION	TEST RESULT	
² Acidity or Alkali	nity	Passes Test	Passes Test	
Appearance		White to almost white, crystalline or finely granulated powder	White to almost white, crystalline or finely granulated powder	
² Appearance of So	olution	Passes Test	Passes Test	
¹ Assay		398.0%-102.0%	99.5%	
² Identification A		Conforms to Reference	Conforms to Reference	
¹ Identification B		Passes Test	Passes Test	
² Identification C		Passes Test	Passes Test	
² Microbial Conten	nt TAMC	$\leq 100 \text{ CFU/g}$	<10 CFU/g	
Proteins		$\leq 0.1 \text{ mg/mL}$	< 0.1 mg/mL	
Substances Impurities	≤ 1.0%	0.1%		
	Unspecified Impurities	≤ 0.3%	<0.3%	
	Total Impurities	≤ 2.0%	0.1%	
Sulfated Ash		≤ 0.1%	<0.1%	
² Water		≤ 1.0%	0.1%	

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		NF COMPENDIA	DCN: 19-002072 v.4.0
1	Analysis	SPECIFICATION	TEST RESULT
² Acidity		Passes Test	Passes Test
² Appearance of	Solution	Passes Test	Passes Test
¹ Assay		98.0 - 102.0%	99.5%
Barium		Passes Test	Passes Test
² Identification A		Conforms to Reference	Conforms to Reference
¹ Identification B		Passes Test	Passes Test
² Identification C		Passes Test	Passes Test
¹ Limit of Lead		≤ 0.5 ppm	<0.5 ppm
	Escherichia coli	Absent	Absent
	Pseudomonas aeruginosa	Absent	Absent
² Microbial	Salmonella species	Absent	Absent
Content	Staphylococcus aureus	Absent	Absent
	TAMC	$^3 \le 100 \text{ CFU/g}$	<10 CFU/g
	TYMC	$\leq 100 \text{ CFU/g}$	<10 CFU/g
	Lactose and 1,6- galactosyl- galactose	≤ 0.6%	0.1%
	Galacturonic Acid	≤ 0.6%	<0.6%
	Dextrose	≤ 0.6%	<0.6%
¹ Related	Tagatose	≤ 0.6%	<0.6%
Substances	Dulcitol	≤ 0.6%	<0.6%
	Arabinose	≤ 0.6%	0.1%
	Any Unspecified Impurity	≤ 0.2%	<0.2%
	Total Impurities	≤ 1.0%	0.1%
Residue on Ignit		≤ 0.1%	<0.1 %
	, Specific Rotation	+78.0° to +81.5°	+80.4°
² Water		≤ 1.0%	0.1%

ADDITIONAL ANALYSES

An	ALYSIS	SPECIFICATION	TEST RESULT
Endotoxins		≤ 2.5 EU/g	<0.5 EU/g
¹ Glucose		≤ 0.1%	<0.1%
	Aluminum (Al)	≤ 400 ppb	<400 ppb
	Cadmium (Cd)	≤ 10 ppb	<10 ppb
	Cobalt (Co)	≤ 50 ppb	<50 ppb
	Chromium (Cr)	≤ 50 ppb	<50 ppb
	Copper (Cu)	≤ 25 ppb	<25 ppb
Two as Matala	Iron (Fe)	≤ 200 ppb	<200 ppb
Trace Metals	Manganese (Mn)	≤ 25 ppb	<25 ppb
	Molybdenum (Mo)	≤ 50 ppb	<50 ppb
	Nickel (Ni)	≤ 50 ppb	<50 ppb
	Selenium (Se)	≤ 50 ppb	<50 ppb
	Vanadium (V)	≤ 50 ppb	<50 ppb
	Zinc (Zn)	≤ 200 ppb	<200 ppb
¹ Residual Ethano	1	≤ 500 ppm	<500 ppm
¹ Residual Isoprop	panol	≤ 5000 ppm	<5000 ppm
¹ Residual Methanol		≤ 100 ppm	<100 ppm
¹ Residual Methyl	Isobutyl Ketone	≤ 500 ppm	<500 ppm

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 18-002374

<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: 1/22/21 Job Title: DA Specialist

Reviewed by: Date: 1/122/21 Job Title: DA Manager

¹Alternate Validated Method

²Analyses are Harmonized

³Specification is more stringent than Compendia Monograph