DCN: BSI-COA-0128 v. 4.1



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22.4								[0.4		

Effective Date:	04-APR-2024	04-APR-2027	: Date of Next Review
Prepared By:	Carissa Albert	BSI-COA-0128 v. 4.0	: Supersedes
QA/QC Approval:	Jaron Hughes	Wayne Talamonti	: Management Approval
Reason for Revision:	See Revision History in MasterControl.		

CERTIFICATE OF ANALYSIS D-GALACTOSE, PLANT DERIVED BIO EXCIPIENT GRADE / GALP-3251-21

LOT: GALP-0124-00062

C₆H₁₂O₆ ♣ F.W. 180.16 g/mol. ♣ CAS# 59-23-4 Manufacturing Date: 03/31/24 Retest Date: 03/31/26 Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 04/29/24 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		³ 98.0%-102.0%	99.4%				
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 Conter	nt TAMC	$\leq 100 \text{ CFU/g}$	<10 CFU/g				
Proteins		$\leq 0.1 \text{ mg/mL}$	<0.1 mg/mL				
	Sum of Impurities A and B	≤ 1.0%	<0.05%				
¹ Related Substances	Unspecified Impurities	≤ 0.3%	<0.05%				
	Total Impurities	≤ 2.0%	<0.05%				
Sulfated Ash		≤ 0.1%	<0.1%				
² Water		≤ 1.0%	0.3%				

		NF COMPENDIA	
	Analysis	SPECIFICATION	TEST RESULT
² Acidity		Passes Test	Passes Test
² Appearance	of Solution	Passes Test	Passes Test
¹ Assay		98.0 - 102.0%	99.4%
Barium		Passes Test	Passes Test
² Identification	n A	Conforms to Reference	Conforms to Reference
¹ Identification	n B	Passes Test	Passes Test
² Identification	n C	Passes Test	Passes Test
¹ Limit of Lea	d	≤ 0.5 ppm	<0.005 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%	<0.05%
	Galacturonic Acid	≤ 0.6%	<0.05%
	Dextrose	≤ 0.6%	<0.05%
¹ Related	Tagatose	≤ 0.6%	<0.05%
Substances	Dulcitol	≤ 0.6%	<0.05%
	Arabinose	≤ 0.6%	<0.05%
	Any Unspecified Impurity	≤ 0.2%	<0.05%
	Total Impurities	≤ 1.0%	<0.05%
Residue on Ig	gnition	≤ 0.1%	<0.1 %
Optical Rotat @ 20°C	ion, Specific Rotation	+78.0° to +81.5°	+80.7°
² Water		≤ 1.0%	0.3%

An	NALYSIS	SPECIFICATION	TEST RESULT
Endotoxins	-	≤ 2.5 EU/g	<1.0 EU/g
¹ Glucose		≤ 0.1%	<0.05%
	Aluminum (Al)	≤ 400 ppb	<400 ppb
	Cadmium (Cd)	≤ 10 ppb	<6 ppb
	Cobalt (Co)	≤ 50 ppb	<5 ppb
	Chromium (Cr)	≤ 50 ppb	<50 ppb
	Copper (Cu)	≤ 25 ppb	<25 ppb
Trace Metals	Iron (Fe)	≤ 200 ppb	<200 ppb
Trace Metals	Manganese (Mn)	≤ 25 ppb	<25 ppb
	Molybdenum (Mo)	≤ 50 ppb	<50 ppb
	Nickel (Ni)	≤ 50 ppb	<20 ppb
	Selenium (Se)	≤ 50 ppb	<50 ppb
	Vanadium (V)	≤ 50 ppb	<10 ppb
	Zinc (Zn)	≤ 200 ppb	<200 ppb
¹ Residual Ethan	ol	≤ 500 ppm	<240 ppm
¹ Residual Isopro	panol	≤ 5000 ppm	<2510 ppm
¹ Residual Metha	anol	≤ 100 ppm	<80 ppm
¹ Residual Methy	yl 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: And Mall Date: 5/7/24 Job Title: QA Tech I

Reviewed by: Ann Bright Date: 5/7/24 Job Title: QA Supervisor

¹Alternate Validated Method

²Analyses are Harmonized

³Specification is more stringent than Compendia Monograph

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