DCN: 20-003401 v.2.0

BIOSPECTRA

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

Effective Date:	01-Mar-2021		01-Mar-2024	: Date of Next Review
		┥	0.1.1.1	
Prepared By:	Jared L Lobb	4	20-003401 v.1.2	: Supersedes
QA/QC Approval:	Jaron Hughes		Wendy Santay	: Management Approval
Reason for Revision:	See Revision History in ensur			

CERTIFICATE OF ANALYSIS

MES MONOHYDRATE

BIO EXCIPIENT GRADE / NEW CODE MESM-3222-50

(HISTORICAL CODE ME3222-K050)

LOT: MESM-0121-0119

C₆H₁₃NO₄S·H₂O ^A F.W. 213.3 g/mol. ^A CAS# 145224-94-8 Manufacturing Date: 3/2/21 Retest Date: 3/31/23 Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 3/4/21 Packaging Site: 100 Majestic Way, Bangor PA, 18013

Analysis		SPECIFICATION	TEST RESULT
Absorbance	260 nm	0.1000 a.u. max.	0.0027 a.u.
(1M)	280 nm	0.1000 a.u. max.	0.0016 a.u.
Appearance and Color		White Crystalline Powder	White Crystalline Powder
Assay		99.5% min.	99.7%
Chloride		0.005% max.	<0.005%
Color (1M, Alkaline)		Colorless	Colorless
DNase		None Detected	None Detected
Enzymes	RNase	None Detected	None Detected
	Protease	None Detected	None Detected
Heavy Metals (as Pb)		2 ppm max.	< 2 ppm
Identification (IR)		Conforms to Reference	Conforms to Reference
Loss on Drying @ 130°C		7 - 10%	9%
pH (5% Soln.)		3.1 - 3.5	3.4
pH (1.0M)		2.7 - 3.7	3.1
pH (0.5M)		2.5 - 4.5	3.3
pK_a		5.9 - 6.3	6.1
Turbidimetry/PVS Limit Test		≤ 1 ppm	≤1 ppm
Residue on Ignition		0.05% max.	<0.01%
Solubility (5%)		Passes Test	Passes Test
Sulfate		0.005% max.	<0.005%
	Arsenic (As)	2 ppm max.	< 2 ppm
Trace Elements	Copper (Cu)	2 ppm max.	< 2 ppm
Trace Elements	Iron (Fe)	2 ppm max.	< 2 ppm
	Lead (Pb)	2 ppm max.	< 2 ppm

8.8%

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-001016

INTENDED USE: 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.

RESIDUAL SOLVENT STATEMENT: Based on the manufacturing process and the controlled handling, storage and analysis of this product, this product complies with the requirements and specifications listed in the current USP method <467> Tables 1, 2, 3, or 4.

Date: 4/1/21 Job Title: QA Manager

Reviewed by: