DCN: 16-001185 v.6.0

BIOSPECTRA

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

Effective Date: 0	01-Mar-2021	01-Mar-2024	: Date of Next Review
Prepared By: Ja	ared L Lobb	16-001185 v.5.0	: Supersedes
QA/QC Approval: Ja	aron Hughes	Wendy Santay	: Management Approval
Peason for Revision: S	See Revision History in ensur		

CERTIFICATE OF ANALYSIS HEPES

BIO EXCIPIENT GRADE / NEW CODE HEPE-3220-25

(HISTORICAL CODE HE3220-K025)

LOT: HEPE-0122-00029

C₈H₁₈N₂O₄S ^ F.W. 238.30 g/mol. ^ CAS# 7365-45-9 Manufacturing Date: 2/12/22 Retest Date: 2/29/24 Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 2/19/22

Packaging Site: 100 Majestic Way, Bangor PA, 18013

Analysis	S	SPECIFICATION	TEST RESULT	
Absorbance (0.1M)	250 nm	0.0500 a.u. max.	0.0087 a.u.	
	260 nm	0.0500 a.u. max.	0.0041 a.u.	
	280 nm	0.0800 a.u. max.	0.0032 a.u.	
Absorbance (0.05M)	250 nm	0.0500 a.u. max.	0.0035 a.u.	
	260 nm	0.0500 a.u. max.	0.0014 a.u.	
	280 nm	0.0800 a.u. max.	0.0010 a.u.	
Appearance and Color		White / Crystals	White / Crystals	
Assay, Dried Basis		99.5% min.	100.2%	
Chloride		0.005% max.	< 0.005%	
Endotoxins		≤ 5 EU/g	< 1 EU/g	
Enzymes	DNase	None Detected	None Detected	
	RNase	None Detected	None Detected	
	Protease	None Detected	None Detected	
Heavy Metals		1 ppm max.	< 1 ppm	
Identification (IR)		Passes Test	Passes Test	
Insoluble Matter		0.01% max.	0.01%	
Loss on Drying		0.5% max.	0.3%	
Microbial Content	TAMC	≤ 100 CFU/g	< 10 CFU/g	
	TYMC	≤ 100 CFU/g	< 10 CFU/g	
pH (5% Soln)		5.0 – 6.5	5.1	
p K a	•	7.45 – 7.65	7.54	

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Anal	YSIS	SPECIFICATION	Test Result
Residue on Ignition	1	0.1% max.	< 0.1%
Solubility (5%)		Passes Test	Passes Test
Solubility (0.05M)		Passes Test	Passes Test
Sulfate		0.005% max.	< 0.005%
Trace Metals	Arsenic (As)	5 ppm max.	< 5 ppm
	Copper (Cu)	5 ppm max.	< 5 ppm
	Iron (Fe)	5 ppm max.	< 5 ppm
	Lead (Pb)	5 ppm max.	< 5 ppm

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-001305

<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.

<u>RESIDUAL SOLVENT STATEMENT:</u> 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.

Prepared by:	_ Date: _	5/13/22	Job Title: OA Specialist
	_		,
Reviewed by:	_ Date: _	0/16/22	Job Title: A Specialist
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