DCN: 16-001185 v.3.0



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

Effective Date: 10-Apr-2020	10-Apr-2023	: Date of Next Review
Prepared By: Kyle Snyder	16-001185 v.2.1	: Supersedes
QA/QC Approval: Carissa McCollian	Hannah Bernier	: Management Approval
Reason for Revision: See Revision History in ensur.		

CERTIFICATE OF ANALYSIS HEPES

BIO EXCIPIENT GRADE / HE3220-SAMPLE COFA

LOT: HE3220-013-0520

C₈H₁₈N₂O₄S A F.W. 238.30 g/mol. A CAS# 7365-45-9 Manufacturing Date: 10/8/2019 Retest Date: 10/31/2021 Manufacturing Site: 100 Majestic Way, Bangor PA, 18013 Packaging Date: Sample CofA Packaging Site: Sample CofA

ANALY	YSIS	SPECIFICATION	TEST RESULT
Absorbance	250 nm	0.0500 a.u. max.	0.0092 a.u.
	260 nm	0.0500 a.u. max.	0.0050 a.u.
	280 nm	0.0800 a.u. max.	0.0044 a.u.
Appearance and Co	lor	White / Crystals	White / Crystals
Assay, Dried Basis		99.5% min.	99.8%
Chloride		0.005% max.	< 0.005%
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.	0.1%
pH (5% Soln)		5.0 - 6.5	5.3 @ 25.0°C
pK_a		7.45 - 7.65	7.50
Residue on Ignition		0.1% max.	<0.1%
Solubility		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

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COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-001305

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.

OVI 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.

Prepared by: Willy fluttey Date: 09/12/20 Job Title: QH Manager

Reviewed by: _______ Date: _______ Date: _______ Job Title: ______ QA Superviser