

# BIOSPECTRA

100 Majestic Way, Bangor, PA 18013 / [www.biospectra.us](http://www.biospectra.us)

Effective Date:	3-Jun-2020	3-Jun-2023	: Date of Next Review
Prepared By:	Amy Hosein	16-001185 v.4.0	: Supersedes
QA/QC Approval:	Wendy Santay	Amy Yenko	: Management Approval
Reason for Revision:	See Revision History in ensur.		

## CERTIFICATE OF ANALYSIS

### HEPES

### BIO EXCIPIENT GRADE / HE3220-K001

### LOT: HE3220-022-0620

C<sub>8</sub>H<sub>18</sub>N<sub>2</sub>O<sub>4</sub>S \* F.W. 238.30 g/mol. \* CAS# 7365-45-9

Manufacturing Date: 4/1/20 Retest Date: 4/30/22

Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 6/22/20 Packaging Site: 100 Majestic Way, Bangor PA, 18013

ANALYSIS		SPECIFICATION	TEST RESULT
Absorbance	250 nm	0.0500 a.u. max.	0.0103 a.u.
	260 nm	0.0500 a.u. max.	0.0055 a.u.
	280 nm	0.0800 a.u. max.	0.0047 a.u.
Appearance and Color		White / Crystals	White / Crystals
Assay, Dried Basis		99.5% min.	100.1%
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.	0.1%
Microbial Content	TAMC	≤ 100 CFU/g	<10 CFU/g
	TYMC	≤ 100 CFU/g	<10 CFU/g
pH (5% Soln)		5.0 – 6.5	5.2
pK <sub>a</sub>		7.45 – 7.65	7.48
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

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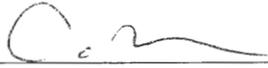
ANALYSIS	SPECIFICATION	TEST RESULT	
Trace Metals	Copper (Cu)	5 ppm max.	< 5 ppm
	Iron (Fe)	5 ppm max.	< 5 ppm
	Lead (Pb)	5 ppm max.	< 5 ppm

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-001305

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.

Prepared by:  Date: 10/23/20 Job Title: QA Supervisor

Reviewed by:  Date: 10/23/20 Job Title: QA Manager