


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

Effective Date:	3-Feb-2025	3-Feb-2028	: Date of Next Review
Prepared By:	Taylor Yurick	BSI-COA-0036 v.6.1	: Supersedes
QA/QC Approval:	Jaron Hughes	Carissa Albert	: Management Approval
Reason for Revision:	See Revision History in MasterControl.		

CERTIFICATE OF ANALYSIS

HEPES

BIO EXCIPIENT GRADE / HEPE-3220

LOT: HEPE-E03-1025-0089

C8H18N2O4S F.W. 238.30 g/mol. CAS# 7365-45-9

Manufacturing Date: 10/20/25 Retest Date: 10/31/27

Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Site: 100 Majestic Way, Bangor PA, 18013

ANALYSIS	SPECIFICATION		TEST RESULT
Absorbance (0.1M)	250 nm	0.0500 a.u. max.	0.0130 a.u.
	260 nm	0.0500 a.u. max.	0.0085 a.u.
	280 nm	0.0800 a.u. max.	0.0080 a.u.
Absorbance (0.05M)	250 nm	0.0500 a.u. max.	0.0067 a.u.
	260 nm	0.0500 a.u. max.	0.0043 a.u.
	280 nm	0.0800 a.u. max.	0.0042 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
	DNase	None Detected	None Detected
	RNase	None Detected	None Detected
Enzymes	Protease	None Detected	None Detected
	1 ppm max.		< 1 ppm
	Passes Test		Passes Test
Heavy Metals	0.01% max.		< 0.01%
Identification (IR)	0.5% max.		0.1%
Insoluble Matter	≤ 100 CFU/g		< 10 CFU/g
Loss on Drying	≤ 100 CFU/g		< 10 CFU/g
Microbial Content	TAMC	≤ 100 CFU/g	
	TYMC	≤ 100 CFU/g	

The information contained herein is the confidential property of BioSpectra. The recipient is responsible for its safe-keeping and the prevention of unauthorized appropriation, use, disclosure and copying.

ANALYSIS	SPECIFICATION	TEST RESULT
pH (5% Soln)	5.0 – 6.5	5.3
pK _a	7.45 – 7.65	7.60
Residue on Ignition	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) Copper (Cu) Iron (Fe) Lead (Pb)	5 ppm max. 5 ppm max. 5 ppm max. 5 ppm max.
		< 5 ppm

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: BSI-ATM-0070

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: Phil McAll Date: 1/30/26 Job Title: QA Tech 111

Reviewed by: Jean Engle Date: 1/30/26 Job Title: QA Supervisor