

HEPES, Free Acid, LBLE, GMP Excipient Grade

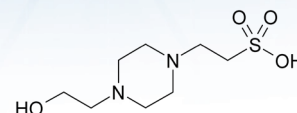
Low Bioburden, Low Endotoxin, GMP Manufactured

INTENDED FOR USE AS AN EXCIPIENT

HEPES is a zwitterionic buffer used to maintain pH of media used in cell cultures. It is one of Good's twelve buffers that has a pK_a value similar to its pH value making it an ideal buffer for pH maintenance. It has only one restriction, it interferes with the Folin protein assay. This buffer can form radicals so it is not suitable for redox studies. HEPES is used as a Good's buffer because it has low UV absorptivity, minimal reactivity, stable pH and is soluble in water.

Lead Time: 3-6 months

Minimum Order Quantity: 250kg



CAS #: 7365-45-9

Molecular Formula: $C_8H_{18}N_2O_4S$

Solubility in Water (g/L): 400

F.W.: 238.30 g/mol

pH @ 20°C: 5.0 - 6.5

Useful pH: 6.8 - 8.2

pK_a @ 20°C: 7.55

BIO EXCIPIENT GRADE | Product Code: HEPE-3250 | Previously: HE3250

$C_8H_{18}N_2O_4S$ • F.W. 238.30 g/mol. • CAS# 7365-45-9



These are general specifications. BioSpectra will customize our products to meet your quality based requirements.

ANALYSIS		SPECIFICATIONS
Absorbance (0.1M)	250 nm	≤ 0.0500 a.u.
	260 nm	≤ 0.0500 a.u.
	280 nm	≤ 0.0800 a.u.
Appearance and Color		White / Crystals
Appearance of Solution (1% Water)		Conforms
Assay, Dried Basis		99.5% min.
Chloride		≤ 50 ppm
Endotoxins		≤ 5 EU/g
Enzymes	DNases (Exo- and endonucleases)	None Detected
	RNase	None Detected
	Protease	None Detected
Heavy Metals (as Pb)		≤ 1 ppm
Identification (IR)		Passes Test
Identification (TLC)		Conforms
Insoluble Matter		≤ 0.01%
Loss on Drying (105°C)		≤ 0.2%
Methanol		≤ 3000 ppm



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ANALYSIS		SPECIFICATIONS
Microbial Content	Bile tolerant Gram negative Bacteria (Absent in 1g)	Conforms
	Candida albicans (Absent in 1g)	Conforms
	Escherichia coli (Absent in 1g)	Conforms
	Pseudomonas aeruginosa (Absent in 1g)	Conforms
	Salmonella species (Absent in 10g)	Conforms
	Staphylococcus aureus (Absent in 1g)	Conforms
	TAMC	≤ 100 CFU/g
TAMC	≤ 50 CFU/g	
pH (5% Soln)		5.0 – 6.5
pH (1% Water)		4.7 – 5.3
pK _a		7.45 – 7.65
Residue on Ignition		≤ 0.1%
Solubility (5%)		Passes Test
Sulfate		≤ 50 ppm
Sulfated Ash (600°C)		≤ 0.10%
Trace Metals	Arsenic (As)	≤ 5 ppm
	Calcium (Ca)	≤ 50 ppm
	Cadmium (Cd)	≤ 5 ppm
	Cobalt (Co)	≤ 5 ppm
	Copper (Cu)	≤ 5 ppm
	Chromium (Cr)	≤ 5 ppm
	Iron (Fe)	≤ 5 ppm
	Lead (Pb)	≤ 1 ppm
	Magnesium (Mg)	≤ 10 ppm
	Manganese (Mn)	≤ 5 ppm
	Nickel (Ni)	≤ 5 ppm
Potassium (K)	≤ 50 ppm	
Zinc (Zn)	≤ 5 ppm	

General Product Description:

- The manufacturing of HEPES, HEPE-3250 is performed at BioSpectra's Bangor and Stroudsburg, PA facilities utilizing multi-use equipment. Equipment used in the manufacturing of HEPE-3250 is cleaned in accordance with BioSpectra's Process Cleaning Validation Master Plan.
- HEPES is a White Crystalline Product.
- Molecular Formula: C₈H₁₈N₂O₄S
- Molecular Weight: 238.30 g/mol.
- CAS Number: 7365-45-9.
- There are no known major food allergens (as defined by FDA and WHO) in the manufacture of this product.
- BioSpectra certifies that all HEPES HEPE-3250 manufactured at BioSpectra and its raw materials are not derived from or come in contact with animal parts, products, and/or byproducts.
- HEPES manufactured at BioSpectra and any raw materials used in the manufacture of HEPES at BioSpectra are not subject to genetic modification.
- Synonyms: N-(2-Hydroxyethyl)piperazine-N'-2-ethanesulfonic acid, 2-[4-(2-hydroxyethyl)piperazin-1-yl]ethanesulfonic acid.

GMP Compliance:

Bio Excipient Grade HEPES, HEPE-3250 is suitable for use as an excipient. It is manufactured in accordance with the ICH-Q7 Good Manufacturing Practice Guide. This grade of HEPES is not suitable to be used as an Active Pharmaceutical Ingredient, Drug Product or Household Item.

Retest Date:

The recommended retest period for HEPES is two years from the date of manufacture.

Storage and Shipping Conditions:

Ship and store in ambient temperature. Store in a clean and dry area. Store in the original container.

Package Sizes:

10kg, 25kg and 50kg pails.

ELEMENTAL IMPURITIES: This product complies with ICHQ3D, USP <232> and USP <233> requirements for Elemental Impurities.

RESIDUAL SOLVENTS: 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.