

BIO SPECTRA

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

Effective Date:	24-Aug-2023	24-Aug-2026	: Date of Next Review
Prepared By:	Amy Yenko	BSI-COA-0017 v.4.1	: Supersedes
QA/QC Approval:	Wayne Talamonti	Carissa Albert	: Management Approval
Reason for Revision:	See Revision History in MasterControl.		

CERTIFICATE OF ANALYSIS
GUANIDINE HYDROCHLORIDE
BIO EXCIPIENT GRADE / NEW CODE GHCL-3220-25
(HISTORICAL CODE GH3220-K025)
LOT#: GHCL-0124-00037

NH₂C(NH)NH₂·HCl * FW: 95.53 g mol. * CAS# 50-01-1

Manufacturing Date: 03/20/24 Retest Date: 03/31/26

Manufacturing Site: 1474 Rockdale Lane, Stroudeburg, PA 18360

Packaging Date: 05/25/24 Packaging Site: 100 Majestic Way, Bangor PA, 18013

ANALYSIS	SPECIFICATIONS	RESULT
Acidity	≤ 0.01%	<0.01%
Appearance and Color	White / Crystals	White / Crystals
Assay (Dried Basis)	99.5 – 101.0%	99.9%
Chloride and Sulfate, <i>Sulfate</i>	≤ 0.005%	<0.005%
Enzymes	DNase	None Detected
	Protease	None Detected
	RNase	None Detected
Identification A, (IR)	Passes Test	Passes Test
Identification B, <i>Absorbance</i>	230nm	≤ 0.2000 a.u.
	260nm	≤ 0.0300 a.u.
	275nm	≤ 0.0300 a.u.
Identification C, <i>Chloride</i>	Meets the Requirements of Test A	Meets the Requirements of Test A
Limit of Nitrate	≤ 0.005%	<0.005%
Loss on Drying	≤ 0.5%	0.1%
Melting Range	184 - 188°C	186 - 187°C
pH (6M)	4.5 - 6.0	5.6 @ 23.1°C
Residue on Ignition	≤ 0.05%	0.01%
Solubility (6M)	Passes Test	Passes Test
Trace Metals	Arsenic (As)	≤ 5 ppm
	Copper (Cu)	≤ 5 ppm
	Iron (Fe)	≤ 5 ppm
	Lead (Pb)	≤ 5 ppm
Water by Karl Fischer	≤ 0.3% w/w	<0.1% w/w
Water Insoluble	≤ 0.05%	<0.05%

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

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: BSI-ATM-0013

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 SOLVENTS 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: Taylor Yuick Date: 6/5/24 Job Title: QA Tech III

Reviewed by: Jason Bligh Date: 6/5/24 Job Title: QA Supervisor