

BIOSPECTRA

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

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|----------------------|-------------------------------|----------------|-----------------------|
| Effective Date: | 30-Jan-2017 | 30-Jan-2020 | : Date of Next Review |
| Prepared By: | Jamie Storm | 16-001152 v2.0 | : Supersedes |
| QA/QC Approval: | Crystal Hamelburg | Chad Pezoldt | : Management Approval |
| Reason for Revision: | See Revision History in ensur | | |

GUANIDINE THIOCYANATE

CERTIFICATE OF ANALYSIS

BIO EXCIPIENT GRADE / GT3220-K001

LOT: GT3220-011-0118

$\text{NH}_2\text{C}(\text{NH})\text{NH}_2\text{-HSCN}$ * F.W. 118.16 g/mol * CAS#: 593-84-0
 Manufacturing Date: 11/01/2016 Retest Date: 11/30/2018
 Manufacturing Site: 100 Majestic Way, Bangor PA, 18013 Packaging Date: 01/15/2018
 Packaging Site: 100 Majestic Way, Bangor PA, 18013

| ANALYSIS | | SPECIFICATION | TEST RESULT |
|----------------------|--------------|------------------|------------------|
| Absorbance | 280nm | 0.300 a.u. max. | 0.0980 a.u. |
| | 300nm | 0.050 a.u. max. | 0.0147 a.u. |
| | 340nm | 0.030 a.u. max. | 0.0040 a.u. |
| Appearance and Color | | White / Crystals | White / Crystals |
| Assay | | 99.5% min. | 99.62% |
| Enzymes | DNase | None Detected | None Detected |
| | RNase | None Detected | None Detected |
| | Protease | None Detected | None Detected |
| Identity (IR) | | Passes Test | Passes Test |
| Loss On Drying | | 0.5% max. | 0.1057% |
| Melting Range | | 115-121°C | 119.5 – 120.6°C |
| pH (5% Solution) | | 5.0 – 7.0 | 5.48 @ 20.4°C |
| Solubility (35%) | | Passes Test | Passes Test |
| 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 |

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-000043

INTENDED USE: Material represented by this Certificate of Analysis is suitable to be used only as the following: ICH Q7 Compliant cGMP Manufactured Excipient for use in further manufacturing. 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 and storage of this product, there is no potential for any of the residual solvents listed in the current USP method <467> Tables 1, 2, 3, or 4 to be present at the specified limits; furthermore, if tested this product would comply with USP/NF requirements.

Prepared by: H. Bennett Date: 1/17/18

Reviewed by: C. M. [unclear] Date: 1/17/18