

# BIOSPECTRA

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

|                      |                                |               |                       |
|----------------------|--------------------------------|---------------|-----------------------|
| Effective Date:      | 15-APR-2020                    | 15-APR-2023   | : Date of Next Review |
| Prepared By:         | Amy Hosein                     | v.1.0         | : Supersedes          |
| QA/QC Approval:      | Wendy Santay                   | Dora Meissner | : Management Approval |
| Reason for Revision: | See Revision History in ensur. |               |                       |

## CERTIFICATE OF ANALYSIS

### UREA

BIO ACTIVE GRADE / UR2220 – K050

LOT: UR2220-016-0820

$\text{NH}_2\text{CONH}_2$  ^ F.W. 60.06 g/mol. ^ CAS# 57-13-6

Manufacture Date: 8/20/20 Retest Date: 8/31/23

Manufacturing Site: 100 Majestic Way, Bangor PA, 18013

Packaging Date: 8/23/20 Packaging Site: 100 Majestic Way, Bangor PA, 18013

Meets or Exceeds USP Specifications

| ANALYSIS                 | SPECIFICATION    | TEST RESULT      |
|--------------------------|------------------|------------------|
| Alcohol Insoluble Matter | 0.04% max.       | <0.04%           |
| Appearance and Color     | White / Crystals | White / Crystals |
| Assay                    | 98.0-102.0%      | 99.9%            |
| Enzymes                  | DNase            | None Detected    |
|                          | RNase            | None Detected    |
|                          | Protease         | None Detected    |
| Heavy Metals             | 10 ppm max.      | <10ppm           |
| Identification A (IR)    | Passes Test      | Passes Test      |
| Identification B         | Passes Test      | Passes Test      |
| Impurities               | Urea RCA         | < 0.1%           |
|                          | Total            | < 2.0%           |
|                          | Unspecified      | < 0.1%           |
| Insoluble Matter         | 0.010% max.      | <0.002%          |
| Loss on Drying           | 1.0% max.        | 0.2%             |
| Melting Range            | 132-135 °C       | 134 - 135 °C     |
| Residue on Ignition      | 0.010% max.      | <0.008%          |
| Trace Metals             | Arsenic (As)     | 5 ppm max.       |
|                          | Copper (Cu)      | 5 ppm max.       |
|                          | Iron (Fe)        | 5 ppm max.       |
|                          | Lead (Pb)        | 5 ppm max.       |

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: 16-000495

CAUTION STATEMENT: For manufacturing, processing, or repacking.

CAUTION STATEMENT: Rx only.

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INTENDED USE: Material represented by this Certificate of Analysis is suitable for use as a non-Sterile Active Pharmaceutical Ingredient 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 a Sterile or Injectable Active Pharmaceutical Ingredient, Drug Product or Household Item.

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.

Prepared by: Jaron Hughes Date: 1/22/21 Job Title: QA Specialist

Reviewed by: CW Date: 1/22/21 Job Title: QA Supervisor