DCN: 16-000062 v.5.2



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

| Effective Date: 4-Dec-2020 | 4-Dec-2023 : Date of Next Review |
|---|------------------------------------|
| Prepared By: Amy Hosein | 16-000062 v.5.1 : Supersedes |
| QA/QC Approval: Carissa McCollian | Wendy Santay : Management Approval |
| Reason for Revision: See Revision History in ensur. | |

CERTIFICATE OF ANALYSIS TRIS HC1

BIO EXCIPIENT GRADE / TH3220-K010

LOT: TH3220-146-0221

NH₂C(CH₂OH)₃ · HCl → F.W. 157.60 g/mol. → CAS# 1185-53-1

Manufacturing Date: 2/15/21 Retest Date: 2/28/23
Manufacturing Site: 1474 Rockdale Lane, Stroudsburg, PA 18360
Packaging Date: 3/1/21 Packaging Site: 100 Majestic Way, Bangor PA, 18013

| Anal | YSIS | SPECIFICATION | TEST RESULT |
|----------------------|----------------|------------------|---------------|
| Absorbance (1M) | 280 nm | 0.06 a.u. max. | <0.06 a.u. |
| Appearance and Color | | White / Crystals | Passes Test |
| Assay | | 99.5% min. | 99.6% |
| Enzymes | DNase | None Detected | None Detected |
| | RNase | None Detected | None Detected |
| | Protease | None Detected | None Detected |
| Heavy Metals | | 2 ppm max. | < 2 ppm |
| Identification (IR) | | Passes Test | Passes Test |
| Insoluble Matter | | 0.001% max. | <0.001% |
| Karl Fischer | | 0.5% max. | 0.3% |
| Melting Range | | 150 − 153 °C | 150-151°C |
| pH (0.5M) | | 4.0 - 5.0 | 4.3 @ 23.3°C |
| pK_a | | 8.0 - 8.4 | 8.2 |
| Residue on Ignition | L | 0.1% max. | <0.1% |
| Solubility 35% | | Passes Test | Passes Test |
| Trace Metals | Arsenic (As) | 1 ppm max. | <1 ppm |
| | Calcium (Ca) | 1 ppm max. | 1 ppm |
| | Copper (Cu) | 1 ppm max. | <1 ppm |
| | Iron (Fe) | 1 ppm max. | <1 ppm |
| | Lead (Pb) | 1 ppm max. | <1 ppm |
| | Magnesium (Mg) | 1 ppm max. | <1 ppm |

DCN: 16-000062 v.5.2

TEST METHOD REFERENCE: DCN: 16-000042

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: 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.

Date: 3/16/21 Job Title: QA Specialist

Date: 3/10/21 Job Title: QA Manager