DCN: 16-000062 v.5.0



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

| Effective Date: 10-Apr-2020 | 10-Apr-2023 : Date of Next Review |
|---|--------------------------------------|
| Prepared By: Kyle Snyder | 16-000062 v.4.1 : Supersedes |
| QA/QC Approval: Carissa McCollian | Hannah Bernier : Management Approval |
| Reason for Revision: See Revision History in ensur. | |

CERTIFICATE OF ANALYSIS TRIS HCl

BIO EXCIPIENT GRADE / TH3220-K010 LOT: TH3220-117-0420

NH₂C(CH₂OH)₃ HCl * F.W. 157.60 g/mol. * CAS# 1185-53-1 Manufacturing Date: 11/27/2019 Retest Date: 11/30/2021 Manufacturing Site: 1474 Rockdale Lane, Stroudsburg, PA 18360

Packaging Date: 4/28/2020 Packaging Site: 100 Majestic Way, Bangor PA, 18013

| Anal | YSIS | SPECIFICATION | TEST RESULT |
|----------------------|----------------|------------------|------------------|
| Absorbance (1M) | 280 nm | 0.06 a.u. max. | 0.01 a.u. |
| Appearance and Color | | White / Crystals | White / Crystals |
| Assay | | 99.5% min. | 99.9% |
| 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. | 0.2% |
| Melting Range | | 150 − 153 °C | 151-152°C |
| pH (0.5M) | | 4.0 - 5.0 | 4.2 @ 23.1°C |
| pK_a | | 8.0 - 8.4 | 8.2 |
| Residue on Ignition | ı | 0.1% max. | <0.1% |
| Solubility 35% | | Passes Test | Passes Test |
| Trace Metals | Arsenic (As) | 1 ppm max. | < 1 ppm |
| | Calcium (Ca) | l 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 |

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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.

Prepared by: Cov Date: 4/29/20 Job Title: OA Supervisor

Reviewed by: White Date: 04/30/20 Job Title: QA Manager