DCN: BSI-COA-0321 v.1.0



| Effective Date:      | 22-Jan-2025                            | 22-Jan-2028    | : Date of Next Review |
|----------------------|--|----------------|-----------------------|
| Prepared By:         | Amy Yencho                             | Not Applicable | : Supersedes          |
| QA/QC Approval:      | Krista Rehrig                          | Carissa Albert | : Management Approval |
| Reason for Revision: | See Revision History in MasterControl. |                |                       |

## CERTIFICATE OF ANALYSIS SODIUM DECANOATE BIOTECH, GMP BIO PHARMA GRADE / NDEC-4201

LOT: NDEC-G13-1124-0021

C<sub>10</sub>H<sub>19</sub>NaO<sub>2</sub> ^ F.W. 194.25 g/mol. ^ CAS# 1002-62-6 Manufacturing Date: 11/20/24 Retest Date: 11/30/26 Manufacturing Site: 100 Majestic Way, Bangor PA, 18013 Packaging Site: 100 Majestic Way, Bangor PA, 18013

| ANALYSIS            | SPECIFICATION                     | TEST RESULT                    |
|---------------------|-----------------------------------|--------------------------------|
| Appearance          | White to off-white powder         | White powder                   |
| Assay, Dried Basis  | 97.0 – 103.0 %                    | 100.7 %                        |
| Identification, IR  | Conforms to Reference<br>Standard | Conforms to Reference Standard |
| Loss on Drying      | ≤ 3.0 %                           | 2.6 %                          |
| pH (10%)            | 9.0 - 11.0                        | 10.1                           |
| Single Impurities   | ≤ 1.0 %                           | 0.2 %                          |
| Sodium              | Passes Test                       | Passes Test                    |
| Solubility in Water | Passes Test                       | Passes Test                    |
| Water, KF           | 1.5 – 3.0 %                       | 2.6 %                          |

COUNTRY OF ORIGIN: U.S.A.

TEST METHOD REFERENCE: DCN: BSI-ATM-0029

<u>INTENDED USE:</u> Material represented by this Certificate of Analysis is suitable for use as a process chemical. It is manufactured in accordance with the IPEC-PQG Joint Good Manufacturing Practice Guide. The material represented by this Certificate of Analysis is not suitable to be used as an Active Pharmaceutical Ingredient, Drug, Drug Product or Household Item.

| Prepared by: Land (h  | Date: 2/4/25 | Job Title: QA Technician          |
|-----------------------|--------------|-----------------------------------|
| Reviewed by: an Allut |              | Job Title: Senior Quality Manager |