## **CERTIFICATE OF ANALYSIS**

Product Name ACETONE

Grade Meets USP/ACS HPLC-UV GRADE Monographs

 Catalog #
 32900HPLC

 Lot #
 C21D05005

 Date of Manufacture:
 04/05/21

**Recommended Retest Date:** Five Years from Date of Manufacture

| TEST                                   | MONO<br>GRAPH    | SPECIFICATION                          | RESULT        |
|--|------------------|--|---------------|
| Assay (corrected for water)            | ACS              | NLT 99.5%                              | 99.57%        |
| Assay (on the anhydrous basis)         | NF               | NLT 99.0%                              | 99.96%        |
| Identification A - Infrared Absorption | NF               | Conforms to Infrared Spectra           | Pass          |
| Identification B - GC                  | NF               | Conforms to Reference Chromatogram     | Pass          |
| Specific Gravity @ 25°C                | NF               | NMT 0.789                              | 0.788         |
| Appearance                             | ACS <sup>+</sup> | Clear liquid with characteristic odor  | Pass          |
| Color (APHA)                           | ACS              | 10 max                                 | 1             |
| Solubility in Water                    | ACS <sup>+</sup> | The solution remains clear for 30 min. | Pass          |
| Residue After Evaporation              | ACS <sup>+</sup> | 0.001%, max                            | 0.000%        |
| Nonvolatile Residue                    | NF <sup>+</sup>  | NMT 2 mg/50mL (0.004%)                 | 0 mg/50mL     |
| Titrable Acid                          | ACS <sup>+</sup> | 0.0003 meq/g, max                      | 0.0001 meq/g  |
| Titrable Base                          | ACS <sup>+</sup> | 0.0006 meq/g, max                      | 0.0001 meq/g  |
| Aldehyde (as HCHO)                     | ACS <sup>+</sup> | 0.002%, max                            | < 0.002%      |
| Isopropyl Alcohol                      | ACS              | 0.05%, max                             | None Detected |
| Methanol                               | ACS              | 0.05%, max                             | 0.04%         |
| Substances Reducing Permanganate       | ACS <sup>+</sup> | To Pass Test                           | Pass          |
| Readily Oxidizable Substances          | NF <sup>+</sup>  | To Pass Test                           | Pass          |
| UV Absorbance                          | ACS/HPLC         | 400nm 0.01 max.                        | 0.00          |
|  |                  | 350nm 0.02 max.                        | 0.00          |
|  |                  | 340nm 0.10 max.                        | 0.04          |
|  |                  | 330nm 1.00 max.                        | 0.61          |



| TEST                            | MONO<br>GRAPH | SPECIFICATION | RESULT |
|---------------------------------|---------------|---------------|--------|
| Liquid Chromatography           | ACS/HPLC      | To Pass Test  | Pass   |
| Absorbance                      | ACS/HPLC      | To Pass Test  | Pass   |
| Gradient Elution                | ACS/HPLC      | To Pass Test  | Pass   |
| Gradient Analysis at 254nm, Max | ACS/HPLC      | To Pass Test  | Pass   |
| Water                           | ACS/NF        | 0.5%, max     | 0.38%  |

<sup>&</sup>lt;sup>†</sup>This test is performed quarterly

## **Certification and Compliance Statements**

This lot of Acetone complies with all of the current requirements listed in the United States Pharmacopeia, American Chemical Society, National Formulary monographs, and meets the requirements for General Use HPLC Grade. This product is not intended for GC or critical HPLC analysis. See Distilled Grade for those Applications.

No chemicals whatsoever are used as solvents at any point in the manufacture, processing or packaging of Acetone. Only Class 2 and Class 3 residual solvents may appear as impurities / related substances / low level contaminants in Acetone. Concentration of Class 2 Option 1 and Class 3 residual solvents is below limits in the current USP/NF General Chapter <467> and ICH Q3C Impurities: Residual Solvents.

This product is not derived, nor does it come in contact with, any materials derived from bovine or other animal sources.

This product is for further commercial manufacturing, laboratory or research use, and may be used as an excipient or a process solvent for pharmaceutical purposes. It is not intended for use as an active ingredient in drug manufacturing nor as a medical device or disinfectant. Appropriate/legal use of this product is the responsibility of the user.

Approved by: D. Simoncelli, Quality Control Chemist

Derh Sint

Date of Approval: 04/05/21