h2/13/25, 1:56-PM/trungtamthuoc.com/SP-NF Acetohydroxamic Acid

Status: Currently Official on 13-Feb-2025
Official Date: Official as of 01-May-2020
Document Type: USP Monographs
DocId: GUID-E6D72F31-782F-4984-A615-A9A161B49957_4_en-US
DOI: https://doi.org/10.31003/USPNF_M540_04_01
DOI Ref: u3fwb

© 2025 USPC Do not distribute

Acetohydroxamic Acid

C2H5NO2

75.07

N-Acetyl hydroxyacetamide;

Acetohydroxamic acid CAS RN[®]: 546-88-3; UNII: 4RZ82L2GY5.

DEFINITION

Acetohydroxamic Acid, dried over phosphorus pentoxide for 16 h, contains NLT 98.0% and NMT 101.0% of acetohydroxamic acid (C₂H_ENO₂).

IDENTIFICATION

Change to read:

• A. Spectroscopic Identification Tests (197), Infrared Spectroscopy: 197K (CN 1-May-2020)

• B.

Sample solution: 20 mg/mL in water

Analysis: To 10 mL of the Sample solution add 2 drops of potassium permanganate TS.

Acceptance criteria: The pink color of the permanganate disappears.

ASSAY

• PROCEDURE

Ferric chloride solution: 20 mg/mL of ferric chloride in 0.1 N hydrochloric acid

Standard solution: $500 \mu g/mL$ of <u>USP Acetohydroxamic Acid RS</u> in 0.1 N hydrochloric acid Sample solution: $500 \mu g/mL$ of Acetohydroxamic Acid, previously dried, in 0.1 N hydrochloric acid

Blank: 0.1 N hydrochloric acid

Analysis

Samples: Standard solutions, Sample solution, and Blank

Transfer 10.0 mL each of the Standard solution, Sample solution, and Blank to separate 100-mL volumetric flasks. To each flask add 50 mL of 0.1 N hydrochloric acid and 10.0 mL of Ferric chloride solution, and dilute with 0.1 N hydrochloric acid to volume. Without delay, concomitantly determine the absorbances of the solutions at the wavelength of maximum absorbance at about 502 nm using the Blank to set the instrument.

Calculate the percentage of acetohydroxamic acid (C₂H₅NO₂) in the portion of Acetohydroxamic Acid taken:

Result =
$$(A_{II}/A_s) \times (C_s/C_{II}) \times 100$$

A,, = absorbance of the Sample solution

A = absorbance of the Standard solution

C_s = concentration of <u>USP Acetohydroxamic Acid RS</u> in the Standard solution (μg/mL)

 C_{II} = concentration of Acetohydroxamic Acid in the Sample solution (µg/mL)

Acceptance criteria: 98.0%-101.0% on the previously dried basis

IMPURITIES

- Residue on Ignition (281): NMT 0.1%
- LIMIT OF HYDROXYLAMINE

Buffer: 1.36 g/L of monobasic potassium phosphate in water, adjusted with 1 M potassium hydroxide to a pH of 7.4

Solution A: 1 mg/mL of pyridoxal 5-phosphate monohydrate in Buffer, prepared in a low-actinic flask fresh before use

Standard stock solution: 2.0 mg/mL of hydroxylamine hydrochloride in water

Standard solutions: Transfer 5.0, 10.0, and 15.0 mL of the *Standard stock solution* to separate 100-mL volumetric flasks, and dilute with water to volume.

2/13/25, 1:56 PM/+run at a mthu a contact Acid

Sample solution: Transfer 1500 mg of Acetohydroxamic Acid, previously dried, to a 100-mL beaker, and dissolve in a sufficient amount of water to cover the electrode of a calibrated pH meter (about 60 mL). While stirring, adjust with 0.05 M potassium hydroxide to a pH of 7.4. Transfer the contents of the beaker, with the aid of small portions of water, to a 100-mL volumetric flask, and dilute with water to volume.

Blank: Water Analysis

Samples: Standard solutions, Sample solution, and Blank

Transfer 2.0 mL of each *Standard solution*, the *Sample solution*, and *Blank* into separate 100-mL volumetric flasks. To each flask add 4.0 mL of *Solution A*. After 8 min, accurately timed, dilute the contents of each flask with *Buffer* to volume.

Immediately determine the fluorescence intensities of the solutions from the *Standard solutions* and the *Sample solution* in a fluorometer at an excitation wavelength of 350 nm and an emission wavelength of 450 nm, setting the instrument to zero with the *Blank*. Determine the best-fit straight line from the fluorescence intensities of the three *Standard solutions* versus the hydroxylamine hydrochloride concentrations, in µg/mL. From the best-fit straight line, determine the concentration, in µg/mL, of hydroxylamine hydrochloride in the *Sample solution*.

Calculate the percentage of hydroxylamine in the portion of Acetohydroxamic Acid taken:

Result =
$$(C_{11}/C) \times (M_{r1}/M_{r2}) \times 100$$

 C_{ij} = concentration of hydroxylamine hydrochloride in the Sample solution (mg/mL)

C = concentration of Acetohydroxamic Acid in the Sample solution (mg/mL)

 M_{c1} = molecular weight of hydroxylamine, 33.03

 M_{c2} = molecular weight of hydroxylamine hydrochloride, 69.50

Acceptance criteria: NMT 0.5%

SPECIFIC TESTS

• Loss on Drying (731)

Analysis: Dry a sample over phosphorus pentoxide for 16 h.

Acceptance criteria: NMT 1.0%

• COMPLETENESS OF SOLUTION (641): A 1.0-g portion dissolves in 10 mL of water to yield a clear solution.

Color of Solution

Sample solution: 200 mg/mL in water

Blank: Water

Instrumental conditions

(See <u>Ultraviolet-Visible Spectroscopy (857)</u>.)

Mode: UV-Vis

Analytical wavelength: Between 400 and 750 nm

Cell: 1 cm

Acceptance criteria: The absorbance is NMT 0.050.

ADDITIONAL REQUIREMENTS

• Packaging and Storage: Preserve in tight containers, and store in a cool, dry place.

USP REFERENCE STANDARDS (11)

USP Acetohydroxamic Acid RS

Auxiliary Information - Please check for your question in the FAQs before contacting USP.

Topic/Question	Contact	Expert Committee
ACETOHYDROXAMIC ACID	Documentary Standards Support	SM32020 Small Molecules 3

Chromatographic Database Information: Chromatographic Database

Most Recently Appeared In:

Pharmacopeial Forum: Volume No. PF 30(5)

Current DocID: GUID-E6D72F31-782F-4984-A615-A9A161B49957_4_en-US

DOI: https://doi.org/10.31003/USPNF_M540_04_01

DOI ref: u3fwb