Calcium Citrate

570.49 $C_{12}H_{10}Ca_3O_{14} \cdot 4H_2O$ 1,2,3-Propanetricarboxylic acid, 2-hydroxy-, calcium salt (2:3), tetrahydrate;

Calcium citrate (3:2), tetrahydrate [5785-44-4].

Calcium Citrate contains four molecules of water of hydration. When dried at 150 ° to constant weight, it contains NL T 97.5% and NMT 100.5% of Ca $_{3}(C_{6}H_{5}O_{7})_{2}$.

IDENTIFICATION

Analysis: Dissolve 0.5 g in a mixture of 10 mL of water and 2.5 mL of 2 N nitric acid. Add 1 mL of mer curic sulfate TS, heat to boiling, and add 1 mL of potassium permanganate

Acceptance criteria: A white precipitate is formed.

Sample: 0.5 g of Calcium Citrate

Analysis: Ignite completely the Sample at as low a temperature as possible, cool, and dissolve the residue in dilute glacial acetic acid (1:10). Filter, and add 10 mL of ammonium oxalate TS to the filtrate.

Acceptance criteria: A voluminous white precipitate that is soluble in hydrochloric acid is formed.

PROCEDURE

Sample solution: Dissolve 350 mg of Calcium Citrate, previously dried at 150° to constant weight, in 12 mL of 0.5 M hydrochloric acid, and dilute with water to about 100 mL. Analysis: While stirring the Sample solution, add 30 mL of 0.05 M edetate disodium VS from a 50-mL buret. Add 15 mL of 1 N sodium hydroxide and 300 mg of hydroxy naphthol blue, and continue the titration to a blue endpoint. Each mL of 0.05 M edetate disodium is equivalent to 8.307 mg of calcium citrate ($Ca_3(C_6H_5O_7)_2$). Acceptance criteria: 97.5%–100.5% on the dried basis

IMPURITIES

ARSENIC, Method I (211)

Test preparation: Dissolve 1 g of Calcium Citrate in 5 mL of 3 N hydrochloric acid, and dilute with water to 35 mL. Acceptance criteria: NMT 3 ppm

HEAVY METALS, Method I (231)

Test preparation: Dissolve 1 g of Calcium Citrate in a mixture of hydrochloric acid and water (2:20). Add 1.5 mL of ammonium hydroxide, and dilute with water to 25 mL.

Acceptance criteria: NMT 20 ppm

LEAD (251)

Test preparation: Dissolve 0.5 g of Calcium Citrate in 20 mL of 3 N hydrochloric acid. Evaporate this solution on a steam bath to 10 mL, dilute with water to 20 mL, and cool. Use 5 mL of Diluted Standard Lead Solution (5 µg of Pb) for the test.

Acceptance criteria: NMT 10 ppm

LIMIT OF FLUORIDE

[NOTE—Prepare and store all solutions in plastic containers.] Standard stock solution: $1000 \ \mu g/mL$ of fluoride ion from USP Sodium Fluoride RS in water

Standard solution: 5 µg/mL of fluoride ion from Standard stock solution. [NOTE—Prepare on the day of use.]

Linearity solution A: Transfer 1.0 mL of the Standard solu-

tion to a 250-mL plastic beaker. Add 50 mL of water, 5 mL of 1 N hydrochloric acid, 10 mL of 1.0 M sodium citrate,

and 10 mL of 0.2 M edetate disodium. If necessar y, adjust with 1 N sodium hydroxide or 1 N hydrochloric acid to a pH of 5.5. Transfer to a 100-mL volumetric flask, and dilute with water to volume. This solution contains 0.05 µg/mL of fluoride.

Linearity solution B: Transfer 5.0 mL of the Standard solution to a 250-mL plastic beaker, and proceed as directed for Linearity solution A beginning with "Add 50 mL of water,".

This solution contains 0.25 µg/mL of fluoride. Linearity solution C: Transfer 10.0 mL of the Standard solution to a 250-mL plastic beaker, and proceed as directed for Linearity solution A beginning with "Add 50 mL of water,". This solution contains 0.50 μ g/mL of fluoride.

Sample solution: Transfer 1.0 g of Calcium Citrate to a 100mL beaker. Add 10 mL of water and, while stirring, 10 mL of 1 N hydrochloric acid. When dissolved, boil rapidly for 1 min, transfer the solution to a 250-mL plastic beaker, and cool in ice water. Add 15 mL of 1.0 M sodium citrate and 10 mL of 0.2 M edetate disodium, and adjust with 1 N sodium hydroxide or 1 N hydrochloric acid to a pH of 5.5. Transfer this solution to a 100-mL volumetric flask, and dilute with water to volume.

Electrode system: Use a fluoride-specific, ion-indicating electrode and a silver-silver chloride reference electrode connected to a pH meter capable of measuring potentials with a minimum reproducibility of ± 0.2 mV (see pH $\langle 791 \rangle$).

Analysis

Samples: Linearity solution A, Linearity solution B, Linearity solution C, and Sample solution

Transfer 50 mL of each Linearity solution A, Linearity solution B, and Linearity solution C to separate 250-mL plastic beakers, and measure the potential of each solution with the Electrode system. Between each reading wash the electrodes with water, and absorb any residual water by blotting the electrodes dry. Plot the logarithms of the fluoride concentrations (0.05, 0.25, and 0.50 µg/mL, respectively) versus potential to obtain a Standard response line.

Transfer 50 mL of the Sample solution to a 250-mL plastic beaker, and measure the potential with the Electrode system. From the measured potential and the Standard response line determine the concentration, C, in μg/mL, of fluoride ion in the Sample solution. Calculate the percentage of fluoride in the specimen taken by multiplying C by 0.01.

Acceptance criteria: NMT 0.003%

• LIMIT OF ACID-INSOLUBLE SUBSTANCES

Sample solution: Dissolve 5 g of Calcium Citrate by heating with a mixture of hydrochloric acid and water (10:50) for 30

Analysis: Filter, wash, and dr y at 105° for 2 h the residue so obtained.

Acceptance criteria: The weight of the residue is NMT 10 mg (0.2%).

SPECIFIC TESTS

• Loss on Drying (731): Dry a sample at 150 ° for 4 h: it loses from 10.0% to 13.3% of its weight.

ADDITIONAL REQUIREMENTS

- PACKAGING AND STORAGE: Preserve in well-closed containers.
- USP REFERENCE STANDARDS (11) USP Sodium Fluoride RS