

layer from the *Standard solution* and *Sample solution* against the aqueous layer from the *Blank*.

Calculate the percentage of citric acid in the portion of Monoglyceride Citrate taken:

$$\text{Result} = (A_U/A_S) \times (V \times C_S/W) \times 100$$

- $A_U$  = absorbance of the *Sample solution*  
 $A_S$  = absorbance of the *Standard solution*  
 $V$  = volume of the *Sample solution* (mL)  
 $C_S$  = concentration of USP Citric Acid RS in the *Standard solution* (mg/mL)  
 $W$  = weight of Monoglyceride Citrate taken to prepare the *Sample solution* (mg)

Acceptance criteria: 14.0%–17.0% on the anhydrous basis

#### IMPURITIES

- **RESIDUE ON IGNITION** (281): NMT 0.3%, determined on 1 g
- **HEAVY METALS**, *Method II* (231): NMT 10 ppm

#### SPECIFIC TESTS

- **FATS AND FIXED OILS**, *Acid Value* (401): 70–100
- **FATS AND FIXED OILS**, *Saponification Value* (401): 260–265
- **WATER DETERMINATION**, *Method I* (921): NMT 0.2%

#### ADDITIONAL REQUIREMENTS

- **PACKAGING AND STORAGE**: Preserve in well-closed containers. No storage requirements specified.
- **USP REFERENCE STANDARDS** (11)  
USP Citric Acid RS

## Monosodium Glutamate

#### DEFINITION

Monosodium Glutamate contains NLT 99.0% and NMT 100.5% of  $C_5H_8NNaO_4 \cdot H_2O$ .

#### IDENTIFICATION

- **A**.  
**Sample solution**: 1 in 30  
**Analysis**: To 1 mL of the *Sample solution* add 1 mL of ninhydrin TS and 100 mg of sodium acetate, and heat in a boiling water bath for 10 min.  
**Acceptance criteria**: An intense, violet blue color is formed.
- **B**.  
**Sample solution**: 1 in 10  
**Analysis**: To 10 mL of the *Sample solution* add 5.6 mL of 1 N hydrochloric acid.  
**Acceptance criteria**: A white, crystalline precipitate of glutamic acid is formed on standing. Precipitation is promoted by agitation. When 6 mL of 1 N hydrochloric acid is added to the turbid solution, the glutamic acid dissolves on stirring.
- **C. IDENTIFICATION TESTS—GENERAL**, *Sodium* (191): It meets the requirements of the pyroantimonate precipitate test.

#### ASSAY

##### PROCEDURE

**Sample**: 250 mg  
**Titrimetric system**  
 (See *Titrimetry* (541).)  
**Mode**: Direct titration  
**Titrant**: 0.1 N perchloric acid VS  
**Blank**: 100 mL of glacial acetic acid with a few drops of water  
**Endpoint detection**: Potentiometric  
**Analysis**: Wet the *Sample* with a few drops of water. Dissolve in 100 mL of glacial acetic acid. Titrate with 0.1 N perchloric acid VS. Perform a blank determination. Calculate the percentage of monosodium glutamate ( $C_5H_8NNaO_4 \cdot H_2O$ ) in the *Sample* taken:

$$\text{Result} = [(V - B) \times N \times F \times 100]/W$$

- $V$  = volume of *Titrant* consumed by the *Sample* (mL)  
 $B$  = volume of *Titrant* consumed by the *Blank* (mL)  
 $N$  = actual normality of the *Titrant* (mEq/mL)  
 $F$  = equivalency factor, 93.56 mg/mEq  
 $W$  = weight of the *Sample* (mg)

Acceptance criteria: 99.0%–100.5%

#### IMPURITIES

- **CHLORIDE AND SULFATE**, *Chloride* (221): A 280-mg portion shows no more chloride than corresponds to 1.0 mL of 0.020 N hydrochloric acid (0.25%).
- **LEAD** (251): NMT 10 ppm
- **HEAVY METALS**, *Method II* (231): NMT 20 ppm

#### SPECIFIC TESTS

##### CLARITY AND COLOR OF SOLUTION

**Sample solution**: 1.0 g in 10 mL of water

**Standard solution**: To 0.2 mL of a solution of sodium chloride containing 10 µg/mL of chloride ion (Cl), add 20 mL of water, and mix. Then add 1 mL of 5 N nitric acid, 0.2 mL of dextrin solution (1 in 50), and 1 mL of silver nitrate TS, and allow to stand for 15 min.

**Analysis**: Compare the *Sample solution* with the *Standard solution* (see *Spectrophotometry and Light-Scattering—Visual Comparison* (851)).

**Acceptance criteria**: The *Sample solution* is colorless and has no more turbidity than the *Standard solution*.

##### OPTICAL ROTATION, *Specific Rotation* (781S)

**Sample solution**: 100 mg/mL in 2 N hydrochloric acid

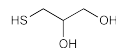
**Acceptance criteria**: +24.8° to +25.3°, determined at 20°

- **pH** (791): 6.7–7.2, in a solution (1 in 20)
- **LOSS ON DRYING** (731): Dry a sample at 100° for 5 h: it loses NMT 0.5% of its weight.

#### ADDITIONAL REQUIREMENTS

- **PACKAGING AND STORAGE**: Preserve in tight containers.

## Monothioglycerol



$C_3H_8O_2S$  108.16  
 1,2-Propanediol, 3-mercapto-.  
 3-Mercapto-1,2-propanediol [96-27-5].

» Monothioglycerol contains not less than 97.0 percent and not more than 101.0 percent of  $C_3H_8O_2S$ , calculated on the anhydrous basis.

**Packaging and storage**—Preserve in tight containers.

**Specific gravity** (841): between 1.241 and 1.250.

**Refractive index** (831): between 1.521 and 1.526.

**pH** (791): between 3.5 and 7.0, in a solution (1 in 10).

**Water**, *Method II* (921): not more than 5.0%.

**Residue on ignition** (281): not more than 0.1%.

**Selenium** (291): 0.003%, a 200-µL test specimen being used.

**Heavy metals**, *Method II* (231): 0.002%.

**Assay**—Dissolve about 400 mg of Monothioglycerol, accurately weighed, in 50 mL of water, and titrate with 0.1 N iodine VS, adding 3 mL of starch TS as the endpoint is approached. Each mL of 0.1 N iodine is equivalent to 10.82 mg of  $C_3H_8O_2S$ .