# SODIUM ERYTHORBATE

Prepared at the 17th JECFA (1973), published in FNP 4 (1978) and in FNP 52 (1992). Metals and arsenic specifications revised at the 61st JECFA (2003). An ADI 'not specified' was established at the 37th JECFA (1990)

SYNONYMS Sodium isoascorbate

## DEFINITION

Chemical names Sodium isoascorbate, sodium D-isoascorbic acid, sodium salt of 2,3didehydro-D-erythro-hexono-1,4-lactone, 3-keto-D-gulofurano-lactone sodium enolate monohydrate

C.A.S. number 6381-77-7

Chemical formula

Structural formula



Formula weight 216.13

Assay Not less than 98% after drying

**DESCRIPTION** White, almost odourless crystalline powder

 $C_6H_7O_6Na \cdot H_2O$ 

### FUNCTIONAL USES Antioxidant

### CHARACTERISTICS

IDENTIFICATION

<u>Solubility</u> (Vol. 4)	Freely soluble in water, very slightly soluble in ethanol	
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<u>Reducing activity</u> A solution of the sample will decolourize a solution of 2,6-dichlorophenolindophenol TS

Test for ascorbate Passes test (Vol. 4)

<u>Test for sodium</u> (Vol. 4) Passes test Test a solution of previously ignited sample, acidified with dilute acetic acid TS, filtered if necessary

#### PURITY

Loss on drying (Vol. 4) Not more than 0.25% (in vacuum over sulfuric acid, 24 h)

Specific rotation (Vol. 4) [alpha] 25, D: Between +95.5° and +98.0° (10% (w/v) solution)

<u>pH</u> (Vol. 4) 5.5 - 8.0 (1 in 10 soln)

<u>Oxalate</u> To a solution of 1 g in 10 ml of water add 2 drops of glacial acetic acid and 5 ml of 10% calcium acetate solution. The solution should remain clear.

Lead (Vol. 4) Not more than 2 mg/kg Determine using an atomic absorption technique appropriate to the specified level. The selection of sample size and method of sample preparation may be based on the principles of the method described in Volume 4, "Instrumental Methods."

METHOD OF<br/>ASSAYDissolve about 0.400 g of the dried sample in a mixture of 100 ml of carbon<br/>dioxide-free water and 25 ml of dilute sulfuric acid TS. Titrate the solution at<br/>once with 0.1 N iodine, adding a few drops of starch TS as indicator as the<br/>end point is approached. Each ml of 0.1 N iodine is equivalent to 10.807 mg<br/>of  $C_6H_7O_6Na \cdot H_2O$ .