

# SODIUM DL-MALATE

*Prepared at the 67<sup>th</sup> JECFA (2006) and published in FAO JECFA Monographs 3 (2006), superseding specifications prepared at the 30<sup>th</sup> JECFA (1986) and published in FNP 52 (1992) and in the Combined Compendium of Food Additive specifications, FAO JECFA monographs 1 (2005). Metals and arsenic specifications were revised at the 59<sup>th</sup> JECFA (2002). An ADI 'not specified' was established at the 23<sup>rd</sup> JECFA (1979).*

## SYNONYMS

Malic acid sodium salt; INS No. 350(ii)

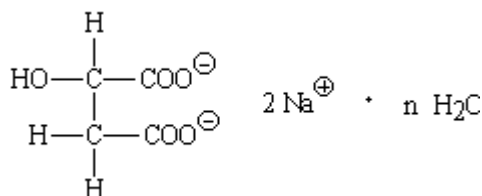
## DEFINITION

Chemical names Disodium DL-malate, hydroxybutanedioic acid disodium salt

C.A.S. number 676-46-0

Chemical formula Hemihydrate:  $C_4H_4Na_2O_5 \cdot 1/2 H_2O$   
Trihydrate:  $C_4H_4Na_2O_5 \cdot 3 H_2O$

Structural formula



Formula weight Hemihydrate: 187.1  
Trihydrate: 232.1

Assay Not less than 98% and not more than 102% on the dried basis

**DESCRIPTION** Odourless white crystalline powder or lumps

**FUNCTIONAL USES** Acidity regulator

## CHARACTERISTICS

### IDENTIFICATION

Solubility (Vol. 4) Freely soluble in water

Test for sodium (Vol. 4) Passes test

Test for malate (Vol. 4) Passes test  
Test 5 ml of a 1 in 20 solution of the sample

## PURITY

Loss on drying (Vol. 4) Hemihydrate: Not more than 7% (130°, 4 h)  
Trihydrate: 20.5% - 23.5% (130°, 4 h)

Alkalinity Not more than 0.2% as Na<sub>2</sub>CO<sub>3</sub>  
Dissolve 1 g of the sample in 20 ml of freshly boiled and cooled water, and add 2 drops of phenolphthalein TS. If a pink colour is produced, add 0.4 ml of 0.1 N sulfuric acid. The colour of the solution disappears.

Fumaric and maleic acid (Vol. 4) Not more than 1.0% of fumaric acid and not more than 0.05% of maleic acid

Lead (Vol. 4) Not more than 2 mg/kg  
Determine using an AAS/ICP-AES technique appropriate to the specified level. The selection of sample size and method of sample preparation may be based on the principles of the methods described in Volume 4.

**METHOD OF ASSAY** Dissolve about 0.25 g of the dried sample, accurately weighed, in 50 ml of glacial acetic acid, and titrate with 0.1 N perchloric acid, determining the endpoint potentiometrically. Each ml of 0.1 N perchloric acid is equivalent to 8.903 mg of C<sub>4</sub>H<sub>4</sub>Na<sub>2</sub>O<sub>5</sub>.