## International Standard



753/4

INTERNATIONAL ORGANIZATION FOR STANDARDIZATION•МЕЖДУНАРОДНАЯ ОРГАНИЗАЦИЯ ПО СТАНДАРТИЗАЦИИ•ORGANISATION INTERNATIONALE DE NORMALISATION

# Acetic acid for industrial use — Methods of test — Part 4: Determination of acetaldehyde monomer content — Titrimetric method

Acide acétique à usage industriel — Méthodes d'essai — Partie 4 : Dosage de l'acétaldéhyde monomère — Méthode titrimétrique

First edition - 1981-10-15

UDC 661.731:543

Ref. No. ISO 753/4-1981 (E)

Descriptors: industrial products, acetic acid, tests, determination of content, acetaldehyde, titration.

### **Foreword**

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Draft International Standards adopted by the technical committees are circulated to the member bodies for approval before their acceptance as International Standards by the ISO Council.

International Standard ISO 753/4 was developed by Technical Committee ISO/TC 47, *Chemistry*, and was circulated to the member bodies in March 1980.

It has been approved by the member bodies of the following countries:

Australia France Poland Austria Germany, F. R. Romania

Belgium Hungary South Africa, Rep. of Brazil India Switzerland

China Italy Thailand

Czechoslovakia Korea, Rep. of United Kingdom Egypt, Arab Rep. of Netherlands USSR

No member body expressed disapproval of the document.

This International Standard has also been approved by the International Union of Pure and Applied Chemistry (IUPAC).

International Standards ISO 753/1 to ISO 753/11 cancel and replace ISO Recommendation R 753-1968, of which they constitute a technical revision.

# Acetic acid for industrial use — Methods of test — Part 4: Determination of acetaldehyde monomer content — Titrimetric method

### 1 Scope and field of application

This part of ISO 753 specifies a titrimetric method for the determination of the acetaldehyde monomer content of acetic acid for industrial use.

The method is applicable to products having acetaldehyde monomer contents equal to or greater than 0.01 % (m/m).

This document should be read in conjunction with ISO 753/1 (see the annex).

 $\mbox{NOTE}-\mbox{For the determination of the total acetaldehyde content, including polymers, see ISO 753/5.}$ 

#### 2 Reference

ISO/R 385, Burettes.

#### 3 Principle

Reaction of the acetaldehyde present in a test portion with an excess of sodium hydrogen sulphite solution, and iodometric titration of the residual sodium hydrogen sulphite.

#### 4 Reaction

CH<sub>3</sub>CHO + NaHSO<sub>3</sub> 
$$\rightarrow$$
 CH<sub>3</sub>  $-$  C  $-$  OH  $|$  SO<sub>3</sub>Na

#### 5 Reagents

During the analysis, use only reagents of recognized analytical grade and only distilled water or water of equivalent purity.

**5.1 Sodium hydrogen sulphite**, approximately 18,2 g/l solution.

Dissolve 1,66 g of disodium disulphite ( $Na_2S_2O_5$ ) in water and dilute to 100 ml.

Prepare this solution at the time of use.

- **5.2 lodine,** standard volumetric solution,  $c(1/2 I_2) = 0.02 \text{ mol/I}.$
- **5.3 Sodium thiosulphate**, standard volumetric solution,  $c(Na_2S_2O_3) = 0.02 \text{ mol/l}.$

#### 5.4 Starch solution.

Triturate 1,0 g of soluble starch with 5 ml of water and, while stirring, pour the mixture into 100 ml of boiling water. Boil for a few minutes and allow to cool.

Discard the solution after 2 weeks.

#### 6 Apparatus

Ordinary laboratory apparatus and

- 6.1 Weighing pipette, of capacity 20 ml.
- **6.2** Two conical flasks, of capacity 250 ml, fitted with ground glass stoppers.
- **6.3** Burettes, of capacity 10 ml, complying with the requirements of ISO/R 385, class A.
- 6.4 Ice-water bath.

#### 7 Procedure

#### 7.1 Test portion

Using the weighing pipette (6.1), weigh, to the nearest 0,01 g, about 10 ml of the laboratory sample and transfer to a 50 ml one-mark volumetric flask containing 10 ml of water.

#### 7.2 Blank test

Carry out a blank test at the same time as the determination, following the same procedure and using the same quantities of all the reagents [except the sodium thiosulphate solution (5.3)] as used for the determination, but omitting the test portion.