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*sec*Butyl alcohol, methyl ethyl ketone, *isobutyl* methyl ketone, *isoamyl* ethyl ketone, diacetone alcohol, and hexylene glycol for industrial use – Determination of acidity to phenolphthalein – Volumetric method

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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.

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No Member Body expressed disapproval of the document.

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1 SCOPE AND FIELD OF APPLICATION

This International Standard specifies a method for the determination of the acidity to phenolphthalein of *sec*butyl alcohol (butan-2-ol), methyl ethyl ketone (butan-2-one), *isobutyl* methyl ketone (4-methylpentan-2-one), *isoamyl* ethyl ketone (5-methylheptan-3-one), diacetone alcohol (4-hydroxy-4-methylpentan-2-one) and hexylene glycol (2-methylpentane-2,4-diol) for industrial use.

2 REFERENCES

ISO/R 758, *Method for the determination of density of liquids at 20 °C.*

ISO ..., *Chemical products for industrial use – Sampling.*¹⁾

3 PRINCIPLE

Dilution of a test portion with water (or with ethanol or propan-2-ol if the sample is not completely soluble in water) and removal of carbon dioxide by passing a stream of nitrogen through the solution.

Titration of the acidity with a standard volumetric sodium hydroxide solution, using phenolphthalein as indicator.

4 REAGENTS

Distilled water, or water of equivalent purity, shall be used in the test.

4.1 Ethanol, 95 % (V/V) or **propan-2-ol**, 99 % (V/V).

4.2 Nitrogen, free from ammonia and carbon dioxide, supplied from a cylinder fitted with a pressure regulator enabling the flow rate to be controlled at 500 ± 50 ml/min.

4.3 Sodium hydroxide, 0,100 M standard volumetric solution.

4.4 Phenolphthalein, 5 g/l ethanolic solution.

Dissolve 0,5 g of phenolphthalein in 100 ml of 95 % (V/V) ethanol and make faintly pink by the addition of dilute sodium hydroxide solution.

5 APPARATUS

Ordinary laboratory apparatus and

5.1 Conical flask, capacity 500 ml, of borosilicate glass.

5.2 Burette, capacity 10 ml, graduated in 0,02 ml divisions.

6 SAMPLING

Follow the principles described in ISO ... Attention is drawn to the following recommendation: place the laboratory sample representative of the material taken from the bulk in a clean, dry, dark coloured glass-stoppered bottle of such a size that it is nearly filled by the sample.

If it is necessary to seal this bottle, care shall be taken to avoid the risk of contamination.

7 PROCEDURE

7.1 Test portion

Take 100,0 ml of the laboratory sample at 20 °C.

7.2 Determination

Place 100 ml of water (in the case of diacetone alcohol or hexylene glycol) or 100 ml of the ethanol or the propan-2-ol (4.1) (in the case of the other products) in the flask (5.1), and pass a stream of the nitrogen (4.2), at a rate of 500 ± 50 ml/min for 10 to 15 min. Add 0,5 ml of the phenolphthalein solution (4.4) and titrate with the standard volumetric sodium hydroxide solution (4.3) until the appearance of a pale pink colour.

Add the test portion (7.1) and pass a stream of the nitrogen (4.2), at a rate of 500 ± 50 ml/min for 10 to 15 min. Add 0,5 ml of the phenolphthalein solution (4.4) and titrate with the standard volumetric sodium hydroxide solution (4.3) until the appearance of a pale pink colour.

1) In preparation.

8 EXPRESSION OF RESULTS

The acidity, expressed as equivalents per kilogram is given by the formula

$$\frac{V}{1\,000\ \rho}$$

and, as a percentage by mass of acetic acid (CH₃COOH) by the formula

$$\frac{0,006\ 0 \times V}{100 \times \rho} \times 100 = \frac{0,006\ 0 \times V}{\rho}$$

where

V is the volume, in millilitres, of the standard volumetric sodium hydroxide solution (4.3) used for the titration;

ρ is the density, in grams per millilitre, of the sample at 20 °C, determined by the method described in ISO/R 758;

0,006 0 is the mass, in grams, of acetic acid corresponding to 1 ml of 0,100 M sodium hydroxide solution.

9 TEST REPORT

The test report shall include the following particulars :

- a) the reference of the method used;
- b) the results and the method of expression used;
- c) any unusual features noted during the determination;
- d) any operation not included in this International Standard or the documents to which reference is made, or regarded as optional.

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