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Surface active agents — Sulfated ethoxylated alcohols and alkylphenols — Determination of content of unsulfated matter

Agents de surface — Sulfates d'alcools et d'alkylphénols éthoxylés — Détermination de la teneur en matière insulfatée

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Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

ISO 8799 was prepared by the European Committee for Standardization (CEN) Technical Committee CEN/TC 276, *Surface active agents*, in collaboration with ISO Technical Committee ISO/TC 91, *Surface active agents*, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

This second edition cancels and replaces the first edition (ISO 8799:1988), which has been technically revised.

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Surface active agents — Sulfated ethoxylated alcohols and alkylphenols — Determination of content of unsulfated matter

1 Scope

This International Standard specifies a method for the determination of the content of unsulfated matter present in ordinary commercial neutralized products of sulfation of ethoxylated alcohols or alkylphenols [alkyl oxyethylene sulfates (ethoxylated alcohol sulfates) or alkylphenol oxyethylene sulfates (ethoxylated alkylphenol sulfates)] containing an average of not more than 20 oxyethylene groups per molecule.

2 Normative references

The following referenced documents are indispensable for the application of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO 607, Surface active agents and detergents — Methods of sample division

ISO 2271, Surface active agents — Detergents — Determination of anionic-active matter by manual or mechanical direct two-phase titration procedure

ISO 3696, Water for analytical laboratory use — Specification and test methods

EN 14480, Surface active agents — Determination of anionic surface active agents — Potentiometric two-phase titration method (Standards/ISO/0301217c-10be-4bb7-8db4-cc5baa42cf8/ISO-8799-2009

3 Principle

From a methanolic solution of the test portion, the unsulfated matter is separated on an ion-exchange column (filled with a mixture of cation-exchange resins and anion-exchange resins).

The unsulfated matter is recovered from the eluate by evaporation and weighing of the residue.

4 Reagents and materials

4.1 General.

WARNING — The procedures described in this International Standard involve the use of hazardous substances. The necessary precautions, as described in regulations covering the handling of hazardous substances, should be taken. Technical, organizational and personal protection measures should be observed.

During the analysis, unless otherwise specified, use only reagents of recognized analytical grade that have been checked in advance so as to not interfere with the analytical results, and water of grade 1 as defined in ISO 3696.

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- **4.2 Methanol**, CH₃OH, (CAS number: 67-56-1).
- **4.3 Hydrochloric acid solution**, c(HCI), about 1 mol/l, (CAS number: 7647-01-0).
- **4.4** Sodium hydroxide solution, c(NaOH), about 2 mol/l, (CAS number 1310-73-2).
- **4.5 Cation-exchange resin**, polystyrene sulfonic acid type, 2 % to 3 % crosslinked, 150 μ m to 330 μ m, hydrogen form.
- **4.6 Anion-exchange resin**, polystyrene quaternary ammonium type, 2 % to 3 % crosslinked, 150 μ m to 330 μ m, chloride form.

5 Apparatus

Ordinary apparatus and the following:

- **5.1** Rotary evaporator, with round-bottom flasks of capacity 250 ml.
- **5.2 Ion-exchange column**: open glass column of internal diameter 25 mm and length 200 mm, provided with a 10 mm to 20 mm layer of glass wool or sintered glass filter and with a stopcock.
- **5.3** Water bath, the temperature of which can be adjusted between 25 °C to 40 °C.

6 Sampling

The test sample shall be prepared and stored in accordance with ISO 607.

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7 Procedure

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7.1 Test portion is iteh ai/catalog/standards/iso/030f2f7c-10be-4bb7-8db4-cc5baa42cfc8/iso-8799-2009

From the laboratory sample, if necessary homogenized by introducing an appropriate and known quantity of water, weigh, to the nearest 1 mg, into a 100 ml beaker, a quantity of the homogeneous laboratory sample corresponding to 5 mmol of anionic-active matter.

7.2 Preparation of the ion-exchange resins

7.2.1 Anion-exchange resin clean-up

Take 100 g of the anion-exchange resin (4.6) and allow to swell in water for 24 h. Transfer the resin to a suitable column and pass 500 ml of the sodium hydroxide solution (4.4) through the column, followed by about 1 000 ml of water. Then pass 400 ml of the hydrochloric acid solution (4.3) through the column and again wash with enough water until the washings have a pH between 5 and 7. The treated resin may be stored in water.

7.2.2 Cation-exchange resin clean-up

Take 100 g of the cation-exchange resin (4.5) and allow it to swell in water for 24 h. Transfer the resin to a suitable column, pass 500 ml of the hydrochloric acid solution (4.3) through the column and wash with water until the washings have a pH between 5 and 7. The treated resin may be stored in water.