
INTERNATIONAL STANDARD



5545

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

Rennet caseins and caseinates – Determination of ash (Reference method)

Caséines présure et caséinates – Détermination des cendres (Méthode de référence)

First edition – 1978-06-15

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UDC 637.147.2 : 543.822

Ref. No. ISO 5545-1978 (E)

Descriptors : caseins, chemical analysis, determination of content, ashes, gravimetric analysis.

FOREWORD

ISO (the International Organization for Standardization) is a worldwide federation of national standards institutes (ISO member bodies). The work of developing International Standards is carried out through ISO technical committees. Every member body interested in a subject for which a technical committee has been set up 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.

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 5545 was developed by Technical Committee ISO/TC 34, *Agricultural food products*, and was circulated to the member bodies in September 1976.

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

Australia	Germany	Poland
Austria	Ghana	Portugal
Bulgaria	Hungary	Romania
Canada	India	South Africa, Rep. of
Chile	Iran	Spain
Czechoslovakia	Israel	Turkey
Egypt, Arab Rep. of	Korea, Rep. of	Yugoslavia
France	Netherlands	

The member bodies of the following countries expressed disapproval of the document on technical grounds :

New Zealand
United Kingdom

NOTE — The method specified in this International Standard has been developed jointly with the IDF (International Dairy Federation) and the AOAC (Association of Official Analytical Chemists, U.S.A.). The text as approved by the above organizations will also be published by FAO/WHO (Code of Principles concerning Milk and Milk products and Associated Standards), by the IDF and by the AOAC (Official Methods of Analysis).

Rennet caseins and caseinates – Determination of ash (Reference method)

1 SCOPE AND FIELD OF APPLICATION

This International Standard specifies a reference method for the determination of the ash of caseins obtained by rennet precipitation and of caseinates, with the exception of ammonium caseinate.

NOTE – For the determination of ash ("fixed ash") of acid caseins, of ammonium caseinates, of their mixtures with rennet casein and with caseinates, and of caseins of unknown type, see ISO 5544.

2 REFERENCES

ISO/R 707, *Milk and milk products – Sampling*.

ISO 3310/1, *Test sieves – Technical requirements and testing – Part 1: Metal wire cloth*.

ISO 5550, *Caseins and caseinates – Determination of water content (Reference method)*.¹⁾

3 DEFINITION

ash of rennet caseins or of caseinates: The substances determined by the procedure described in this International Standard and expressed as a percentage by mass.

4 PRINCIPLE

Incineration of a test portion at 825 ± 25 °C. Weighing of the residue.

5 APPARATUS

5.1 Analytical balance.

5.2 Silica or platinum dish, about 70 mm diameter and 25 to 50 mm deep.

5.3 Electrical furnace with air circulation, capable of being controlled at 825 ± 25 °C.

5.4 Desiccator, containing an effective desiccant.

5.5 Grinding device, for grinding the laboratory sample, if necessary (see 7.1.4), without development of undue heat and without loss or absorption of moisture. A hammer-mill shall not be used.

5.6 Test sieve, wire cloth, diameter 200 mm, nominal size of aperture 500 µm, with receiver, complying with ISO 3310/1.

6 SAMPLING

See ISO/R 707.

7 PROCEDURE

7.1 Preparation of the test sample

7.1.1 Thoroughly mix the laboratory sample by repeatedly shaking and inverting the container (if necessary, after having transferred all of the laboratory sample to an air-tight container of sufficient capacity to allow this operation to be carried out).

7.1.2 Transfer about 50 g of the thoroughly mixed laboratory sample to the test sieve (5.6).

7.1.3 If the 50 g portion directly passes or almost completely passes the sieve, use for the determination the sample as prepared in 7.1.1.

7.1.4 Otherwise, grind the 50 g portion, using the grinding device (5.5), until it passes the sieve. Immediately transfer all the sieved sample to an air-tight container of sufficient capacity and mix thoroughly by repeatedly shaking and inverting. During these operations, take precautions to avoid any change in the water content of the product.

7.1.5 After the test sample has been prepared, the determination (7.4) should be proceeded with as soon as possible.

7.2 Preparation of the dish

Heat the dish (5.2) in the electrical furnace (5.3), controlled at 825 ± 25 °C, for 30 min. Allow the dish to cool in the desiccator (5.4) to the temperature of the balance room and weigh to the nearest 0,1 mg.

7.3 Test portion

Weigh, to the nearest 0,1 mg, directly in or by difference into the prepared dish, approximately 3 g of the test sample (7.1).

1) At present at the stage of draft.

7.4 Determination

Heat the dish with its contents on a low flame until the test portion is completely charred, taking care that it does not burst into flame.

Transfer the dish to the electrical furnace (5.3), controlled at $825 \pm 25^\circ\text{C}$, and heat for at least 1 h until all carbon has disappeared from the dish. Allow the dish to cool in the desiccator (5.4) to the temperature of the balance room and weigh to the nearest 0,1 mg.

Repeat the operations of heating in the electrical furnace (5.3), cooling and weighing, until the mass remains constant to within 1 mg or begins to increase. Record the minimum mass.

8 EXPRESSION OF RESULTS

8.1 Method of calculation and formula

8.1.1 The ash of the sample, as a percentage by mass, is equal to

$$\frac{m_1 - m_2}{m_0} \times 100$$

where

- m_0 is the mass, in grams, of the test portion;
- m_1 is the mass, in grams, of the dish and residue;
- m_2 is the mass, in grams, of the prepared dish.

Calculate the ash to the nearest 0,01 % and report the final result to the nearest 0,1 %.

8.1.2 To calculate the ash of the sample on the dry basis, as a percentage by mass, multiply the result obtained in accordance with 8.1.1 by

$$\frac{100}{100 - M}$$

where M is the water content of the sample determined according to ISO 5550.

8.2 Precision

8.2.1 Repeatability

The difference between two single results obtained on identical test material by one analyst using the same apparatus within a short time interval will exceed 0,15 g of ash per 100 g of product on average not more than once in 20 cases in the normal and correct operation of the method.

8.2.2 Reproducibility

The difference between two single and independent results obtained by two operators working in different laboratories on identical test material will exceed 0,25 g of ash per 100 g of product on average not more than once in 20 cases in the normal and correct operation of the method.

9 TEST REPORT

The test report shall show the method used and the result obtained; it shall also mention all operating conditions not specified in this International Standard, or regarded as optional, as well as any circumstances that may have influenced the result.

The report shall include all details necessary for complete identification of the sample.