

BHUTAN STANDARD
FRUIT AND VEGETABLE PRODUCTS-
DETERMINATION OF SOLUBLE SOLIDS-
REFRACTOMETRIC METHOD



ICS 67.05

© Copyright 2019
BHUTAN STANDARD BUREAU
The National Standards Body of Bhutan
THIMPHU

BTS 277: XXXX ISO 2173: 2003

NATIONAL FOREWORD

This Bhutan Standard which is identical with ISO 2173: 2003 FRUIT AND VEGETABLES PRODUCTS-DETERMINATION OF SOLUBLE SOLIDS-REFRACTOMETRIC METHOD Standard issued by the International Organization for Standardization was adopted by Bhutan Standards Bureau by Food and Agriculture technical committee (TC 02) and approved by the Bhutan Standards Bureau Board (BSB Board) on xxxx, 2019.

The text of the ISO Standard has been approved as suitable for publication as Bhutan Standard without deviation. Certain conventions are however, not identical to those used in Bhutan Standard.

Attention is particularly drawn to the following:

a) Where the words “ISO Standard” appear referring to this standard, they should be read as “Bhutan Standard”.

b) Wherever page numbers are quoted, they are “ISO Standard” page numbers.

**Fruit and vegetable products —
Determination of soluble solids —
Refractometric method**

*Produits dérivés des fruits et légumes — Détermination du résidu sec
soluble — Méthode réfractométrique*

BSB TC use only! Not for SALE!



PDF disclaimer

This PDF file may contain embedded typefaces. In accordance with Adobe's licensing policy, this file may be printed or viewed but shall not be edited unless the typefaces which are embedded are licensed to and installed on the computer performing the editing. In downloading this file, parties accept therein the responsibility of not infringing Adobe's licensing policy. The ISO Central Secretariat accepts no liability in this area.

Adobe is a trademark of Adobe Systems Incorporated.

Details of the software products used to create this PDF file can be found in the General Info relative to the file; the PDF-creation parameters were optimized for printing. Every care has been taken to ensure that the file is suitable for use by ISO member bodies. In the unlikely event that a problem relating to it is found, please inform the Central Secretariat at the address given below.

BSB TC use only! Not for SALE!

© ISO 2003

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized in any form or by any means, electronic or mechanical, including photocopying and microfilm, without permission in writing from either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office
Case postale 56 • CH-1211 Geneva 20
Tel. + 41 22 749 01 11
Fax + 41 22 749 09 47
E-mail copyright@iso.org
Web www.iso.org

Published in Switzerland

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 2173 was prepared by Technical Committee ISO/TC 34, *Food products*, Subcommittee SC 3, *Fruit and vegetable products*.

This second edition cancels and replaces the first edition (ISO 2173:1978), which has been technically revised.

Fruit and vegetable products — Determination of soluble solids — Refractometric method

1 Scope

This International Standard specifies a refractometric method for the determination of the soluble solids in fruit and vegetable products.

This method is particularly applicable to thick products, to products containing suspended matter, and to products rich in sugar. If the products contain other dissolved substances, the results will be only approximate; nevertheless, for convenience the result obtained by this method can be considered conventionally as the soluble solids content.

NOTE For the determination of the soluble solids in fruit juices (not containing suspended matter) and in concentrated juices (clarified), the pycnometric method specified in ISO 2172 is applicable.

2 Terms and definitions

For the purposes of this document, the following terms and definitions apply.

2.1

soluble solids determined by the refractometric method

concentration of sucrose in an aqueous solution which has the same refractive index as the product analysed, under specified conditions of preparation and temperature

NOTE This concentration is expressed as a mass fraction in percent.

3 Principle

The refractive index of a test solution is measured at $20\text{ °C} \pm 0,5\text{ °C}$ using a refractometer. The refractive index is correlated with the amount of soluble solids (expressed as sucrose concentration) using tables, or by direct reading on the refractometer of the mass fraction of soluble solids.

4 Reagents

Use only reagents of recognized analytical grade.

4.1 Water

The water used shall have been distilled twice in borosilicate glass apparatus, or shall be water of at least equivalent purity.

5 Apparatus

Usual laboratory apparatus and, in particular, the following.

5.1 Refractometer

Use one of the following.

5.1.1 Refractometer indicating the refractive index, by means of a scale graduated in 0,001, in order to allow readings to be estimated to 0,000 2.

This refractometer shall be adjusted so that at $20\text{ }^{\circ}\text{C} \pm 0,5\text{ }^{\circ}\text{C}$ it registers a refractive index of 1,333 0 for distilled water.

5.1.2 Refractometer indicating the mass fraction of sucrose, by means of scale graduated in 0,10 %.

This refractometer shall be adjusted so that at $20\text{ }^{\circ}\text{C} \pm 0,5\text{ }^{\circ}\text{C}$ it registers a mass fraction of soluble solids (sucrose) of zero for distilled water.

5.2 Means for circulating water, to maintain the temperature of the prisms of the refractometer (5.1.1 or 5.1.2) constant to within $\pm 0,5\text{ }^{\circ}\text{C}$, in the neighbourhood of $20\text{ }^{\circ}\text{C}$, which is the reference temperature (see 8.1).

5.3 Beaker, of capacity 250 ml.

6 Sampling

It is important that the laboratory receive a sample which is truly representative and has not been damaged or changed during transport or storage.

7 Procedure

7.1 Preparation of test solution

7.1.1 Clear liquid products

Thoroughly mix the laboratory sample and use it directly for the determination.

7.1.2 Semi-thick products (purees, etc.)

Thoroughly mix the laboratory sample. Press a part of the sample through a gauze folded in four, rejecting the first drops of the liquid and reserving the remainder of the liquid for the determination.

7.1.3 Thick products (jams, jellies, etc.)

Weigh into the tared beaker (5.3), to the nearest 0,01 g, a suitable quantity (up to 40 g) of the laboratory sample and add 100 ml to 150 ml of water. Heat the contents of the beaker to boiling and allow to boil gently for 2 min to 3 min, stirring with a glass rod. Cool the contents and mix thoroughly.

After 20 min, weigh to the nearest 0,01 g, then filter through a fluted filter or a Büchner funnel into a dry vessel. Reserve the filtrate for the determination.

7.1.4 Frozen products

After thawing the sample and removing, if necessary, stones, pips and hard seed-cavity walls, mix the product with the liquid formed during the thawing process and proceed as described in 7.1.2 or 7.1.3 as appropriate.

7.1.5 Dried products

Cut a part of the laboratory sample into small pieces. Remove, if necessary, stones, pips and hard seed-cavity walls, and mix carefully. Then weigh into a tared beaker, to the nearest 0,01 g, 10 g to 20 g of the sample. Add 5 to 10 times this mass of water and place on a boiling water bath for 30 min, stirring from time to time with a glass rod. If necessary, prolong the heating time until a homogeneous mixture is obtained. Cool the contents of the beaker and mix well.

After 20 min, weigh to the nearest 0,01 g, then filter into a dry vessel. Reserve the filtrate for the determination.

If the test solution is too dark to be read in the refractometer, dilute the test solution with concentrated sugar solution; never use water for this purpose. Mix weighed amounts of the solution under examination and a solution of pure sugar of about the same strength (see reference [1]).

7.2 Determination

Adjust the water circulation (5.2) in order to operate at the required temperature (between 15 °C and 25 °C) and allow it to flow to bring the prisms of the refractometer (5.1.1 or 5.1.2) to the same temperature, which shall remain constant to within $\pm 0,5$ °C during the determination.

Bring the test solution (7.1) to the measuring temperature. Put a small quantity of test solution (2 or 3 drops are sufficient) onto the fixed prism of the refractometer (5.1.1 or 5.1.2) and immediately adjust the movable prism. Suitably illuminate the field of view. The use of a sodium vapour lamp allows more precise results to be obtained (especially in the case of coloured and dark products).

Bring the line dividing the light and dark parts of the surface into the field of view to the crossing of the threads. Read the value of the refractive index or the mass fraction of sucrose, according to the instrument used (5.1.1 or 5.1.2).

8 Expression of results

8.1 Corrections

8.1.1 If the determination has been carried out at a temperature other than $20\text{ °C} \pm 0,5\text{ °C}$, the following corrections are required.

a) For the scale indicating the refractive index (5.1.1), apply the formula:

$$n_D^{20} = n_D^t + 0,0013(t - 20)$$

where

n_D^{20} is the refractive index at 20 °C;

n_D^t is the refractive index at the temperature of measurement;

t is the temperature of measurement, in degrees Celsius.

b) For the scale indicating the mass fraction of sucrose (5.1.2), correct the result according to Table A.1.

8.1.2 If the determination has been carried out for the products with added salt, correct the refractometer reading, expressed as a concentration of sucrose at $20\text{ °C} \pm 0,5\text{ °C}$, for added salt by the following formula (see reference [2]):

$$S = (R - N) \times 1,016$$

where

- S is the mass fraction of soluble solids, in percent, as sucrose, corrected for added NaCl;
- R is the refractometer reading, as a mass fraction in percent, as sucrose;
- N is the total chloride content, as a mass fraction in percent, expressed as NaCl;
- 1,016 is the correction factor for added salt.

8.1.3 If the determination has been carried out for the highly acidic products, such as citrus juices and concentrated citrus juices, correct the refractometer reading, expressed as a mass fraction of sucrose at $20\text{ °C} \pm 0,5\text{ °C}$, by making the following addition to the refractometric reading (see reference [3]):

$$0,012 + 0,193 \times M - 0,000\,4 \times M^2$$

where M is the total acidity expressed in grams per 100 g, at pH = 8,1, expressed as anhydrous citric acid (see reference [4]). The calculated values for this expression are given in Table A.2.

8.2 Calculation method

8.2.1 Refractometer with refractive index scale

8.2.1.1 Read from Table A.3 the mass fraction of soluble solids corresponding to the value read in accordance with 7.2, corrected if necessary in accordance with 8.1.1 a). In the case of liquid or semi-thick products (7.1.1 or 7.1.2), the mass fraction of soluble solids is equal to the number found. If the determination has been carried out on a diluted solution (7.1.3 or 7.1.5), the mass fraction of soluble solids, in percent, is equal to

$$(P \cdot m_1) / m_0$$

where

- P is the mass fraction of soluble solids in the diluted solution, in percent;
- m_0 is the mass, in grams, of the sample before dilution (7.1.3 or 7.1.5);
- m_1 is the mass, in grams, of the sample after dilution (7.1.3 or 7.1.5).

8.2.1.2 If the determination has been carried out on a dark solution (see 7.1.5) that has to be diluted with concentrated sugar solution, the mass fraction of soluble solids, in percent, is equal to (see reference [1])

$$\frac{[(m_W + m_B) C - m_B D]}{m_W}$$

where

- m_W is the mass, in grams, of the sample diluted with sugar solution;
- m_B is the mass, in grams, of sugar solution used in the dilution;
- C is the mass fraction of soluble solids, in percent, in the mixture ($m_W + m_B$), obtained from the refractive index;
- D is the mass fraction of soluble solids, in percent, in the pure sugar solution, obtained from its refractive index.

Express the result to one decimal place.

8.2.2 Refractometer with sucrose scale

In the case of liquid or semi-thick products (7.1.1 or 7.1.2), the mass fraction of soluble solids, in percent, as sucrose, is equal to the value read in accordance with 8.1.1 b). If the determination has been carried out on a diluted solution (7.1.3 or 7.1.5), calculate the mass fraction of soluble solids by means of the formula given in 8.2.1.1, or the formula in 8.2.1.2 for a dark solution diluted with a sugar solution (see 7.1.5).

Express the result to one decimal place.

9 Repeatability

The absolute difference between two independent single test results, obtained using the same method on identical test material in the same laboratory by the same operator using the same equipment within a short interval of time, will in not more than 5 % of cases be greater than 0,5 g of soluble solids per 100 g or per 100 ml of product.

10 Test report

The test report shall specify:

- all information necessary for the complete identification of the sample;
- the sampling method used, if known;
- the test method used, with reference to this International Standard;
- all operating details not specified in this International Standard, or regarded as optional, together with details of any incidents which may have influenced the test result(s);
- the test result(s) obtained;
- if the repeatability has been checked, the final quoted result obtained.

Annex A (normative)

Correction and conversion tables

Table A.1 — Correction of readings of the refractometer with scale indicating sucrose for a temperature different from 20 °C ± 0,5 °C

Temperature °C	Scale reading for soluble solids, % (by mass)									
	5	10	15	20	25	30	40	50	60	70
Corrections to be subtracted										
15	0,29	0,31	0,33	0,34	0,34	0,35	0,37	0,38	0,39	0,40
16	0,24	0,25	0,26	0,27	0,28	0,28	0,30	0,30	0,31	0,32
17	0,18	0,19	0,20	0,21	0,21	0,21	0,22	0,23	0,23	0,24
18	0,13	0,13	0,14	0,14	0,14	0,14	0,15	0,15	0,16	0,16
19	0,06	0,06	0,07	0,07	0,07	0,07	0,08	0,08	0,08	0,08
Corrections to be added										
21	0,07	0,07	0,07	0,07	0,08	0,08	0,08	0,08	0,08	0,08
22	0,13	0,14	0,14	0,15	0,15	0,15	0,15	0,16	0,16	0,16
23	0,20	0,21	0,22	0,22	0,23	0,23	0,23	0,24	0,24	0,24
24	0,27	0,28	0,29	0,30	0,30	0,31	0,31	0,31	0,32	0,32
25	0,35	0,36	0,37	0,38	0,38	0,39	0,40	0,40	0,40	0,40

Table A.2 — Acid correction (for citrus juices and citrus juice concentrates)

Total acid pH = 8,1 expressed as anhydrous citric acid (g acid/100 g)	Correction value ^a	Total acid pH = 8,1 expressed as anhydrous citric acid (g acid/100 g)	Correction value ^a
0,2	0,04	4,2	0,81
0,4	0,08	4,4	0,85
0,6	0,12	4,6	0,89
0,8	0,16	4,8	0,93
1,0	0,20	5,0	0,97
1,2	0,24	5,2	1,01
1,4	0,28	5,4	1,04
1,6	0,32	5,6	1,07
1,8	0,36	5,8	1,11
2,0	0,39	6,0	1,15
2,2	0,43	6,2	1,19
2,4	0,47	6,4	1,23
2,6	0,51	6,6	1,27
2,8	0,55	6,8	1,30
3,0	0,58	7,0	1,34
3,2	0,62		
3,4	0,66		
3,6	0,70		
3,8	0,74		
4,0	0,78		

^a The correction values given should be added to the refractometer readings with scale indicating sucrose obtained at 20 °C ± 0,5 °C

Table A.3 — Refractive index and corresponding mass fraction of soluble solids (sucrose)

Refractive index n_D^{20}	Soluble solids (sucrose) % (by mass)	Refractive index n_D^{20}	Soluble solids (sucrose) % (by mass)	Refractive index n_D^{20}	Soluble solids (sucrose) % (by mass)	Refractive index n_D^{20}	Soluble solids (sucrose) % (by mass)
1,333 0	0	1,367 2	22	1,407 6	44	1,455 8	66
1,334 4	1	1,368 9	23	1,409 6	45	1,458 2	67
1,335 9	2	1,370 6	24			1,460 6	68
1,337 3	3	1,372 3	25	1,411 7	46	1,463 0	69
1,338 8	4			1,413 7	47	1,465 4	70
1,340 3	5	1,374 0	26	1,415 8	48		
		1,375 8	27	1,417 9	49	1,467 9	71
1,341 8	6	1,377 5	28	1,420 1	50	1,470 3	72
1,343 3	7	1,379 3	29			1,472 8	73
1,344 8	8	1,381 1	30	1,422 2	51	1,475 3	74
1,346 3	9			1,424 3	52	1,477 8	75
1,347 8	10	1,382 9	31	1,426 5	53		
		1,384 7	32	1,428 6	54	1,480 3	76
1,349 4	11	1,386 5	33	1,430 8	55	1,482 9	77
1,350 9	12	1,388 3	34			1,485 4	78
1,352 5	13	1,390 2	35	1,433 0	56	1,488 0	79
1,354 1	14			1,435 2	57	1,490 6	80
1,355 7	15	1,392 0	36	1,437 4	58		
		1,393 9	37	1,439 7	59	1,493 3	81
1,357 3	16	1,395 8	38	1,441 9	60	1,495 9	82
1,358 9	17	1,397 8	39			1,498 5	83
1,360 5	18	1,399 7	40	1,444 2	61	1,501 2	84
1,362 2	19			1,446 5	62	1,503 9	85
1,363 8	20	1,401 6	41	1,448 8	63		
		1,403 6	42	1,451 1	64		
1,365 5	21	1,405 6	43	1,453 5	65		

Bibliography

- [1] AOAC Official Method 932.14, *Solids in sirups*. AOAC Official Methods of Analysis, 1995, 44.1.04
- [2] AOAC Official Method 970.59, *Solids (Soluble) in Tomato Products: Refractive index method*. AOAC Official Methods of Analysis, 1995, 42.1.10
- [3] EN 12143:1996, *Fruit and vegetable juices — Estimation of soluble solids content — Refractometric method*
- [4] ISO 750:1998, *Fruit and vegetable products — Determination of titratable acidity*
- [5] ISO 2172:1983, *Fruit juice — Determination of soluble solids content — Pycnometric method*

BSB TC use only! Not for SALE!

BSB TC use only! Not for SALE!

SUB COMMITTEE ON JUICES

(TC 02/SC 05)

Organization

Representative(s)

National Food Testing Laboratory, BAFRA

Mr. Kanjur Wangdi

(Convener)

Bhutan Agro Industry Limited

Mrs. Nim Dem Hingmang

Bhutan Agro Industry Limited

Mrs.Jigme Wangmo

Bhutan Fruit Products Private limited

Ms.Karma Sonam

Bhutan Milk and Agro Pvt. Limited

Dr. M. Sharma

Bhutan Milk and Agro Pvt. Limited

Ms.Kinga Delma

National Food Testing Laboratory, BAFRA

Mr. Anil Rai

National Post Harvest Center, Paro

Mr. Sonam Tobgay

National Post Harvest Center, Paro

Mr. Dorji Rinchen

Zimdra Food Pvt. Limited, P/ling

Mr. Umesh Parsad

Zimdra Food Pvt. Limited, P/ling

Mr. Rinchen Chedup

Bhutan Standards Bureau

Mr. Sonam Phuntsho,
Director General
(Ex-officio member)

Member Secretary

Ms. Tashi Choden
Standardization Division
Bhutan Standards Bureau

BTS 277: XXXX ISO 2173: 2003

FOOD AND AGRICULTURE TECHNICAL COMMITTEE

(TC 02)

Organization

Representative(s)

National Dairy Research Development Centre

Mr.Phuntsho.T.Norbu
(Chairperson)

Bhutan Agriculture and Food Regulatory Authority

Mr. Jamyang Phuntsho

Bhutan Agriculture and Food Regulatory Authority

Mrs. Gyem Bidha

Bhutan Agro Industry Limited

Mrs. Nim Dem Hingmang

Bhutan Agro Industry Limited

Mrs. Jigme Wangmo

Bhutan Exporters Association

Mr. Dorji Tshering

Bhutan Livestock Development Corporation Limited

Mr. Sithar Dorji

Bhutan Livestock Development Corporation Limited

Mr. Pema Khandu

Department of Agriculture, MoAF

Mrs.Pema Choden

Department of Agriculture, MoAF

Mr. Jimba Rabgyel

Department of Agriculture and Marketing Cooperatives

Mr. Dawa Tshering

Department of Agriculture and Marketing Cooperatives

Mr.Tashi Tshering

Department of Industry, MoEA

Mr.Tashi Dorji

National Post Harvest Center, Paro

Mr.Dechen Tshering

Office of Consumer Protection, MoEA

Mr. Jigme Dorji

Office of Consumer Protection, MoEA

Mrs. Chencho Zangmo

Bhutan Standards Bureau

Mr. Sonam Phuntsho,
Director General
(Ex-officio member)

Member Secretary

Ms. Tashi Choden
Standardization Division
Bhutan Standards Bureau