INTERNATIONAL STANDARD

ISO 6321

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AMENDMENT 1 1998-03-15

Animal and vegetable fats and oils — Determination of melting point in open capillary tubes (slip point)

AMENDMENT 1: Palm oil samples

iTeh Scorps gras d'origines animale et végétale — Détermination du point de fusion en tube capillaire ouvert AMENDEMENT 1: Échantillons d'huile de palme

ISO 6321:1991/Amd 1:1998 https://standards.iteh.ai/catalog/standards/sist/f872a379-fce2-4eff-b08c-407c36dd2cf8/iso-6321-1991-amd-1-1998



Foreword

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Amendment 1 to International Standard ISO 6321:1991 was prepared by Technical Committee ISO/TC 34, *Agricultural food products*, Subcommittee SC 11, *Animal and vegetable fats and oils*.

ISO 6321:1991/Amd 1:1998

Annex A forms an integral part of this international Standardrds/sist/f872a379-fce2-4eff-b08c-407c36dd2cf8/iso-6321-1991-amd-1-1998

Annex B is for information only.

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Animal and vegetable fats and oils — Determination of melting point in open capillary tubes (slip point)

AMENDMENT 1: Palm oil samples

Page 1, clause 1

Add the following phrase before the Notes:

"A method for the determination of the melting point of palm oil samples is given in annex A."

Page 5, subclause 10.1

Add the following phrase at the end of this subclause:

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"The results of interlaboratory tests on palm oil samples are given in annex B."

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Add annexes A and B.

Annex A

(normative)

Method for palm oil samples

Melt the sample and filter through a filter paper. Conduct the filtration in an oven set at 60 °C to avoid any crystallization of the sample. Leave the filtered sample in the oven for 10 min until it is free of air bubbles.

Dip at least three clean capillary tubes into the liquid sample so that columns of fat approximately 10 mm high are obtained in the tubes. Immediately chill the columns of fat by holding and rolling the ends of the tubes containing the sample pressed against a piece of ice, until the fat has solidified. Do not allow the open end of the tube to touch the ice. Wipe the tubes against a piece of tissue paper as quickly as possible. Place the tubes in a test tube which is held in a beaker of water that has been equilibrated at 10 °C \pm 1 °C in a thermostated water bath. Transfer the beaker to the water bath and hold for 16 h at 10 °C \pm 1 °C.

For the determination, follow the procedure as given in 8.3.1 to 8.3.3 Regulate the rise in temperature in the water jacket to 1 °C/min, slowing down to 0,5 °C/min as the slip point is reached. Note the temperature value indicated by the thermometer as soon as the fat rises in each of the tubes.

Note the arithmetic mean of the three readings obtained and take this as the result of one determination.

Annex B

(informative)

Results of interlaboratory tests on palm oil samples

Samples		Slip point			
		MS 817:1989	ISO 6	ISO 6321	
		AOCS CC 3-251)	Method A	Method B	
	1	36,8	38,2	36,5	
Palm oil, RBD ²⁾	2	35,3	37,4	35,5	
	3	35,2	37,7	35,5	
	4	36,6	38,0	36,5	
	5	35,6	37,5	35,5	
Palm olein, RBD	1	22,3	24,4	25,5	
	2	22,2	24,4	25,5	
	3	22,5	24,3	25,5	
	4	22,5	24,2	24,9	
	5	22,3	24,2	24,9	
	1	51,6	51,8	51,5	
• 🗂	2	52,8		52,8	
Palm stearin, RBD	l Gh	SIA46,0ARI	PRE44,8 IEW	45,0	
	4	52,3	52,8	53,4	
	5	(stağğards.	iteh.a52,8	51,5	
Crude palm oil https:	1	35,8	35,6	26,0	
	2	ISC35321:1991/An	<u>rd 1:1998</u> 36,6	26,0	
	//stænda	rds.iteh.ai/catglag/standards/	sist/f872a37 <mark>36</mark> ,fae2-4eff-b08	c- 26,0	
	4	407c36dd23586jso-6321-19	91-amd-1-35,88	26,0	
	5	35,8	36,8	26,0	
Crude palm kernel oil	1	27,8	27,7	27,6	
	2	26,6	27,8	27,6	
	3	26,7	26,7	27,0	
	4	26,8	26,7	27,0	
	5	27,0	27,5	27,4	
Palm kernel oil, RBD	1	27,8	27,8	28,2	
	2	27,8	27,6	27,6	
	3	27,7	27,5	28,0	
	4	27,8	27,2	28,0	
	5	27,6	27,3	27,8	
Palm kernel olein, RBD	1	26,2	25,8	26,0	
	2	23,4	23,3	23,8	
	3	23,5	23,4	23,8	
	4	23,4	23,4	23,8	
	5	24,6	24,4	24,5	
	1	32,2	32,2	33,0	
	2	32,2	32,8	33,0	
Palm kernel stearin, RBD	3	39,3	38,5	39,4	
	4	33,3	33,0	33,2	
	5	32,3	33,6	33,2	

Table B.1 — Comparison of methods

	Palm oil	Palm olein	Palm stearin
No. of laboratories retained after eliminating outliers	10	11	11
Mean (°C)	37,4	20,5	52,1
Standard deviation of repeatability s_r (°C)	0,23	0,15	0,09
Coefficient of variation of repeatability	0,6	0,7	0,2
Repeatability limit, $2,8 \times s_r$ (°C)	0,64	0,42	0,25
Standard deviation of reproducibility s_R (°C)	0,78	0,98	0,54
Coefficient of variation of reproducibility	2,1	4,8	1,0
Reproducibility limit, 2,8 × s_R (°C)	2,2	2,7	1,5

Table B.2 — Statistical results of interlaboratory tests

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