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**Isobutene-isoprene rubber (IIR) —  
Evaluation procedure**

*Caoutchouc isobutène-isoprène (IIR) — Méthode d'évaluation*

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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.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular, the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see [www.iso.org/directives](http://www.iso.org/directives)).

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. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see [www.iso.org/patents](http://www.iso.org/patents)).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT), see [www.iso.org/iso/foreword.html](http://www.iso.org/iso/foreword.html).

This document was prepared by Technical Committee ISO/TC 45, *Rubber and rubber products*, Subcommittee SC 3, *Raw materials (including latex) for use in the rubber industry*.

This seventh edition cancels and replaces the sixth edition (ISO 2302:2014), which has been technically revised.

The main changes compared to the previous edition are as follows:

- the normative references have been updated in [Clause 2](#), [5.3](#) and [Clause 10 d](#)), where ISO 247 has been replaced by ISO 247-1 and ISO 247-2;
- the normative references have been updated in [Clause 2](#), [6.2.2.3 j](#)) and [7.1](#), where ISO 3417 has been replaced by ISO 6502-2;
- a footnote has been added to [Table 1](#) allowing the possibility to replace TMDT by an alternative curative.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at [www.iso.org/members.html](http://www.iso.org/members.html).

# Isobutene-isoprene rubber (IIR) — Evaluation procedure

**WARNING** — Users of this document should be familiar with the normal laboratory practice. This document does not purport to address all of the safety problems, if any, associated with its use. It is the responsibility of the user to establish appropriate safety and health practices.

## 1 Scope

This document specifies:

- the physical and chemical tests on raw rubbers, and
- the standardized materials, a standardized test formulation, equipment, and processing methods for evaluating the vulcanization characteristics of all types of isobutene-isoprene rubber (IIR).

## 2 Normative references

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements 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 37, *Rubber, vulcanized or thermoplastic — Determination of tensile stress-strain properties*

ISO 247-1:2018, *Rubber — Determination of ash — Part 1: Combustion method*

ISO 247-2:2018, *Rubber — Determination of ash — Part 2: Thermogravimetric analysis (TGA)*

ISO 248-1, *Rubber, raw — Determination of volatile-matter content — Part 1: Hot-mill method and oven method*

ISO 248-2, *Rubber, raw — Determination of volatile-matter content — Part 2: Thermogravimetric methods using an automatic analyser with an infrared drying unit*

ISO 289-1, *Rubber, unvulcanized — Determinations using a shearing-disc viscometer — Part 1: Determination of Mooney viscosity*

ISO 1795:2017, *Rubber, raw natural and raw synthetic — Sampling and further preparative procedures*

ISO 2393, *Rubber test mixes — Preparation, mixing and vulcanization — Equipment and procedures*

ISO 6502-1, *Rubber — Measurement of vulcanization characteristics using curemeters — Part 1: Introduction*

ISO 6502-2, *Rubber — Measurement of vulcanization characteristics using curemeters — Part 2: Oscillating disc curemeter*

ISO 23529, *Rubber — General procedures for preparing and conditioning test pieces for physical test methods*

## 3 Terms and definitions

No terms and definitions are listed in this document.

ISO and IEC maintain terminological databases for use in standardization at the following addresses:

- ISO Online browsing platform: available at <https://www.iso.org/obp>
- IEC Electropedia: available at <http://www.electropedia.org/>

## 4 Sampling and further preparative procedure

- 4.1 Select the sample from the lot in accordance with ISO 1795.
- 4.2 Take a laboratory sample of approximately 1,5 kg by the method described in ISO 1795.
- 4.3 Prepare test samples in accordance with ISO 1795.

## 5 Physical and chemical tests on raw rubber

### 5.1 Mooney viscosity

Prepare a test sample in accordance with the preferred procedure in ISO 1795, i.e. without milling, cutting the test sample directly from the laboratory sample. The test sample shall be as free as possible from air and pockets that can trap air against the rotor and die surface.

If agreed between the interested parties or if the condition of the test sample (e.g. excessive porosity) makes milling necessary, it shall be performed in accordance with paragraphs 1 and 2 of ISO 1795:2017, 7.3.2.2.

Determine the Mooney viscosity in accordance with ISO 289-1 on this test portion, as ML(1+8) at 125 °C.

### 5.2 Volatile matter

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Determine the volatile-matter content by the hot-mill method or by the oven method as specified in ISO 248-1 or by the method specified in ISO 248-2.

### 5.3 Ash

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Determine the ash in accordance with either method A, method B or method C of ISO 247-1:2018, or method A of ISO 247-1:2018.

## 6 Preparation of test mixes for evaluation of isobutene-isoprene rubbers

### 6.1 Standard test formulation

The standard test formulation is given in [Table 1](#). The materials shall be national or international standard reference materials. If no standard reference material is available, the materials to be used shall be agreed by the parties concerned.

**Table 1 — Standard test formulation**

Material	Parts by mass
Isobutene-isoprene rubber (IIR)	100,00
Stearic acid <sup>b</sup>	1,00
Industry reference black <sup>a</sup>	50,00
Zinc oxide <sup>b</sup>	3,00
Sulfur <sup>b</sup>	1,75
<sup>a</sup> The current industry reference black shall be used. <sup>b</sup> Powder materials shall be used (standard curing ingredients used in the industry). <sup>c</sup> It is possible to replace TMTD by an appropriate amount of an alternative curative (e.g. TBzTD) to prevent release of toxic chemical from the mixture prepared for evaluation of isobutene-isoprene rubber.	

Table 1 (continued)

Material	Parts by mass
Tetramethylthiuram disulfide(TMTD) <sup>b,c</sup>	1,00
Total	156,75
<p><sup>a</sup> The current industry reference black shall be used.</p> <p><sup>b</sup> Powder materials shall be used (standard curing ingredients used in the industry).</p> <p><sup>c</sup> It is possible to replace TMTD by an appropriate amount of an alternative curative (e.g. TBzTD) to prevent release of toxic chemical from the mixture prepared for evaluation of isobutene-isoprene rubber.</p>	

## 6.2 Procedure

### 6.2.1 Equipment and procedure

The equipment and procedure for the preparation, mixing, and vulcanization shall be in accordance with ISO 2393.

### 6.2.2 Mixing procedures

#### 6.2.2.1 General

Three alternative mixing procedures are specified, but in accordance with ISO 2393, the laboratory internal mixer procedure is preferred.

- method A: mill mixing;
- method B: single-stage mixing using a laboratory internal mixer (the preferred procedure);
- method C: two-stage mixing using a laboratory internal mixer for initial mixing and a mill for final mixing.

NOTE These procedures can give unidentical results.

#### 6.2.2.2 Method A — Mixing with a laboratory mill

The standard laboratory mill-batch mass, in grams, shall be based on four times the formulation mass (i.e.  $4 \times 156,75 \text{ g} = 627 \text{ g}$ ). The surface temperature of the rolls shall be maintained at  $45 \text{ °C} \pm 5 \text{ °C}$  throughout the mixing.

A good rolling bank at the nip of the rolls shall be maintained during mixing. If this is not obtained with the nip settings specified hereunder, small adjustments to the mill openings may be necessary.

	Duration (min)	Cumulative time (min)
a) Band the rubber with the mill opening set at 0,65 mm.	1,0	1,0
b) Mix the carbon black and the stearic acid and add evenly across the mill rolls at a uniform rate. Increase the mill opening at intervals to maintain a constant rolling bank. When all the carbon black has been incorporated, make a 3/4 cut from each side.	10,0	11,0

Do not cut the batch while free carbon black is evident in the bank or on the milling surface. Be certain to return to the batch any materials that drop through the mill.

- |   |     |      |
|---|-----|------|
| c) Add the zinc oxide, the sulfur, and the TMTD.  | 3,0 | 14,0 |
| d) Make three alternating 3/4 cuts from each side.  | 2,0 | 16,0 |
| e) Cut the batch from the mill. Set the mill opening to 0,8 mm and pass the rolled batch endwise through the mill six times.  | 2,0 | 18,0 |
| f) Sheet the batch to approximately 6 mm and check-weigh the batch (see ISO 2393). If the mass of the batch differs from the theoretical value by more than $\begin{matrix} +0,5 \\ -1,5 \end{matrix}$ %, discard the batch and re-mix. |     |      |
| g) Remove sufficient material for curemeter testing.  |     |      |
| h) Sheet the batch to approximately 2,2 mm for preparing test sheets or to the appropriate thickness for preparing ISO ring or dumbbell test pieces in accordance with ISO 37.  |     |      |
| i) After mixing, condition the batch for at least 2 h but not more than 24 h, if possible at standard laboratory temperature and humidity as defined in ISO 23529.  |     |      |

### 6.2.2.3 Method B — Single stage mixing using a laboratory internal mixer

For laboratory internal mixers having a nominal capacity of 65 cm<sup>3</sup> to about 2 000 cm<sup>3</sup>, the batch mass shall be equal to the nominal mixer capacity, in cubic centimetres, multiplied by the density of the compound. For each batch mixed, the laboratory internal mixer conditions shall be the same during the preparation of a series of identical mixes. At the beginning of each series of test mixes, a machine-conditioning batch shall be mixed using the same formulation as the mixes under test. The laboratory internal mixer shall be allowed to cool down to 60 °C between the end of one test batch and the start of the next. The temperature control conditions shall not be altered during the mixing of a series of test batches.

The mixing technique shall be such as to obtain a good dispersion of all the ingredients.

The temperature of the batch discharged on completion of mixing shall not exceed 120 °C. If necessary, adjust the batch mass or the mixer head starting temperature so that this condition is met

In the following procedure, compounding materials other than rubber and carbon black, may be added to the batch more precisely and with greater ease if they are previously blended together in the proportions required by the formulation. Such blends may be made using one of the following:

- a mortar and pestle;
- a double-cone mixer (mix for 10 min with the intensifier bar turning);
- a blender (mix for five periods of 3 s each, scraping the inside of the blender to dislodge material stuck to the sides after each 3 s period) (a “Waring”-type blender has been found suitable for this method).

**CAUTION — If the mixing periods are longer than 3 s, the stearic acid may melt, thus preventing good dispersion.**

A general mixing procedure for the laboratory internal mixer is as follows:

	Duration (min)	Cumulative time (min)
a) Load the rubber, lower the ram, and allow the rubber to be masticated.	0,5	0,5
b) Raise the ram and add the zinc oxide, sulfur, stearic acid and TMTD, taking care to avoid any loss. Then add the carbon black. Sweep down the mixer throat and lower the ram.	1,0	1,5



- |  |     |     |
|--|-----|-----|
| c) Allow the batch to mix.   | 2,0 | 3,5 |
| d) Raise the ram, clean the mixer throat and the top of the ram, and lower the ram.  | 0,5 | 4,0 |
| e) Allow the batch to mix.   | 1,0 | 5,0 |
| f) Discharge the batch and immediately check the temperature with a suitable measuring device.   |     |     |
| g) The final temperature of the discharged batch after the cumulative mixing time of 5 min shall not exceed 120 °C. If it does, discard the batch and repeat the procedure with a different batch mass or starting temperature.  |     |     |
| h) Pass the batch through a mill set at 50 °C ± 5 °C twice with a 3,0 mm mill opening.   |     |     |
| i) Check the batch mass (see ISO 2393) and record. If it differs from the theoretical value by more than $\begin{matrix} +0,5 \\ -1,5 \end{matrix}$ %, discard the batch and re-mix.   |     |     |
| j) Cut out a test piece for determining the vulcanization characteristics in accordance with ISO 6502-2 or ISO 6502-1, if required. Condition the test piece for 2 h to 24 h at 23 °C ± 3 °C before testing.   |     |     |
| k) If required, sheet the batch to approximately 2,2 mm for preparing test sheets or to the appropriate thickness for preparing ISO ring or dumbbell test pieces in accordance with ISO 37. To obtain the effects of mill direction, pass the folded batch four times between mill rolls set at 50 °C ± 5 °C and at the appropriate mill opening. Cool on a flat, dry surface. |     |     |
| l) After mixing and prior to vulcanizing, condition the batch for at least 2 h but not more than 24 h, if possible at standard laboratory temperature and humidity as defined in ISO 23529.  |     |     |

For a miniature internal mixer having a nominal mixing capacity of 65 cm<sup>3</sup>, a batch mass corresponding to 0,47 times the formulation mass (i.e.  $0,47 \times 156,75 = 73,67$  g) has been found to be suitable.

Mix with the head temperature of the miniature internal mixer maintained at 60 °C ± 3 °C and a starting (unloaded) rotor speed of 6,3 rad/s to 6,6 rad/s (60 rev/min to 63 rev/min).

Prepare the rubber by passing it once through a laboratory mill with the roll temperature set at 50 °C ± 5 °C and an opening of 0,5 mm. Cut the sheet thus produced into strips 25 mm wide.

For an internal mixer having a nominal capacity of 1 170 cm<sup>3</sup> ± 40 cm<sup>3</sup>, a batch mass corresponding to (8,5 × 156,75 g = 1 332 g) has been found to be suitable.

The speed of the fast rotor shall be set at 7 rad/s to 8 rad/s (67 rev/min to 87 rev/min).

#### 6.2.2.4 Method C — Two-stage mixing using a laboratory internal mixer for initial mixing and a mill for final mixing

##### 6.2.2.4.1 General

The mixing technique shall be such as to obtain a good dispersion of all the ingredients.

The temperature of the batch discharged on completion of mixing shall be between 150 °C and 170 °C. If necessary, adjust the batch mass or the mixer head starting temperature so that this condition is met.

During final mixing, a good rolling bank at the nip of the rolls shall be maintained. If this is not obtained with the nip setting specified, small adjustments to the mill openings may be necessary.

A general mixing procedure for the laboratory internal mixer is given in [6.2.2.4.2](#) and [6.2.2.4.3](#).