

---

---

**Rubber — Standard reference elastomers  
(SREs) for characterizing the effect of  
liquids on vulcanized rubbers**

*Caoutchouc — Élastomères de référence normalisés (SRE) pour la  
caractérisation de l'effet des liquides sur les caoutchoucs vulcanisés*

STANDARDSISO.COM : Click to view the full PDF of ISO 13226:2011



STANDARDSISO.COM : Click to view the full PDF of ISO 13226:2011



**COPYRIGHT PROTECTED DOCUMENT**

© ISO 2011

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](mailto:copyright@iso.org)  
Web [www.iso.org](http://www.iso.org)

Published in Switzerland

## Contents

Page

Foreword .....	iv
Introduction.....	v
<b>1 Scope</b> .....	<b>1</b>
<b>2 Normative references</b> .....	<b>1</b>
<b>3 Composition</b> .....	<b>2</b>
<b>4 Preparation</b> .....	<b>2</b>
<b>5 Description</b> .....	<b>2</b>
<b>6 Test sheet properties</b> .....	<b>2</b>
<b>7 Designation</b> .....	<b>2</b>
<b>8 Storage</b> .....	<b>3</b>
<b>Annex A (normative) Acrylic rubbers: SRE-ACM/1 and SRE-ACM/1X</b> .....	<b>4</b>
<b>Annex B (normative) Acrylonitrile-butadiene rubbers: SRE-NBR 28/P and SRE-NBR 28/PX</b> .....	<b>6</b>
<b>Annex C (normative) Acrylonitrile-butadiene rubbers: SRE-NBR 28/S, SRE-NBR 28/SX, SRE-NBR 34/S and SRE-NBR 34/SX</b> .....	<b>9</b>
<b>Annex D (normative) Acrylonitrile-butadiene rubbers: SRE-NBR/M</b> .....	<b>13</b>
<b>Annex E (normative) Acrylonitrile-butadiene rubbers: SRE-NBR/L</b> .....	<b>14</b>
<b>Annex F (normative) Chlorobutyl rubbers: SRE-CIIR/1</b> .....	<b>15</b>
<b>Annex G (normative) Chloroprene rubbers: SRE-CR/1</b> .....	<b>16</b>
<b>Annex H (normative) Ethylene-propylene rubbers: SRE-EPM/1</b> .....	<b>17</b>
<b>Annex I (normative) Fluoropolymer rubbers: SRE-FKM/1</b> .....	<b>18</b>
<b>Annex J (normative) Fluoropolymer rubbers: SRE-FKM/2X</b> .....	<b>19</b>
<b>Annex K (normative) Hydrogenated acrylonitrile-butadiene rubbers: SRE-HNBR/1 and SRE-HNBR/1X</b> .....	<b>22</b>
<b>Annex L (normative) Natural rubbers: SRE-NR/1</b> .....	<b>26</b>
<b>Annex M (normative) Silicone rubbers: SRE-MQ/1</b> .....	<b>27</b>
<b>Annex N (normative) Silicone rubbers: SRE-VMQ1 and SRE-VMQ/1X</b> .....	<b>28</b>

## 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 13226 was prepared by Technical Committee ISO/TC 45, *Rubber and rubber products*, Subcommittee SC 2, *Testing and analysis*.

This third edition cancels and replaces the second edition (ISO 13226:2005), which has been revised to eliminate errors and inconsistencies. It also incorporates the Technical Corrigendum ISO 13226:2005/Cor.1:2007.

STANDARDSISO.COM : Click to view the full PDF of ISO 13226:2011

## Introduction

The materials covered by this International Standard are specified in Annexes A to N. The following standard reference elastomers are included:

a) Acrylic rubbers:

- SRE-ACM/1 and SRE-ACM/1X

b) Acrylonitrile-butadiene rubbers:

- SRE-NBR 28/P and SRE-NBR 28/PX
- SRE-NBR 28/S, SRE-NBR 28/SX, SRE-NBR 34/S and SRE-NBR 34/SX
- SRE-NBR/M
- SRE-NBR/L

c) Chlorobutyl rubbers:

- SRE-CIIR/1

d) Chloroprene rubbers:

- SRE-CR/1

e) Ethylene-propylene rubbers:

- SRE-EPM/1

f) Fluoropolymer rubbers:

- SRE-FKM/1
- SRE-FKM/2X

g) Hydrogenated acrylonitrile-butadiene rubbers:

- SRE-HNBR/1 and SRE-HNBR/1X

h) Natural rubbers:

- SRE-NR/1

i) Silicone rubbers:

- SRE-MQ/1
- SRE-VMQ/1 and SRE-VMQ/1X

[STANDARDSISO.COM](https://standardsiso.com) : Click to view the full PDF of ISO 13226:2011

# Rubber — Standard reference elastomers (SREs) for characterizing the effect of liquids on vulcanized rubbers

**WARNING** — Persons using this International Standard should be familiar with normal laboratory practice. This standard 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 and to ensure compliance with any national regulatory conditions.

**IMPORTANT** — Certain procedures specified in this International Standard might involve the use or generation of substances, or the generation of waste, that could constitute a local environmental hazard. Reference should be made to appropriate documentation on safe handling and disposal after use.

## 1 Scope

This International Standard specifies requirements for vulcanized rubbers in sheet form for use as standards in characterizing the effect of test liquids and service fluids. Details of the individual standard reference elastomers are listed in the annexes.

The compounding and preparation ensure that the property profile agrees sufficiently with that of the material group represented, while the simple formulation ensures reliable reproducibility.

The specified property changes of the SRE when in contact with a fluid under specified conditions may be included as supplementary data in specifications for the fluid concerned.

This International Standard is not designed to provide formulations of elastomeric-product compositions for actual service.

## 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 37, *Rubber, vulcanized or thermoplastic — Determination of tensile stress-strain properties*

ISO 48, *Rubber, vulcanized or thermoplastic — Determination of hardness (hardness between 10 IRHD and 100 IRHD)*

ISO 1817, *Rubber, vulcanized or thermoplastic — Determination of the effect of liquids*

ISO 2230, *Rubber products — Guidelines for storage*

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

ISO 2781:2008, *Rubber, vulcanized or thermoplastic — Determination of density*

ISO 7619-1, *Rubber, vulcanized or thermoplastic — Determination of indentation hardness — Part 1: Durometer method (Shore hardness)*

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

### 3 Composition

The materials shall be homogeneous mixes of the ingredients shown in the relevant annex, in the proportions shown, weighed to the accuracy required in ISO 2393.

All ingredients shall be of recognized rubber quality.

The identification of proprietary materials as suitable ingredients does not exclude the use of other materials that can be shown to meet the requirements of the standard.

If necessary, the quantity of carbon black or silica may be adjusted from one consignment of rubber to another to give properties within the limits specified in the annexes. If the filler content is adjusted, the details shall be declared.

### 4 Preparation

The mixing of compounds and the vulcanization of test sheets shall be carried out in accordance with ISO 2393 and, where appropriate, as modified by detailed conditions and procedures given in the annexes. If a mould release agent is needed, dry PTFE mould lubricant shall be used.

### 5 Description

The vulcanized test sheets shall be  $2\text{ mm} \pm 0,2\text{ mm}$  thick when measured in accordance with ISO 23529, and free from surface defects and internal voids when viewed with normally corrected vision.

A "lot" of sheets is those sheets made from a single batch of rubber mix vulcanized under the same conditions. A "pressing" is the quantity of sheets produced at the same time in a single moulding operation.

### 6 Test sheet properties

The test sheets shall be characterized by one or more physical properties in accordance with the relevant ISO standard test method. Properties and tolerances for individual materials are given in the annexes.

All sheets shall be tested for compliance with the requirements of Clause 5. A sample sheet from each lot shall be tested for the properties of interest for compliance with the tolerances given in the appropriate annex. The actual values obtained shall be reported.

The test pieces necessary for determining material properties shall be taken from the test sheets in such a way that a minimum distance from the edge of the sheets is maintained. Dumb-bell test pieces shall be taken with their longitudinal axis parallel to the direction of milling of the moulding blank.

### 7 Designation

Each elastomer shall be designated as ISO 13226 SRE-A/B where ISO 13226 is this International Standard, SRE is the abbreviation for standard reference elastomer and, after the hyphen, A is a set of code-letters designating the polymer type and B is a descriptor identifying the particular compound. Some designations end with the letter X to signify that additional test requirements apply.

## 8 Storage

Test sheets shall be stored in accordance with ISO 2230. After one year, they shall be retested or disposed of. If the sheets still meet the test requirements, they may be stored and used for a further year.

NOTE The yearly tests apply irrespective of the type of rubber used.

STANDARDSISO.COM : Click to view the full PDF of ISO 13226:2011

## Annex A (normative)

### Acrylic rubbers: SRE-ACM/1 and SRE-ACM/1X

#### A.1 Purpose

These SREs are representative of ACM materials such as are used, for instance, for parts in contact with petroleum products in the mechanical-engineering and automobile sectors.

They are used for the characterization of service fluids such as mineral oils, fuels, lubricants, hydraulic fluids, coolants and refrigerants with regard to their effect on vulcanized acrylic rubbers.

The changes in mass, volume, hardness, tensile strength and elongation at break of the SRE when in contact with the service fluid under specified conditions may be included as supplementary data in specifications for the fluid concerned.

#### A.2 Composition

**Table A.1 — Composition of the SRE**

Ingredients	Parts by mass
Acrylic rubber <sup>a</sup>	100,0
Stearic acid	1,0
Pentaerythrite stearate <sup>b</sup>	2,0
Octylated diphenylamine (ODPA) <sup>c</sup>	2,0
Carbon black, N550	65,0
Sodium stearate	4,0
Quaternary ammonium salts <sup>d</sup>	2,0
<b>Total</b>	<b>176,0</b>
<sup>a</sup> HyTemp 4051 <sup>®</sup> from Zeon Chemicals L.P., or equivalent. <sup>b</sup> Struktol WB 222 <sup>®</sup> from Schill & Seilacher, or equivalent. <sup>c</sup> Vulkanox OCD/SG <sup>®</sup> from Lanxess AG, or equivalent. <sup>d</sup> HyTemp NPC-50 <sup>®</sup> from Zeon Chemicals L.P., or equivalent.	

#### A.3 Recommended mixing procedure

Mix on a two-roll mill (see Clause A.6).

#### A.4 Vulcanization

Condition the sheeted compound at ambient temperature for 12 h to 48 h.

Press-cure at 180 °C ± 2 °C for 10 min ± 1 min.

Post-cure at 175 °C ± 2 °C for 4 h ± 0,5 h.

## A.5 Test sheet properties

SRE-ACM/1 material shall fulfil the basic property given in Table A.2. Additional properties may be specified (see Table A.3).

SRE-ACM/1X material shall fulfil both the basic and the additional properties.

**Table A.2 — Basic property**

Property	Unit	Requirement	Test method
Increase in mass in test liquid B for fuels, as in ISO 1817	%	26 to 29	ISO 1817, three type 2 dumb-bell test pieces <sup>a</sup> immersed for 22 h ± 0,5 h at 23 °C ± 2 °C Test piece/test liquid volume ratio: 1/(30 ± 5)
<sup>a</sup> As specified in ISO 37.			

**Table A.3 — Additional properties in the initial state**

Property	Unit	Requirement	Test method
Tensile strength	MPa	12 to 16	ISO 37, five type 2 dumb-bell test pieces
Elongation at break	%	140 to 220	
Hardness	Shore A	69 to 74	ISO 7619-1, three type 2 dumb-bell test pieces <sup>a</sup> , three plies
	IRHD	69 to 74	ISO 48, three type 2 dumb-bell test pieces <sup>a</sup> , three plies
Density	Mg/m <sup>3</sup>	1,30 to 1,34	ISO 2781:2008, method A, three test pieces
<sup>a</sup> As specified in ISO 37.			

## A.6 Mixing procedures

The mixing procedure given in Table A.4 can be used to produce SRE-ACM/1X material using a two-roll mill.

**Table A.4 — Mixing on a two-roll mill**

Roll diameter: 200 mm		
Working width: 395 mm		
Speed of rolls: 18/22 min <sup>-1</sup>		
Surface temperature of rolls: 70 °C ± 5 °C		
Mass of rubber: 650 g		
Mixing step	Elapsed time min	Nip opening mm
Band rubber	0	2,0 ± 0,5
Add stearic acid, pentaerythrite stearate, ODPa and carbon black	1	2,2 ± 0,5
Make 3/4 cuts (four from each side)	11	
Add sodium stearate and quaternary ammonium salts <sup>a</sup>	13	
Make 3/4 cuts (six from each side)	17	
Turn the rolled sheet (three times)	19	
Sheet off	21	
Final temperature of sheet: approx. 75 °C		
<sup>a</sup> It is recommended that some of the final cuts be made during the addition of the sodium stearate to finalize the mixing procedure so that the sheet is taken off not later than 6 min ± 1 min after the accelerator has been added.		

## Annex B (normative)

### Acrylonitrile-butadiene rubbers: SRE-NBR 28/P and SRE-NBR 28/PX

#### B.1 Purpose

These SREs are representative of peroxide-cured NBR materials such as are used, for instance, for parts in contact with petroleum products in the mechanical-engineering and automobile sectors.

They are used for the characterization of service fluids such as mineral oils, fuels, lubricants, hydraulic fluids, coolants and refrigerants with regard to their effect on vulcanized nitrile rubbers.

The changes in mass, volume, hardness, tensile strength and elongation at break of the SRE when in contact with the service fluid under specified conditions may be included as supplementary data in specifications for the fluid concerned.

#### B.2 Composition

**Table B.1 — Composition of the SRE**

Ingredients	Parts by mass
NBR with (28 ± 0,5) % by mass of acrylonitrile <sup>a</sup>	100,0
<i>N</i> -(1,3-dimethylbutyl)- <i>N'</i> -phenyl- <i>p</i> -phenylenediamine (6PPD) <sup>b</sup>	0,5
Zinc oxide, precipitated <sup>c</sup>	5,0
Carbon black, N550	70,0
Dicumyl peroxide (40 % by mass) <sup>d</sup>	3,0
<b>Total</b>	<b>178,5</b>
<sup>a</sup> Perbunan NT 2845 <sup>®</sup> from Lanxess AG, or equivalent. <sup>b</sup> Vulkanox 4020 <sup>®</sup> from Lanxess AG, or equivalent. <sup>c</sup> Zinkoxyd aktiv <sup>®</sup> from Lanxess AG, or equivalent. <sup>d</sup> Perkadox BC 40 <sup>®</sup> from Akzo Nobel Chemicals B.V., or equivalent.	

#### B.3 Recommended mixing procedure

Prepare a masterbatch without the peroxide in an internal mixer followed by homogenizing and addition of peroxide on a two-roll mill (see B.6.1).

Alternatively, mix completely on a two-roll mill (see B.6.2).

#### B.4 Vulcanization

Condition the sheeted compound at ambient temperature for 20 h to 24 h.

Press-cure at 170 °C ± 2 °C for 20 min ± 1 min.

## B.5 Test sheet properties

SRE-NBR 28/P material shall fulfil the basic property given in Table B.2. Additional properties may be specified (see Table B.3).

SRE-NBR 28/PX material shall fulfil both the basic and the additional properties.

**Table B.2 — Basic property**

Property	Unit	Requirement	Test method
Increase in mass in test liquid B for fuels, as in ISO 1817	%	25 to 28	ISO 1817, three type 2 dumb-bell test pieces <sup>a</sup> immersed for 22 h ± 0,5 h at 23 °C ± 2 °C Test piece/test liquid volume ratio: 1/(30 ± 5)
<sup>a</sup> As specified in ISO 37.			

**Table B.3 — Additional properties in the initial state**

Property	Unit	Requirement	Test method
Tensile strength	MPa	20 to 25	ISO 37, five type 2 dumb-bell test pieces
Elongation at break	%	170 to 220	
Hardness	Shore A	79 to 84	ISO 7619-1, three type 2 dumb-bell test pieces <sup>a</sup> , three plies
	IRHD	79 to 84	ISO 48, three type 2 dumb-bell test pieces <sup>a</sup> , three plies
Density	Mg/m <sup>3</sup>	1,21 to 1,25	ISO 2781:2008, method A, three test pieces
<sup>a</sup> As specified in ISO 37.			

## B.6 Mixing procedures

The following mixing procedures can be used to produce SRE-NBR 28/PX material:

### B.6.1 Internal mixer and subsequent two-roll mill

Table B.4 gives an example of a mixing procedure for preparation of a masterbatch without peroxide, using an internal mixer fitted with intermeshing blades:

**Table B.4 — Mixing on an internal mixer**

Mixing-chamber volume <sup>a</sup> : 4,6 dm <sup>3</sup>	
Chamber filled to: (65 ± 5) % <sup>b</sup>	
Speed <sup>c</sup> : 30 min <sup>-1</sup>	
Mass of rubber: 2 000 g	
Mixing step	Elapsed time min
Add rubber	0
Add zinc oxide and 6PPD	2
Add carbon black	3
Dump	8
Final temperature of masterbatch: approx. 120 °C	
<sup>a</sup> Determined by means of wheat grains.	
<sup>b</sup> When using a mixer fitted with non-intermeshing blades, fill the chamber to (80 ± 5) %.	
<sup>c</sup> With cooling fully operative.	

Table B.5 gives an example of a mixing procedure for homogenization of the masterbatch and addition of peroxide, using a two-roll mill:

**Table B.5 — Homogenization on a two-roll mill**

Roll diameter: 250 mm Working width: 410 mm Speed of rolls: 0 min to 13 min: 12/18 min <sup>-1</sup> 14 min to 25 min: 12/12 min <sup>-1</sup> Surface temperature of rolls: 50 °C ± 5 °C		
Mixing step	Elapsed time min	Nip opening mm
Band hot masterbatch	0	3,0 ± 0,5
Make 3/4 cuts (14 from each side)	1	
Turn the rolled sheet (eight times)	8	
Make 3/4 cuts (two from each side)	13	
Add peroxide	14	
Make 3/4 cuts (five from each side)	16	
Turn the rolled sheet (four times)	19	
Make 3/4 cuts (five from each side)	21	
Turn the rolled sheet (four times)	23	
Sheet off	25	1,5 ± 0,5
Final temperature of sheet: approx. 75 °C		

**B.6.2 Mixing using a two-roll mill**

**Table B.6 — Mixing on a two-roll mill**

Roll diameter: 200 mm Working width: 395 mm Speed of rolls: 20/24 min <sup>-1</sup> Surface temperature of rolls: 50 °C ± 5 °C Mass of rubber: 1 000 g		
Mixing step	Elapsed time min	Nip opening mm
Band rubber	0	2,0 ± 0,5
Add zinc oxide and 6PPD	3	
Make 3/4 cuts (three from each side)	4	
Add carbon black	6	Set stepwise to 3,2 ± 0,5
Add peroxide	17	
Make 3/4 cuts (six from each side)	19	3,2 ± 0,5
Turn the rolled sheet (four times)	23	
Sheet off	26	1,5 ± 0,5
Final temperature of sheet: approx. 90 °C		

## Annex C (normative)

### Acrylonitrile-butadiene rubbers: SRE-NBR 28/S, SRE-NBR 28/SX, SRE-NBR 34/S and SRE-NBR 34/SX

#### C.1 Purpose

These SREs are representative of low-sulfur-cured NBR materials such as are used, for instance, for parts in contact with petroleum products in the mechanical-engineering and automobile sectors.

They are used for the characterization of service fluids such as mineral oils, fuels, lubricants, hydraulic fluids, coolants and refrigerants with regard to their effect on vulcanized nitrile rubbers.

In order to cover a wide range of service fluids, two different acrylonitrile (ACN) contents and thus a different swelling behaviour of the SREs are specified as follows:

- ISO 13226 SRE-NBR 28/S and SRE-NBR 28/SX (ACN content 28 %);
- ISO 13226 SRE-NBR 34/S and SRE-NBR 34/SX (ACN content 34 %).

The changes in mass, volume, hardness, tensile strength and elongation at break of the SRE when in contact with the service fluid under specified conditions may be included as supplementary data in specifications for the fluid concerned.

#### C.2 Composition

**Table C.1 — Composition of the SRE**

Ingredients	Parts by mass	
	28/S 28/SX	34/S 34/SX
NBR (ACN content 28 %) <sup>a</sup>	100,0	—
NBR (ACN content 34 %) <sup>b</sup>	—	100,0
Polymerized 2,2,4-trimethyl-1,2-dihydroquinoline (TMQ) <sup>c</sup>	2,0	2,0
Zinc oxide, precipitated <sup>d</sup>	5,0	5,0
Stearic acid	1,0	1,0
Carbon black, N550	65,0	65,0
Tetramethylthiuram disulfide (TMTD) <sup>e</sup>	2,5	2,5
<i>N</i> -cyclohexylbenzothiazole-2-sulfenamide (CBS) <sup>f</sup>	1,5	1,5
Sulfur	0,2	0,2
<b>Total</b>	177,2	177,2
<sup>a</sup> Perbunan NT 2845 <sup>®</sup> from Lanxess AG, or equivalent. <sup>b</sup> Perbunan NT 3445 <sup>®</sup> from Lanxess AG, or equivalent. <sup>c</sup> Vulkanox HS/LG <sup>®</sup> from Lanxess AG, or equivalent. <sup>d</sup> Zinkoxyd aktiv <sup>®</sup> from Lanxess AG, or equivalent. <sup>e</sup> Vulkacit Thiuram/C <sup>®</sup> from Lanxess AG, or equivalent. <sup>f</sup> Vulkacit CZ/C <sup>®</sup> from Lanxess AG, or equivalent.		

### C.3 Recommended mixing procedures

Prepare a masterbatch without the TMTD, CBS and sulfur in an internal mixer followed by homogenizing and addition of vulcanizing ingredients on a two-roll mill (see C.6.1).

To obtain good sulfur dispersion, use about 10 % of the rubber to make a sulfur masterbatch.

Alternatively, mix completely on a two-roll mill. Prepare a sulfur masterbatch and store it for between 30 min and 24 h before proceeding with the main mix (see C.6.2).

### C.4 Vulcanization

Condition the sheeted compound at ambient temperature for 2 h to 24 h.

Press-cure at  $160\text{ °C} \pm 2\text{ °C}$  for  $20\text{ min} \pm 1\text{ min}$ .

### C.5 Test sheet properties

SRE-NBR 28/S and SRE-NBR 34/S materials shall fulfil the basic property given in Table C.2. Additional properties may be specified (see Table C.3).

SRE-NBR 28/SX and SRE-NBR 34/SX materials shall fulfil both the basic and the additional properties.

Table C.2 — Basic property

Property	Unit	Requirement		Test method
		28/S 28/SX	34/S 34/SX	
Increase in mass in test liquid B for fuels, as in ISO 1817	%	26 to 29	17 to 20	ISO 1817, three type 2 dumb-bell test pieces <sup>a</sup> immersed for $22\text{ h} \pm 0,5\text{ h}$ at $23\text{ °C} \pm 2\text{ °C}$ Test piece/test liquid volume ratio: $1/(30 \pm 5)$
<sup>a</sup> As specified in ISO 37.				

Table C.3 — Additional properties in the initial state

Property	Unit	Requirement		Test method
		28/SX	34/SX	
Tensile strength	MPa	20 to 25	20 to 25	ISO 37, five type 2 dumb-bell test pieces
Elongation at break	%	300 to 400	300 to 400	
Hardness	Shore A	76 to 81	77 to 82	ISO 7619-1, three type 2 dumb-bell test pieces <sup>a</sup> , three plies
	IRHD	76 to 81	77 to 82	ISO 48, three type 2 dumb-bell test pieces <sup>a</sup> , three plies
Density	Mg/m <sup>3</sup>	1,19 to 1,23	1,20 to 1,24	ISO 2781:2008, method A, three test pieces
<sup>a</sup> As specified in ISO 37.				

## C.6 Mixing procedures

The following mixing procedures can be used to produce SRE-NBR 28/SX and SRE-NBR 34/SX materials:

### C.6.1 Internal mixer and subsequent two-roll mill

Use an internal mixer to prepare a masterbatch, without the vulcanizing ingredients, e.g. as specified in Table C.4.

Homogenize using a two-roll mill and add the vulcanizing ingredients, e.g. as specified in Table C.5.

To achieve good sulfur dispersion, which is essential for good tensile characteristics, take about 10 % of the rubber and mix a sulfur batch, e.g. as specified in Table C.6 (with properly adjusted working width and/or nip opening). Add this batch on the two-roll mill during step 5 (i.e. the addition of sulfur, TMTD and CBS) in Table C.5.

**Table C.4 — Mixing procedure for preparation of a masterbatch without vulcanizing ingredients, using an internal mixer fitted with intermeshing blades (example)**

Mixing-chamber volume <sup>a</sup> : 4,6 dm <sup>3</sup> Chamber filled to: (65 ± 5) % <sup>b</sup> Speed <sup>c</sup> : 30 min <sup>-1</sup> Mass of rubber: 2 000 g	
Mixing step	Elapsed time min
Add rubber	0
Add zinc oxide and TMQ	2
Add carbon black and stearic acid	3
Dump	8
Final temperature of masterbatch: approx. 120 °C	
<sup>a</sup> Determined by means of wheat grains. <sup>b</sup> When using a mixer fitted with non-intermeshing blades, fill the chamber to (80 ± 5) %. <sup>c</sup> With cooling fully operative.	

**Table C.5 — Mixing procedure for homogenization of masterbatch and addition of vulcanizing ingredients, using a two-roll mill (example)**

Roll diameter: 250 mm Working width: 410 mm Speed of rolls: 0 min to 13 min: 12/18 min <sup>-1</sup> 14 min to 25 min: 12/12 min <sup>-1</sup> Surface temperature of rolls: 50 °C ± 5 °C		
Mixing step	Elapsed time min	Nip opening mm
Band hot masterbatch	0	3,0 ± 0,5
Make 3/4 cuts (14 from each side)	1	
Turn the rolled sheet (eight times)	8	
Make 3/4 cuts (two from each side)	13	
Add sulfur, TMTD and CBS	14	
Make 3/4 cuts (five from each side)	16	
Turn the rolled sheet (four times)	19	
Make 3/4 cuts (five from each side)	21	
Turn the rolled sheet (four times)	23	
Sheet off	25	1,5 ± 0,5
Final temperature of sheet: approx. 90 °C. To ensure homogeneous sulfur distribution, the final temperature shall not be less than 90 °C. Increase the surface temperature of the rolls if necessary.		

### C.6.2 Mixing using a two-roll mill

To ensure good sulfur distribution, prepare a sulfur batch (masterbatch) at a roll surface temperature of  $80\text{ °C} \pm 5\text{ °C}$ , e.g. as specified in Table C.6.

Store the sulfur batch for at least 30 min, but no longer than 24 h.

Continue the preparation of the mix at a roll surface temperature of  $50\text{ °C} \pm 5\text{ °C}$ , e.g. as specified in Table C.7.

**Table C.6 — Mixing procedure for preparation of a sulfur batch (example)**

Roll diameter: 200 mm		
Working width: 395 mm		
Speed of rolls: $20/24\text{ min}^{-1}$		
Surface temperature of rolls: $80\text{ °C} \pm 5\text{ °C}$		
Mass of rubber: 1 000 g		
Mixing step	Elapsed time min	Nip opening mm
Pass the rubber twice	0	$\leq 1$
Band rubber	1	$2,5 \pm 0,5$
Add sulfur	2	
Make 3/4 cuts (three from each side)	4	
Sheet off	7	
Final temperature of sheet: approx. $90\text{ °C}$ . To ensure homogeneous sulfur distribution, the final temperature shall not be less than $90\text{ °C}$ . Increase the surface temperature of the rolls if necessary.		

**Table C.7 — Mixing procedure for preparation of mix with sulfur batch (see Table C.6) (example)**

Roll diameter: 200 mm		
Working width: 395 mm		
Speed of rolls: $20/24\text{ min}^{-1}$		
Surface temperature of rolls: $50\text{ °C} \pm 5\text{ °C}$		
Mixing step	Elapsed time min	Nip opening mm
Pass sulfur batch between rolls	0	$2,5 \pm 0,5$
Add antioxidant	1	
Make 3/4 cuts (two from each side)	2,5	
Add zinc oxide	3	
Make 3/4 cuts (two from each side)	5	
Add carbon black and stearic acid	6	Set stepwise to $3,2 \pm 0,5$
Make 3/4 cuts (four from each side)	24	$3,2 \pm 0,5$
Add TMTD and CBS	26	
Make 3/4 cuts (six from each side)	28	
Turn the rolled sheet (four times)	30	
Sheet off	31	$1,5 \pm 0,5$
Final temperature of sheet: approx. $90\text{ °C}$		

## Annex D (normative)

### Acrylonitrile-butadiene rubbers: SRE-NBR/M

#### D.1 Purpose

Representative of a medium-nitrile-content acrylonitrile-butadiene rubber.

#### D.2 Composition

Table D.1 — Composition of the SRE

Ingredients	Parts by mass
Medium-acrylonitrile-butadiene rubber (ACN content 33 %) <sup>a</sup>	100
Stearic acid	1,5
Zinc oxide	5,0
Carbon black, N550	55 (nominal)
Sulfur masterbatch <sup>b</sup>	1,9
Dibenzothiazyl disulfide (MBTS) masterbatch <sup>c</sup>	2,0
<b>Total</b>	<b>165,4</b>
<p><sup>a</sup> Medium-nitrile-content, hot-polymerized Breon N33H80 from Zeon Chemicals Europe Ltd. has been found to be suitable.</p> <p><sup>b</sup> Masterbatch containing 80 % sulfur and 20 % of an ethylene-propylene rubber.</p> <p><sup>c</sup> Masterbatch containing 75 % active accelerator and 25 % of an ethylene-propylene rubber.</p>	

#### D.3 Vulcanization

Press-cure at 155 °C ± 2 °C for 20 min ± 15 s.

#### D.4 Test sheet properties

SRE-NBR/M material shall fulfil the basic property given in Table D.2.

Table D.2 — Basic property

Property	Unit	Requirement	Test method
Increase in volume in test liquid 101, as in ISO 1817	%	19 to 22	ISO 1817, three test pieces immersed for $(48_{-1}^0)$ h at 100 °C ± 1 °C Test piece/test liquid volume ratio: 1/(30 ± 5)

No additional properties are specified.

## Annex E (normative)

### Acrylonitrile-butadiene rubbers: SRE-NBR/L

#### E.1 Purpose

Representative of a low-nitrile-content acrylonitrile-butadiene rubber.

#### E.2 Composition

**Table E.1 — Composition of the SRE**

Ingredients	Parts by mass
Low-acrylonitrile-butadiene rubber (ACN content 18 %) <sup>a</sup>	100
Stearic acid	1,5
Zinc oxide	5,0
Carbon black, N550	67 (nominal)
Sulfur masterbatch <sup>b</sup>	1,9
Dibenzothiazyl disulfide (MBTS) masterbatch <sup>c</sup>	2,0
<b>Total</b>	<b>177,4</b>
<sup>a</sup> Low-nitrile-content Paracril 18.80 from Uniroyal Chemical has been found to be suitable. <sup>b</sup> Masterbatch containing 80 % sulfur and 20 % of an ethylene-propylene rubber. <sup>c</sup> Masterbatch containing 75 % active accelerator and 25 % of an ethylene-propylene rubber.	

#### E.3 Vulcanization

Press-cure at 155 °C ± 2 °C for 30 min ± 15 s.

#### E.4 Test sheet properties

SRE-NBR/L material shall fulfil the basic property given in Table E.2.

**Table E.2 — Basic property**

Property	Unit	Requirement	Test method
Increase in volume in test liquid B, as in ISO 1817	%	53 to 59	ISO 1817, three test pieces immersed for (48 <sup>0</sup> <sub>-1</sub> ) h at 40 °C ± 1 °C Test piece/test liquid volume ratio: 1/(30 ± 5)

No additional properties are specified.

## Annex F (normative)

### Chlorobutyl rubbers: SRE-CIIR/1

#### F.1 Purpose

Representative of a chlorobutyl rubber.

#### F.2 Composition

Table F.1 — Composition of the SRE

Ingredients	Parts by mass
Chlorobutyl rubber <sup>a</sup>	100
Stearic acid	1,5
Zinc oxide	5,0
Carbon black, N550	25 (nominal)
Carbon black, N990 <sup>b</sup>	120 (nominal)
Sulfur masterbatch <sup>c</sup>	0,94
Petroleum jelly	3,0
Mineral oil <sup>d</sup>	3,0
Dibenzothiazyl disulfide (MBTS) masterbatch <sup>e</sup>	2,0
<b>Total</b>	<b>260,44</b>
<p><sup>a</sup> Chlorobutyl 1066 from Exxon has been found to be suitable.</p> <p><sup>b</sup> Thermax MT N990<sup>®</sup> from Evonik Carbon Black GmbH, or equivalent.</p> <p><sup>c</sup> Masterbatch containing 80 % sulfur and 20 % of an ethylene-propylene rubber.</p> <p><sup>d</sup> UK DEF STAN 91-44, or equivalent.</p> <p><sup>e</sup> Masterbatch containing 75 % active accelerator and 25 % of an ethylene-propylene rubber.</p>	

#### F.3 Vulcanization

Press-cure at 155 °C ± 2 °C for 25 min ± 15 s.

#### F.4 Test sheet properties

SRE-CIIR/1 material shall fulfil the basic property given in Table F.2.

Table F.2 — Basic property

Property	Unit	Requirement	Test method
Increase in volume in test liquid B, as in ISO 1817	%	93 to 100	ISO 1817, three test pieces immersed for (48 <sub>-1</sub> <sup>0</sup> ) h at 40 °C ± 1 °C Test piece/test liquid volume ratio: 1/(30 ± 5)

No additional properties are specified.

## Annex G (normative)

### Chloroprene rubbers: SRE-CR/1

#### G.1 Purpose

Representative of a general-purpose chloroprene rubber.

#### G.2 Composition

Table G.1 — Composition of the SRE

Ingredients	Parts by mass
Chloroprene rubber <sup>a</sup>	100
Stearic acid	0,5
Lightly calcined magnesia	4,0
Zinc oxide	5,0
Carbon black, N550	17 (nominal)
Organic accelerator <sup>b</sup>	0,75 to 1,0
Antioxidant <sup>c</sup>	2,0
<b>Total</b>	129,25 to 129,5
<p><sup>a</sup> A slow-crystallizing form of a non-sulfur-modified polymer is required. Neoprene WRT from DuPont Performance Elastomers has been found to be suitable.</p> <p><sup>b</sup> Vulkacit CRV from Lanxess AG has been found to be suitable.</p> <p><sup>c</sup> Octamine from Uniroyal Chemical has been found to be suitable.</p>	

#### G.3 Vulcanization

Press-cure at 160 °C ± 2 °C for 25 min ± 15 s.

#### G.4 Test sheet properties

SRE-CR/1 material shall fulfil the basic property given in Table G.2.

Table G.2 — Basic property

Property	Unit	Requirement	Test method
Increase in volume in test liquid C, as in ISO 1817	%	76 to 84	ISO 1817, three test pieces immersed for $(48_{-1}^0)$ h at 40 °C ± 1 °C Test piece/test liquid volume ratio: 1/(30 ± 5)

No additional properties are specified.

## Annex H (normative)

### Ethylene-propylene rubbers: SRE-EPM/1

#### H.1 Purpose

Representative of a general-purpose ethylene-propylene rubber.

#### H.2 Composition

**Table H.1 — Composition of the SRE**

Ingredients	Parts by mass
Ethylene-propylene copolymer rubber <sup>a</sup>	100
Stearic acid	0,5
Carbon black, N550	55 (nominal)
Sulfur masterbatch <sup>b</sup>	0,44
1,1-dibutylperoxy-3,3,5-trimethylcyclohexane (40 % active powder) <sup>c</sup>	7,5
<b>Total</b>	<b>163,44</b>
<sup>a</sup> Keltan 3300 A ethylene-propylene copolymer from DSM Elastomers has been found to be suitable. <sup>b</sup> Masterbatch containing 80 % sulfur and 20 % of an ethylene-propylene rubber. <sup>c</sup> Trigonox 29-40 BPD from Akzo Nobel Chemicals B.V. has been found to be suitable.	

#### H.3 Vulcanization

Press-cure at 140 °C ± 2 °C for 10 min ± 15 s.

#### H.4 Test sheet properties

SRE-EPM/1 material shall fulfil the basic property given in Table H.2.

**Table H.2 — Basic property**

Property	Unit	Requirement	Test method
Increase in volume in test liquid 103 (tri- <i>n</i> -butyl phosphate: density at 20 °C 0,975 Mg/m <sup>3</sup> to 0,976 Mg/m <sup>3</sup> ), as in ISO 1817	%	20 to 23	ISO 1817, three test pieces immersed for 22 h ± 0,25 h at 100 °C ± 1 °C Test piece/test liquid volume ratio: 1/(30 ± 5)

No additional properties are specified.

## Annex I (normative)

### Fluoropolymer rubbers: SRE-FKM/1

#### I.1 Purpose

Representative of a fluoropolymer rubber.

#### I.2 Composition

Table I.1 — Composition of the SRE

Ingredients	Parts by mass
Fluoropolymer rubber <sup>a</sup>	100
Lightly calcined magnesia <sup>b</sup>	3,0
Carbon black, N990 <sup>c</sup>	30 (nominal)
2,5-di( <i>t</i> -butylperoxy)-2,5-dimethylhexane (50 % active powder)	3,0
Coagent <sup>d</sup>	3,0
<b>Total</b>	139,0
<sup>a</sup> Viton type GF from DuPont Performance Elastomers has been found to be suitable. <sup>b</sup> Maglite D from Elastochem has been found to be suitable. <sup>c</sup> Thermax MT N990 <sup>®</sup> from Evonik Carbon Black GmbH, or equivalent. <sup>d</sup> Triallyl isocyanurate has been found to be suitable for use with Viton type GF.	

#### I.3 Vulcanization

Press-cure at 175 °C ± 2 °C for 8 min ± 15 s.

Post-cure at 230 °C ± 5 °C for (24<sup>0</sup><sub>-2</sub>) h.

#### I.4 Test sheet properties

SRE-FKM/1 material shall fulfil the basic property given in Table I.2.

Table I.2 — Basic property

Property	Unit	Requirement	Test method
Increase in volume in test liquid E (toluene), as in ISO 1817	%	8 to 11	ISO 1817, three test pieces immersed for 168 h ± 2 h at 40 °C ± 1 °C Test piece/test liquid volume ratio: 1/(30 ± 5)

No additional properties are specified.

## Annex J (normative)

### Fluoropolymer rubbers: SRE-FKM/2X

#### J.1 Purpose

This SRE is representative of FKM materials such as are used, for instance, for parts in contact with petroleum products in the mechanical-engineering and automobile sectors.

It is used for the characterization of service fluids such as mineral oils, fuels, lubricants, hydraulic fluids, coolants and refrigerants with regard to their effect on vulcanized fluoropolymer rubbers.

The changes in mass, volume, hardness, tensile strength and elongation at break of the SRE when in contact with the service fluid under specified conditions may be included as supplementary data in specifications for the fluid concerned.

#### J.2 Composition

Table J.1 — Composition of the SRE

Ingredients	Parts by mass
Vinylidene fluoride-hexafluoropropylene copolymer <sup>a</sup>	100,0
Magnesium oxide, high activity <sup>b</sup>	3,0
Calcium hydroxide, small particle size <sup>c</sup>	2,0
Carbon black, N990 <sup>d</sup>	25,0
Accelerator: organic phosphonium salt, e.g. triphenylbenzyl-phosphonium chloride <sup>e</sup>	0,44
Crosslinking agent: 2,2-bis(4-hydroxyphenyl)-hexafluoropropane (bisphenol AF) <sup>f</sup>	1,35
<b>Total</b>	<b>131,79</b>
<p>The following brands have been found to be suitable:</p> <p><sup>a</sup> Tecnoflon N935<sup>®</sup> from Solvay, or Viton A 500<sup>®</sup> from Du Pont Performance Elastomers or Fluorel FC 2230<sup>®</sup> from Dyneon.</p> <p><sup>b</sup> Elastomag 170<sup>®</sup> from Nordmann, Rassmann GmbH &amp; Co.</p> <p><sup>c</sup> Calcium hydroxide from Boehringer AG.</p> <p><sup>d</sup> Thermax MT N990<sup>®</sup> from Evonik Carbon Black GmbH.</p> <p><sup>e</sup> Intercure 2<sup>®</sup> from Interbusiness, Milan, or an equivalent accelerator such as Tecnoflon FOR M2 from Ausimont. In the latter case, take 1,45 phr of Tecnoflon FOR M2 and reduce the amount of rubber by 1 g per 100 g of rubber.</p> <p><sup>f</sup> 2,2-Bis(4-hydroxyphenyl)-hexafluoropropane from Hoechst, or an equivalent crosslinking agent such as Tecnoflon FOR M1 from Ausimont or Viton Curative No. 30 from DuPont Performance Elastomers. The bisphenol content of both of the last two agents is 50 %, so take 2,76 phr and reduce the amount of rubber by 1 g per 100 g of rubber.</p>	

#### J.3 Recommended mixing procedure

A two-roll mill can be used (see Clause J.6).

#### J.4 Vulcanization

Condition the sheeted compound at ambient temperature for about 24 h. Prepare samples for cure. The direction of milling shall be marked on the slabs.

Press-cure at  $180\text{ °C} \pm 2\text{ °C}$  for  $7\text{ min} \pm 1\text{ min}$ .

Post-cure at  $220\text{ °C} \pm 5\text{ °C}$  for  $16\text{ h} \pm 2\text{ h}$  in an air-circulating oven with air exchange.

#### J.5 Test sheet properties

SRE-FKM/2X material shall fulfil both the basic and the additional properties given in Tables J.2 and J.3.

**Table J.2 — Basic property**

Property	Unit	Requirement	Test method
Increase in mass in test liquid C for fuels, as in ISO 1817	%	6 to 9	ISO 1817, three type 2 dumb-bell test pieces <sup>a</sup> immersed for $(24 \pm \frac{b}{2})$ h at $60\text{ °C} \pm 2\text{ °C}$ Test piece/test liquid volume ratio: $1/(30 \pm 5)$
<sup>a</sup> As specified in ISO 37.			

**Table J.3 — Additional properties in the initial state**

Property	Unit	Requirement	Test method
Tensile strength	MPa	12 to 18	ISO 37, five type 2 dumb-bell test pieces
Elongation at break	%	250 to 330	
Hardness	IRHD	67 to 73	ISO 48, microtest, three test pieces
Density	Mg/m <sup>3</sup>	1,83 to 1,87	ISO 2781:2008, method A, three test pieces

## J.6 Mixing procedures

The mixing procedure given in Table J.4 can be used to produce SRE-FKM/2X material using a two-roll mill.

**Table J.4 — Mixing on a two-roll mill**

Roll diameter: 150 mm		
Working width: 350 mm		
Speed of rolls: 18/22 min <sup>-1</sup>		
Surface temperature: 50 °C ± 5 °C		
Mass of rubber: 1 000 g		
Mixing step	Elapsed time min	Nip opening mm
Band rubber (first ingredient in Table J.1)	0	2,0 ± 0,5
Blend the other ingredients (2 to 6) together and add evenly across the rolls at a constant rate Make 3/4 cuts (four from each side)	2	3,0 ± 1,0
Turn the rolled sheet (six times)	12	0,5 ± 0,2
Sheet off	18	2,2 ± 0,5
Final temperature of sheet: 60 °C to 70 °C		
Refine mixed stock after about 24 h storage on the mill under the same conditions, as follows:		
Band batch	0	3,0 ± 1,0
Make 3/4 cuts (three from each side) Turn the rolled sheet (six times)	2	0,5 ± 0,2
Sheet off	8	2,2 ± 0,5

## Annex K (normative)

### Hydrogenated acrylonitrile-butadiene rubbers: SRE-HNBR/1 and SRE-HNBR/1X

#### K.1 Purpose

These SREs are representative of saturated HNBR materials such as are used, for instance, for parts in contact with petroleum products in the mechanical-engineering and automobile sectors.

They are used for the characterization of service fluids such as mineral oils, fuels, lubricants, hydraulic fluids, coolants and refrigerants with regard to their effect on vulcanized fully hydrogenated nitrile rubbers.

The changes in mass, volume, hardness, tensile strength and elongation at break of the SRE when in contact with the service fluid under specified conditions may be included as supplementary data in specifications for the fluid concerned.

#### K.2 Composition

**Table K.1 — Composition of the SRE**

Ingredients	Parts by mass
HNBR (ACN content 39 %) <sup>a</sup>	100,0
Zinc oxide, precipitated <sup>b</sup>	2,0
Magnesium oxide <sup>c</sup>	2,0
Styrenated diphenylamine (SDPA) <sup>d</sup>	1,0
Carbon black, N550	50,0
Triallyl isocyanurate (TAIC) <sup>e</sup>	1,5
1,3-bis( <i>t</i> -butylperoxyisopropyl)-benzene (40 % by mass) <sup>f</sup>	7,5
<b>Total</b>	<b>164,0</b>
<sup>a</sup> Therban A 3907 <sup>®</sup> from Lanxess AG, or equivalent. <sup>b</sup> Zinkoxyd aktiv <sup>®</sup> from Lanxess AG, or equivalent. <sup>c</sup> Maglite DE <sup>®</sup> from Merck & Co, or equivalent. <sup>d</sup> Vulkanox DDA <sup>®</sup> from Lanxess AG, or equivalent. <sup>e</sup> Perkalink 301 <sup>®</sup> from Akzo Nobel Chemicals B.V., or equivalent. <sup>f</sup> Perkadox 14-40 <sup>®</sup> from Akzo Nobel Chemicals B.V., or equivalent.	

#### K.3 Recommended mixing procedure

Prepare a masterbatch without the peroxide in an internal mixer followed by homogenizing and addition of peroxide on a two-roll mill (see K.6.1).

Alternatively, sheet off the masterbatch on a two-roll mill, cut into strips, cool and re-introduce into the internal mixer and add peroxide (see K.6.2).