



**SLOVENSKI STANDARD**  
**SIST ISO 4611:1996**

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**Polimerni materiali - Ugotavljanje vplivov vlažne toplote, vodne prhe in slane meglice**

Plastics -- Determination of the effects of exposure to damp heat, water spray and salt mist

**iTeh STANDARD PREVIEW**

Plastiques -- Détermination des effets d'une exposition à la chaleur humide, au brouillard d'eau et au brouillard salin

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**Ta slovenski standard je istoveten z: ISO 4611:1987**

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**ICS:**

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# INTERNATIONAL STANDARD

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

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## Plastics — Determination of the effects of exposure to damp heat, water spray and salt mist

*Plastiques — Détermination des effets d'une exposition à la chaleur humide, au brouillard  
d'eau et au brouillard salin*

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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.

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. They are approved in accordance with ISO procedures requiring at least 75 % approval by the member bodies voting.

International Standard ISO 4611 was prepared by Technical Committee ISO/TC 61, *Plastics*.

This second edition cancels and replaces the first edition (ISO 4611:1980), of which it constitutes a minor revision.

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Users should note that all International Standards undergo revision from time to time and that any reference made herein to any other International Standard implies its latest edition, unless otherwise stated.

# Plastics — Determination of the effects of exposure to damp heat, water spray and salt mist

## 0 Introduction

**0.1** Various test methods are available for the exposure of plastics to different aggressive agents acting in a combined and simultaneous fashion, such as natural weathering. Other test methods are available for the purpose of a separate evaluation of the action of individual aggressive agents. Among the latter there are, for example, tests for the resistance to specific chemicals and to radiations of a definite spectral range.

For some applications, it may be desirable to evaluate the behaviour of the materials in a hot damp atmosphere just below the saturation limit of water vapour, as well as in the presence of the liquid phase.

In these conditions not only water absorption or leaching of some ingredients of the composition may be observed but also degradation phenomena due to hydrolysis, exudation of plasticizers, etc.

It may also be desirable sometimes to evaluate the behaviour of materials in the presence of a highly corrosive electrolyte, such as a sodium chloride solution (salt mist), which is the principal aggressive agent present in marine environments and of particular importance in the case of nautical applications. It is well known that sodium chloride has no noticeable action on the polymers that are the basic components of plastics, and that salt solutions, owing to their higher osmotic pressure, are normally absorbed by plastics to a lesser degree than pure water, but it cannot be assumed *a priori* that they have no action on composite materials, containing fillers, reinforcing components or pigments, for instance.

Furthermore, the evaluation of the effect of salt mist can be very important for finished or semi-finished articles which, while basically consisting of plastic materials, do contain some metallic elements, such as moulded-in inserts, thin laminated foils, surface coatings applied by electro-plating or other procedures, or, lastly, metal cores sheathed with plastics by extrusion or by dipping in pastes or fluidized-bed powders.

**0.2** Methods and equipment for obtaining reproducible aggressive environments of the above types are well known

and have been described by International Standards relevant to other materials and IEC (International Electrotechnical Commission) standards relevant to electrical and electronic components. The same equipment and procedures described in these standards can also be employed for plastics, with appropriate care and adjustments.

**0.3** The present International Standard is intended to provide general guidance only, on the choice of suitable equipment and procedures for obtaining the exposure conditions described above and for the preparation of test specimens. It also only gives general guidance on the properties to be evaluated. Specific details are given in the various ISO and IEC Publications.

For the expression of results, the present International Standard follows, as far as possible, the same criteria adopted in the existing test methods for the exposure to chemicals (see ISO 175) and to natural weathering or artificial light (see ISO 4582).

**0.4** These tests are intended to yield data about the effects of the described exposures on the materials; however, a direct correlation between the experimental results and the behaviour in service is not to be inferred.

## 1 Scope and field of application

**1.1** This International Standard specifies the exposure conditions of plastics to

- damp heat,
- water spray,
- salt mist,

and the methods for the evaluation of the change of some significant characteristics after given exposure stages.

**1.2** This International Standard is, in general, suitable for all plastics in the form of standard test specimens, and finished articles or parts thereof.

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1.3 This International Standard considers separately methods for the determination of

- change of mass,
- change of dimensions and appearance,
- change of physical properties.

## 2 References

ISO 175, *Plastics — Determination of the effects of liquid chemicals including water.*

ISO 293, *Plastics — Compression moulding test specimens of thermoplastic materials.*

ISO 294, *Plastics — Injection moulding test specimens of thermoplastic materials.*

ISO 295, *Plastics — Compression moulding test specimens of thermosetting materials.*

ISO 2557-1, *Plastics — Amorphous thermoplastic moulding materials — Preparation of test specimens with a defined level of shrinkage — Part 1: Test specimens in the form of parallelepipedic bars (Injection moulding and compression moulding).*

ISO 2557-2, *Plastics — Amorphous thermoplastics — Preparation of test specimens with a specified reversion — Part 2: Plates.*

ISO 2818, *Plastics — Preparation of test specimens by machining.*

ISO 3205, *Preferred test temperatures.*

ISO 3768, *Metallic coatings — Neutral salt spray test (NSS test).*

ISO 4582, *Plastics — Determination of changes in colour and variation in properties after exposure to daylight under glass, natural weathering or artificial light.*

IEC Publication 68, *Basic environmental testing procedures:*

68-2-3, *Test Ca: Damp heat, steady state.*

68-2-11, *Test Ka: Salt mist.*

68-2-30, *Test Db: Damp heat, cyclic (12 + 12-hour cycle).*

68-2-38, *Test Z/AD: Composite temperature/humidity cyclic test.*

## 3 Principle

Determination of one or more properties before and after given periods of exposure in the specified environmental conditions, and observation of any change in appearance. If required, the determination of one or more properties may be carried out after exposure and a subsequent drying treatment or a recondi-

tioning treatment carried out with the aim of obtaining the same state of equilibrium with atmospheric humidity as that of the initial specimens.

## 4 General test conditions

### 4.1 Environmental conditions and equipment

#### 4.1.1 Damp heat

The preferred test conditions are those described by the IEC Publications referred to in 4.1.1.1 and 4.1.1.2. Different temperature and/or humidity conditions may be specified, however, in the relevant product specifications or by agreement between the interested parties.

##### 4.1.1.1 Steady state test

A suitable test method is specified in IEC Publication 68-2-3.

The following conditions are specified:

Temperature:  $40 \pm 2 \text{ }^\circ\text{C}$

Relative humidity:  $(93 \pm \frac{2}{3}) \%$

##### 4.1.1.2 Cyclic test

If a cyclic test is required, the conditions specified in IEC Publication 68-2-30 can be adopted.

This Publication specifies 12 h + 12 h cycles with temperature change from:

$25 \pm 3 \text{ }^\circ\text{C}$

to a)  $40 \pm 2 \text{ }^\circ\text{C}$

b)  $55 \pm 2 \text{ }^\circ\text{C}$

Relative humidity is maintained at  $(93 \pm 3) \%$  at the higher temperature and not less than 95 % during the rest of the cycle.

NOTE — For composite temperature/humidity cycles, with the addition of a number of excursions to sub-zero temperatures, some guidance can be found in IEC Publication 68-2-38.

#### 4.1.2 Water spray

The main difference between this exposure condition and that for damp heat/steady state (see 4.1.1) is the constant presence of the liquid phase, in the form of small water droplets.

Suitable equipment for obtaining these conditions is substantially identical to that for the salt mist exposure (see 4.1.3) and is described in the relevant specifications.

Distilled or deionized water, having a pH between 6 and 7, shall be used instead of the salt solution.

The temperature in the test enclosure shall be  $40 \pm 2 \text{ }^\circ\text{C}$ .

### 4.1.3 Salt mist

Suitable equipment and procedure for the exposure to salt mist (or salt spray) are described in ISO 3768.

The general operating conditions are as follows:

Temperature inside the cabinet .....  $35 \pm 2$  °C

Sodium chloride solution:

- concentration .....  $50 \pm 5$  g/l
- pH ..... 6,5 to 7,2

The solution is prepared by dissolving sodium chloride of recognized analytical grade in distilled or deionized water.

The purity requirements are specified in ISO 3768.

Quantity of "mist" collected over a period of 24 h on a horizontal collecting area of 8 000 mm<sup>2</sup> ..... 1 to 2 ml/h

These conditions comply also with IEC Publication 68-2-11.

NOTE — In this test, the 35 °C temperature value has been retained, even though this value is not included among those recommended in ISO 3205, because it is specified in the ISO 3768 test referred to and by the majority of existing national standards.

### 4.2 Duration of tests

The test duration shall be as specified in the relevant standard or as agreed upon by the interested parties, with reference to the intended application.

It is recommended that the time periods be chosen from the following standard scale:

24, 48, 96, 144, 168 h

and, for long periods:

1, 2, 4, 8, 16, 26, 52, 78 weeks.

### 4.3 Test specimens

See 5.2, 6.2 and 7.2.

### 4.4 Conditioning

Unless otherwise agreed by the interested parties, the test specimens shall be conditioned before testing for at least 86 h at  $23 \pm 2$  °C and  $(50 \pm 5)$  % r.h.

NOTE — For certain materials, which are known to approach rapidly, or on the contrary very slowly, the state of equilibrium of temperature and, above all, of humidity, shorter or longer conditioning periods may be specified in the particular specifications concerning them (see the annex).

### 4.5 Treatment after exposure

The exposed specimens shall be tested either

- a) directly after the exposure, or

- b) after the exposure and subsequent drying or reconditioning

The first procedure shall be used when it is required to know the state of the material while it still contains the amount of water it had absorbed by the end of the exposure. The second procedure shall be used when it is required to determine the changes in properties of the material as a result of the exposure only. In the case of reconditioning, the specimens shall be brought back, as far as possible, to the same state as the initial state before exposure with regard to equilibrium with atmospheric humidity (see 4.4).

#### 4.5.1 Test after exposure only

After rinsing with distilled or deionized water, if necessary, and wiping dry, the exposed specimens shall be brought to  $23 \pm 2$  °C in a closed container; usually 4 h are sufficient for this purpose.

#### 4.5.2 Test after exposure and drying or reconditioning

After rinsing and wiping dry, the specimens shall be dried or reconditioned to equilibrium with the same atmospheric conditions as before the exposure (see 4.4), taking due account of the procedures described in A.3.1 and A.3.2 of the annex. Unless otherwise specified in the relevant product standard or agreed upon between the interested parties, the specimens shall be dried in an oven at  $50 \pm 2$  °C for 24 h and cooled to  $23 \pm 2$  °C in a desiccator.

## 5 Change of mass

### 5.1 General

5.1.1 In this type of test, the changes of mass are, at least partially, due to water absorption, and they are, therefore, particularly influenced by the conditioning and drying/reconditioning of the test specimens.

It is important, therefore, that the precise conditions of test be specified in the relevant product specifications.

Normally the specimens are weighed immediately after exposure, rinsing and wiping dry, or are treated as described in 4.5.1.

If the determination of mass after drying or reconditioning is required, the specimens shall be dried or reconditioned as specified in 4.5.2.

NOTE — This test is not applicable to cellular materials.

5.1.2 The changes of mass are generally proportional to the surface area of the test specimens, but are influenced by their thickness.

IMPORTANT — It is emphasized that the comparison of different plastics by means of this test is valid only if the test specimens used are of the same shape, the same dimensions and in as nearly as possible the same state (of surface, internal stresses, etc.).

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## 5.2 Test specimens

The test specimens may be obtained directly by moulding or by machining. In the latter case, the cut surfaces shall be smooth and shall not show any trace of charring that may be due to the method of preparation.

## 5.2.1 Moulding and extrusion compounds

The test specimens shall be in the form of a square of  $50 \pm 1$  mm side and  $3 \pm 0,2$  mm thick. Rectangular specimens having the same surface area (for example  $100 \text{ mm} \times 25 \text{ mm}$ , i.e.  $2\,500 \text{ mm}^2$ ) may be used.

The specimens may be cut from a sheet of the same thickness, or prepared by compression or injection moulding or by extrusion under the conditions given in the relevant specification for the material or under the conditions prescribed by the supplier of the material.

Moulding materials may be, alternatively, directly moulded in a mould of the prescribed dimensions.

The state of internal stress, if necessary, shall be determined according to ISO 2557-1 and/or ISO 2557-2, as applicable.

NOTE — The general principles for preparing moulded and machined test specimens are the subject of the following International Standards: ISO 293, ISO 294, ISO 295, ISO 2557-1, ISO 2557-2, ISO 2818.

## 5.2.2 Sheet

The specimens shall be  $50 \pm 1$  mm square or rectangular, having the same surface area, and shall be cut from the sheet under test.

If the nominal thickness of the sheet under test is equal to or less than 25 mm, the thickness of the test specimen shall be the same as that of the sheet.

If the nominal thickness is greater than 25 mm and in the absence of special provisions in the relevant specification, the thickness of the test specimen shall be reduced to 25 mm by machining on one surface only.

The machined surface shall not be directly exposed to water or salt spray.

A complete description of the machining, if any, shall be included in the test report.

## 5.2.3 Semi-finished and finished products (other than sheets)

The specimens shall be as similar in shape and size as possible to the specimens described in 5.2.1, and be prepared according to the product specification or as agreed upon between the interested parties.

If any machining is necessary, a complete description thereof shall be included in the test report.

## 5.2.4 Number of test specimens

At least three specimens shall be tested.

## 5.3 Conditioning

See 4.4.

## 5.4 Procedure

5.4.1 Determine the mass ( $m_1$ ) of each specimen to the nearest 0,001 g.

5.4.2 Expose the specimens to the test environment chosen from those listed in clause 4.

5.4.3 Rinse, if necessary (for example in the case of exposure to salt mist), and wipe the specimens dry.

5.4.4 Immediately determine the mass ( $m_2$ ) of each specimen to the nearest 0,001 g.

5.4.5 Dry or recondition the specimens, if required, according to 4.5.2 and determine the mass ( $m_3$ ) of each specimen to the nearest 0,001 g.

## 5.5 Expression of results

5.5.1 The change of mass per unit area, in grams per square metre, is given by the formula

$$\frac{m_2 - m_1}{S} \quad \text{or} \quad \frac{m_3 - m_1}{S} \quad \text{respectively}$$

where

$m_1$ ,  $m_2$  and  $m_3$  are as defined in 5.4.1, 5.4.4 and 5.4.5;

$S$  is the initial total surface area (including the edges of the specimens) in square metres.

5.5.2 The change of mass, as a percentage, is given by the formula

$$\frac{m_2 - m_1}{m_1} \times 100 \quad \text{or} \quad \frac{m_3 - m_1}{m_1} \times 100 \quad \text{respectively.}$$

The change is positive for increase of mass and negative for reduction of mass.

5.5.3 Calculate the mean value of the test results for all the specimens from a given sample.

## 6 Change of dimensions and appearance

## 6.1 General

Change of dimensions may result from either volume changes due to water absorption or leaching of some components, or to relaxation of internal moulded-in stresses, or both.



It is important, therefore, that the precise conditions of test be specified in the relevant product specifications.

For anisotropic materials, such as calendered or extruded sheets or extruded rods, the changes in linear dimensions in the machine direction (lengthwise) and in the transverse direction (crosswise) may be different; it is therefore necessary to determine the changes in both directions.

Where it is desired to differentiate between the effect of the relaxation of moulded-in stresses and that of the action of water, the test may also be performed using a set of annealed specimens.

## 6.2 Test specimens

Prepare the test specimens as described in 5.2. For anisotropic materials, the sides shall be parallel to the machine and transverse directions, respectively (see 6.1).

NOTE — The dimensional measurements may be made on the same specimens used for the determination of the changes in mass, immediately after weighing them.

## 6.3 Conditioning

See 4.4.

## 6.4 Procedure

**6.4.1** Measure the thickness of each specimen at four marked points to an accuracy of 0,01 mm using a dial micrometer, and calculate the mean value ( $\bar{d}_1$ ).

Measure individually the four sides of the square or the rectangle to an accuracy of 0,1 mm and calculate the mean values of the dimensions in the two perpendicular directions respectively (length  $\bar{l}_1$  and width  $\bar{b}_1$ ). For less regularly shaped specimens, for example from semi-finished or finished products, measure the most meaningful dimensions.

**6.4.2** Expose the specimens to the test environment chosen from those listed in clause 4.

**6.4.3** Rinse, if necessary (for example in the case of exposure to salt mist), and wipe the specimens dry.

**6.4.4** Remeasure the specimens in the same manner as before exposure, i.e. determine the dimensions after exposure ( $\bar{l}_2$ ,  $\bar{b}_2$  and  $\bar{d}_2$ ).

NOTE — If the specimens have undergone severe warpage, the linear measurements should be carried out with a tape measure.

**6.4.5** Note any change in appearance.

**6.4.6** Dry or recondition the specimens, if required, as described in 4.5.2 and determine the dimensions after drying or reconditioning ( $\bar{l}_3$ ,  $\bar{b}_3$ ,  $\bar{d}_3$ ).

## 6.5 Expression of results

**6.5.1** The results may be expressed in either of two ways:

a) as the percentage change in dimension (after exposure), based on the initial dimension, using, as applicable, the formulae

$$\frac{\bar{l}_2 - \bar{l}_1}{\bar{l}_1} \times 100, \quad \frac{\bar{b}_2 - \bar{b}_1}{\bar{b}_1} \times 100, \quad \frac{\bar{d}_2 - \bar{d}_1}{\bar{d}_1} \times 100$$

or (after exposure and drying or reconditioning)

$$\frac{\bar{l}_3 - \bar{l}_1}{\bar{l}_1} \times 100, \quad \frac{\bar{b}_3 - \bar{b}_1}{\bar{b}_1} \times 100, \quad \frac{\bar{d}_3 - \bar{d}_1}{\bar{d}_1} \times 100$$

The change is positive for increase and negative for decrease in the dimension.

b) as the percentage of the final dimension relative to the initial dimension, using, as applicable, the formulae

$$\frac{\bar{l}_2}{\bar{l}_1} \times 100, \quad \frac{\bar{b}_2}{\bar{b}_1} \times 100, \quad \frac{\bar{d}_2}{\bar{d}_1} \times 100$$

or  
 $\frac{\bar{l}_3}{\bar{l}_1} \times 100, \quad \frac{\bar{b}_3}{\bar{b}_1} \times 100, \quad \frac{\bar{d}_3}{\bar{d}_1} \times 100$  respectively.

In this case, 100 % means no change, values lower than 100 % a reduction and values higher than 100 % an increase in the dimensions.

Report any apparent deformation such as warpage, curling, delamination, or visible signs of surface degradation such as:

- change in colour and/or gloss, presence of crazing, cracks;
- blisters;
- exudation of plasticizers, tackiness;
- blooming of solid ingredients;
- corrosion of metal components, if any;

and assign, if possible, a qualification such as slight, moderate, severe, etc.

## 7 Change of other physical properties

### 7.1 General

Any physical property may be determined; the most significant are usually the mechanical, optical and electrical properties.

It is important that the precise conditions of test be specified in the relevant product specifications.