BS EN ISO 899-1:2003 +A1:2015

Incorporating Corrigendum No. 1

Plastics — Determination of creep behaviour —

Part 1: Tensile creep

ICS 83.080.01



National foreword

This British Standard is the UK implementation of EN ISO 899-1:2003+A1:2015. It is identical to ISO 899-1:2003, incorporating amendment 1:2015. It supersedes BS EN ISO 899-1:2003 which is withdrawn.

The UK participation in its preparation was entrusted to Technical Committee PRI/21, Testing of plastics.

A list of organizations represented on this committee can be obtained on request to its secretary.

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Amendments/corrigenda issued since publication

This British Standard was published under the authority of the Standards Policy and Strategy Committee on 18 June 2003

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Amd. No.	Date	Comments
14606 Corrigendum No. 1	31 July 2003	Replacing the EN ISO foreword page and incorporating the Annex ZA page
	30 April 2015	Implementation of ISO amendment 1:2015 with CEN endorsement A1:2015: Subclause A.2 updated

ISBN 978 0 580 77079 1

EUROPEAN STANDARD NORME EUROPÉENNE

EUROPÄISCHE NORM

EN ISO 899-1:2003+A1

March 2015

ICS 83.080.01

English version

Plastics - Determination of creep behaviour - Part 1: Tensile creep (ISO 899-1:2003)

Plastiques - Détermination du comportement au fluage -Partie 1: Fluage en traction (ISO 899-1:2003) Kunststoffe - Bestimmung des Kriechverhaltens - Teil 1: Zeitstand-Zugversuch (ISO 899-1:2003)

This European Standard was approved by CEN on 23 May 2003.

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BS EN ISO 899-1:2003+A1:2015

ISO 899-1:2003+A1:2015

CORRECTED 2003-07-16

Foreword

This document (EN ISO 899-1:2003) has been prepared by Technical Committee ISO/TC 61 "Plastics" in collaboration with Technical Committee CEN/TC 249 "Plastics", the secretariat of which is held by IBN.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by December 2003, and conflicting national standards shall be withdrawn at the latest by December 2003.

This document supersedes EN ISO 899-1:1996.

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Endorsement notice

The text of ISO 899-1:2003 has been approved by CEN as EN ISO 899-1:2003 without any modifications.

NOTE Normative references to International Standards are listed in Annex ZA (normative).

Foreword to amendment A1

This document (EN ISO 899-1:2003/A1:2015) has been prepared by Technical Committee ISO/TC 61 "Plastics" in collaboration with Technical Committee CEN/TC 249 "Plastics" the secretariat of which is held by NBN.

This Amendment to the European Standard EN ISO 899:2003 shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by September 2015, and conflicting national standards shall be withdrawn at the latest by September 2015.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CEN [and/or CENELEC] shall not be held responsible for identifying any or all such patent rights.

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Endorsement notice

The text of ISO 899-1:2003/Amd 1:2015 has been approved by CEN as EN ISO 899-1:2003/A1:2015 without any modification.

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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.

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 899-1 was prepared by Technical Committee ISO/TC 61, *Plastics*, Subcommittee SC 2, *Mechanical properties*.

This second edition cancels and replaces the first edition (ISO 899-1:1993), which has been technically revised.

ISO 899 consists of the following parts, under the general title *Plastics* — *Determination of creep behaviour*:

- Part 1: Tensile creep
- Part 2: Flexural creep by three-point loading

Plastics — Determination of creep behaviour —

Part 1:

Tensile creep

1 Scope

- **1.1** This part of ISO 899 specifies a method for determining the tensile creep of plastics in the form of standard test specimens under specified conditions such as those of pretreatment, temperature and humidity.
- **1.2** The method is suitable for use with rigid and semi-rigid non-reinforced, filled and fibre-reinforced plastics materials (see ISO 472 for definitions) in the form of dumb-bell-shaped test specimens moulded directly or machined from sheets or moulded articles.
- **1.3** The method is intended to provide data for engineering-design and research and development purposes. Data for engineering-design purposes requires the use of extensometers to measure the gauge length of the specimen. Data for research or quality-control purposes may use the change in distance between the grips (nominal extension).
- **1.4** Tensile creep may vary significantly with differences in specimen preparation and dimensions and in the test environment. The thermal history of the test specimen can also have profound effects on its creep behaviour (see Annex A). Consequently, when precise comparative results are required, these factors must be carefully controlled.
- **1.5** If tensile-creep properties are to be used for engineering-design purposes, the plastics materials should be tested over a broad range of stresses, times and environmental conditions.

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 62:1999, Plastics — Determination of water absorption

ISO 291:1997, Plastics — Standard atmospheres for conditioning and testing

ISO 472:1999, Plastics — Vocabulary

ISO 527-1:1993, Plastics — Determination of tensile properties — Part 1: General principles

ISO 527-2:1993, Plastics — Determination of tensile properties — Part 2: Test conditions for moulding and extrusion plastics

ISO 10350-1:1998, Plastics — Acquisition and presentation of comparable single-point data — Part 1: Moulding materials

ISO 11403-1:2001, Plastics — Acquisition and presentation of comparable multipoint data — Part 1: Mechanical properties

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 472 and the following apply.

3.1

creep

increase in strain with time when a constant force is applied

3.2

initial stress

 σ

tensile force per unit area of the initial cross-section within the gauge length

NOTE 1 It is given by the equation

$$\sigma = \frac{F}{A}$$

where

F is the force, in newtons;

A is the average initial cross-sectional area within the narrow (gauge) section of the specimen, in square millimetres.

NOTE 2 It is expressed in megapascals.

3.3

extension

 $(\Delta L)_t$

increase in the distance between the gauge marks, expressed in millimetres, at time t

NOTE It is given by the equation

$$(\Delta L)_t = L_t - L_0$$

where

 L_t is the gauge length, in millimetres, at any given time t during the test;

 L_0 is the original gauge length, in millimetres, of the specimen after application of a preload but prior to application of the test load.

3.4

nominal extension

 $(\Delta L^*)_t$

increase in the distance between the grips (increase in grip separation)

NOTE It is given by the equation

$$(\Delta L^*)_t = L_t^* - L_0^*$$

where

 L_t^* is the distance between the grips at any given time t during the test, in millimetres;

 L^*_0 is the initial distance between the grips, expressed in millimetres, holding the specimen after application of a preload but prior to application of the test load.

3.5

tensile-creep strain

 \mathcal{E}_{t}

change in the distance between the gauge marks, relative to the initial distance, produced by the applied load at any given time t during a creep test

NOTE 1 It is given by the equation

$$\varepsilon_t = \frac{(\Delta L)_t}{L_0}$$

NOTE 2 It is expressed as a dimensionless ratio or as a percentage.

3.6

nominal tensile-creep strain

 \mathcal{E}^{\star}_{t}

change in the distance between the grips, relative to the initial distance, produced by the applied load at any given time t during a creep test

NOTE 1 It is given by the equation

$$\varepsilon_t^* = \frac{(\Delta L^*)_t}{L^*_0}$$

NOTE 2 It is expressed as a dimensionless ratio or as a percentage.

3.7

tensile-creep modulus

 E_{t}

ratio of initial stress to tensile-creep strain, calculated as in 7.1.1

3.8

nominal tensile-creep modulus

 E^*_t

ratio of initial stress to nominal tensile-creep strain, calculated as in 7.1.2

3.9

isochronous stress-strain curve

Cartesian plot of stress versus creep strain, at a specific time after application of the test load

3.10

time to rupture

period of time the specimen is under full load until rupture

3.11

creep-strength limit

initial stress which will just cause rupture ($\sigma_{B,t}$) or will produce a specified strain ($\sigma_{\varepsilon,t}$) at a specified time t, at a given temperature and relative humidity

3.12

recovery from creep

decrease in strain at any given time after completely unloading the specimen, expressed as a percentage of the strain just prior to the removal of the load

4 Apparatus

4.1 Gripping device, capable of ensuring that the direction of the load applied to the test specimen coincides as closely as possible with the longitudinal axis of the specimen. This ensures that the test specimen is subjected to simple stress and that the stresses in the loaded section of the specimen may be assumed to be uniformly distributed over cross-sections perpendicular to the direction of the applied load.

NOTE It is recommended that grips be used that will allow the specimen to be fixed in place, correctly aligned, prior to applying the load. Self-locking grips which allow the specimen to move as the load increases are not suitable for this test

- **4.2 Loading system**, capable of ensuring that the load is applied smoothly, without causing transient overloading, and that the load is maintained to within \pm 1 % of the desired load. In creep-to-rupture tests, provision shall be made to prevent any shocks which occur at the moment of rupture being transmitted to adjacent loading systems. The loading mechanism shall allow rapid, smooth and reproducible loading.
- **4.3 Extension-measuring device**, comprising any contactless or contact device capable of measuring the extension of the specimen gauge length or the increase in the distance between the clamp grips under load without influencing the specimen behaviour by mechanical effects (e.g. undesirable deformations, notches), other physical effects (e.g. heating of the specimen) or chemical effects.

In the case of contactless (optical) measurement of the strain, the longitudinal axis of the specimen shall be perpendicular to the optical axis of the measuring device. To determine the increase in length of the test specimen, an extensometer shall be used which records the increase in the distance between the clamp grips. The accuracy of the extension-measuring device shall be better than \pm 0,01 mm.

For creep-to-rupture tests, it is recommended that the extension be measured by means of a contactless optical system operating on the cathetometer principle. Automatic indication of time to rupture is highly desirable. The gauge length shall be marked on the specimen, either by attaching (metal) clips with scratched-on gauge marks, or by ruling the gauge marks with an inert, thermally stable paint.

Electrical-resistance strain gauges are suitable only if the material tested is of such a nature as to permit such strain gauges to be attached to the specimen by means of adhesive and only if the adhesion quality is constant during the duration of the test. The modulus of the strain gauge when bonded to the specimen shall be such that the specimen is not reinforced.

- **4.4** Time-measurement device, accurate to 0,1 %.
- **4.5 Micrometer**, reading to 0,01 mm or closer, for measuring the initial thickness and width of the test specimen.

5 Test specimens

Use test specimens of the same shape and dimensions as specified for the determination of tensile properties by the relevant material standard or, by default, as specified in ISO 527-2.

6 Procedure

6.1 Conditioning and test atmosphere

Condition the test specimens as specified in the International Standard for the material under test. In the absence of any information on conditioning, use the most appropriate set of conditions specified in ISO 291, unless otherwise agreed by the interested parties.

The creep behaviour will be affected not only by the thermal history of the specimen under test, but also by the temperature and (where applicable) humidity used in conditioning. If the specimen is not in humidity equilibrium, creep will be affected in the following way: a specimen which is too dry will produce an additional strain due to water absorption during the test and a specimen which is too humid will contract due to water desorption. It is recommended that a conditioning time $\geq t_{90}$ (see ISO 62) be used.

Conduct the test in the same atmosphere as used for conditioning, unless otherwise agreed upon by the interested parties, e.g. for testing at elevated or low temperatures. Ensure that the variation in temperature during the duration of the test remains within \pm 2 °C.

6.2 Measurement of test-specimen dimensions

Measure the dimensions of the conditioned test specimens in accordance with ISO 527-1:1993, Subclause 9.2.

6.3 Mounting the test specimens

Mount a conditioned and measured specimen in the grips and set up the extension-measuring device as required.

6.4 Selection of stress value

Select a stress value appropriate to the application envisaged for the material under test, and calculate, using the equation given in 3.2, the load to be applied to the test specimen.

If the initial strain is specified instead of the stress, the stress value may be calculated using Young's modulus for the material (see ISO 527-1).

6.5 Loading procedure

6.5.1 Preloading

When it is necessary to preload the test specimen prior to increasing the load to the test load, for example in order to eliminate backlash by the test gear, take care to ensure that the preload does not influence the test results. Do not apply the preload until the temperature and humidity of the test specimen (gripped in the test apparatus) correspond to the test conditions. Measure the gauge length after application of the preload. Maintain the preload during the whole duration of the test.

6.5.2 Loading

Load the test specimen smoothly so that full loading of the specimen is reached between 1 s and 5 s after the beginning of the application of the load. Use the same rate of loading for each of a series of tests on one material.

Take the total load (including the preload) to be the test load.

6.6 Extension-measurement schedule

Record the point in time at which the specimen is fully loaded as t = 0. Unless the extension is automatically and/or continuously recorded, choose the times for making individual measurements as a function of the creep curve obtained from the particular material under test. It is preferable to use the following measurement schedule:

1 min, 3 min, 6 min, 12 min and 30 min;

1 h, 2 h, 5 h, 10 h, 20 h, 50 h, 100 h, 200 h, 500 h, 1 000 h, etc.

If discontinuities are suspected or observed in the creep-strain versus time plot, take readings more frequently.

6.7 Time measurement

Measure, to within \pm 0,1 % or \pm 2 s (whichever is the less severe tolerance), the total time which has elapsed up to each creep measurement.

6.8 Temperature and humidity control

Unless temperature and relative humidity (where applicable) are recorded automatically, record them at the beginning of the test and then at least three times a day initially. When it has become evident that the conditions are stable within the specified limits, they may be checked less frequently (but at least once a day).

6.9 Measurement of recovery rate (optional)

Upon completion of non-rupture tests, remove the load rapidly and smoothly and measure the recovery rate using, for instance, the same schedule as was used for creep measurement.

7 Expression of results

7.1 Method of calculation

7.1.1 Tensile-creep modulus, E_t

Calculate the tensile-creep modulus, E_t , by dividing the initial stress, σ , by the tensile-creep strain, ε_t , at each of the selected measurement times.

It is given, in megapascals, by the equation

$$E_t = \frac{\sigma}{\varepsilon_t} = \frac{F \cdot L_0}{A \cdot (\Delta L)_t}$$

where

F is the applied force, in newtons;

 L_0 is the initial gauge length, in millimetres;

A is the initial cross-sectional area, in square millimetres, of the specimen;

 $(\Delta L)_t$ is the extension, in millimetres, at time t.

7.1.2 Nominal tensile-creep modulus, E_t^*

Calculate the nominal tensile-creep modulus, E_t^* , by dividing the initial stress, σ , by the nominal tensile-creep strain, ε_t^* , at each of the selected measurement times.

It is given, in megapascals, by the equation

$$E_t^* = \frac{\sigma}{\varepsilon_t^*} = \frac{F \cdot L_0^*}{A \cdot (\Delta L_t^*)_t}$$

where

F is the applied force, in newtons;

 L_0^* is the initial distance between the grips, in millimetres;

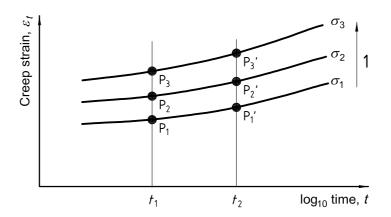
A is the initial cross-sectional area of the specimen, in square millimetres;

 $(\Delta L^*)_t$ is the increase in the distance between the grips, in millimetres, at time t.

7.2 Presentation of results

7.2.1 Creep curves

If testing is carried out at different temperatures, the raw data should preferably be presented, for each temperature, as a series of creep curves showing the tensile strain plotted against the logarithm of time, one curve being plotted for each initial stress used (see Figure 1).



Key

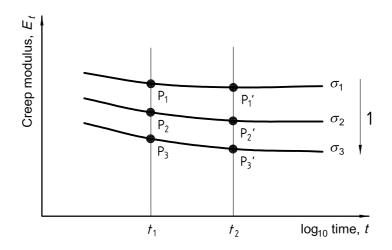
1 increasing stress

Figure 1 — Creep curves

The data may also be presented in other ways, e.g. as described in 7.2.2 and 7.2.3, to provide information required for particular applications.

7.2.2 Creep-modulus/time curves

For each initial stress used, the tensile-creep modulus, calculated in accordance with 7.1.1, may be plotted against the logarithm of the time under load (see Figure 2).



Key

1 increasing stress

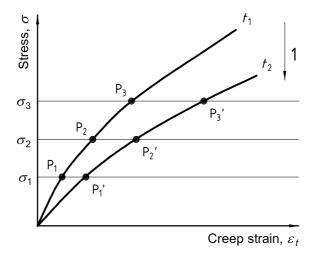
Figure 2 — Creep-modulus/time curves

If testing is carried out at different temperatures, plot a series of curves for each temperature.

7.2.3 Isochronous stress-strain curves

An isochronous stress-strain curve is a Cartesian plot showing how the strain depends on the applied load, at a specific point in time after application of the load. Several curves are normally plotted, corresponding to times under load of 1 h, 10 h, 100 h, 1 000 h and 10 000 h. Since each creep test gives only one point on each curve, it is necessary to carry out the test at, at least, three different stresses, and preferably more, to obtain an isochronous curve (see ISO 11403-1).

To obtain an isochronous stress-strain curve for a particular time under load (say 10 h) from a series of creep curves as shown in Figure 1, read off, from each creep curve, the strain at 10 h, and plot these strain values (x-axis) against the corresponding stress values (y-axis). Repeat the process for other times to obtain a series of isochronous curves (see Figure 3).



Key

1 increasing time

Figure 3 — Isochronous stress-strain curves

If testing is carried out at different temperatures, plot a series of curves for each temperature.

7.2.4 Three-dimensional representation

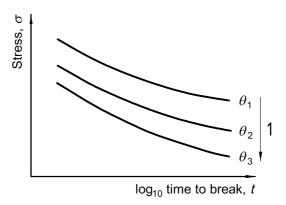
A relationship of the form $\varepsilon = f(t, \sigma)$ exists between the different types of curve (see Figures 1 to 3) that can be derived from the raw creep-test data. This relationship can be represented as a surface in a three-dimensional space (see Reference [1] in the Bibliography).

All the curves that can be derived from the raw creep-test data form part of this surface. Because of the experimental error inherent in each measurement, the points corresponding to the actual measurements normally do not lie on the curves but just off them.

The surface $\varepsilon = f(t, \sigma)$ can therefore be generated by deriving a number of the curves which form it, but a number of sophisticated smoothing operations are usually necessary. Computer techniques permit this to be done rapidly and reliably.

7.2.5 Creep-to-rupture curves

Creep-to-rupture curves allow the prediction of the time to failure at any stress. They may be plotted as stress against log time to break (see Figure 4) or log stress against log time to break.



Key

1 increasing temperature

NOTE The stress, σ , may also be plotted on a logarithmic scale.

Figure 4 — Creep-to-rupture curves

7.3 Precision

The precision of this test method is not known because interlaboratory data are not available. When interlaboratory data are obtained, a precision statement will be added at the following revision.

8 Test report

The test report shall include the following particulars:

- a) a reference to this part of ISO 899;
- b) a complete description of the material tested, including all pertinent information on composition, preparation, manufacturer, tradename, code number, date of manufacture, type of moulding and any annealing;
- c) the dimensions of each test specimen;

- d) the method of preparation of the test specimens;
- e) the directions of the principal axes of the test specimens with respect to the dimensions of the product or some known or inferred orientation in the material;
- f) details of the atmosphere used for conditioning and testing;
- g) which tensile creep modulus, E_t or E_t^* , was calculated;
- h) the creep-test data for each temperature at which testing was carried out, presented in one or more of the graphical forms described in 7.2, or in tabular form;
- i) if recovery-rate measurements are made, the time-dependent strain after unloading the test specimen (see 6.9).

Annex A (informative)

Physical-ageing effects on the creep of polymers

A.1 General

Physical ageing takes place when a polymer is cooled from an elevated temperature at which the molecular mobility is high to a lower temperature at which relaxation times for molecular motions are long in comparison with the storage time at that temperature. Under these circumstances, changes in the structure will take place over a long period of time, involving rearrangement in the shape and packing of molecules as the polymer approaches the equilibrium structural state for the lower temperature. Associated with this ageing process, there is a progressive decrease in the molecular mobility of the polymer, even when the temperature remains constant. As a direct consequence of this, the creep deformation produced by an applied stress will depend upon the age of the polymer, creep rates being lower in more highly aged material.

This is illustrated in Figure A.1 which shows creep compliance curves for PVC specimens of different ages. Each of these specimens has been rapidly cooled from a temperature of 85 °C (close to $T_{\rm g}$) and stored at the test temperature of 23 °C for different times $t_{\rm e}$ prior to load application. The physical age of a specimen is then defined by the time $t_{\rm e}$ and it can be seen that the older the specimen the further its creep curve is shifted on the time axis.

A.2 Creep at elevated temperatures

The influence of physical ageing on creep behaviour is more complicated when measurements are made at elevated temperatures following a storage period at a lower temperature. It is well known that an increase in temperature leads to an increase in molecular mobility and thus a higher rate of creep deformation. In addition to this, changes in molecular structure take place on heating that are associated with a reduction in the physical age of the polymer and lead to a further increase in mobility. Creep deformation at the higher temperature is, therefore, more rapid than expected from the temperature increase alone. With increasing time, physical ageing is reactivated, and the associated progressive decrease in mobility thus leads to a shift in creep behaviour to longer time, as described in A.1, and thus to a dependence of creep behaviour on dwell time at the high temperature prior to load application. The timescales associated with the changes in physical age depend on the age of the polymer prior to the temperature increase and the magnitudes of the temperature increase and the glass-transition temperature.

Illustrations of the transient changes in creep behaviour that can occur with dwell time at the elevated temperature are shown in Figures A.2 and A.3. In Figure A.2, PVC specimens were stored at 23 °C for 200 h prior to heating to the test temperature of 44 °C. Creep curves were then measured after different dwell times, $t_{\rm e2}$, at 44 °C prior to load application. The shift in creep behaviour to longer times is interpreted as the reactivation of physical ageing at 44 °C before loading following the reduction in age state from that at 23 °C resulting from the increase in temperature. In Figure A.3, creep tests were carried out under the same conditions but following a storage period of greater than 1 year at 23 °C prior to heating to the test temperature. The progressive reduction in the structural age of the specimens is observed here as a shift in the curves to shorter creep times and arises because of the more extensive structural changes that have taken place through physical ageing at 23 °C before heating that are not fully overcome by the relatively short times, $t_{\rm e}$, at the temperature prior to loading.

A further issue needs to be considered in the analysis of creep data at elevated temperatures. The shape of a creep curve at the elevated temperature will change, if during the reactivation of physical ageing, significant changes in age take place in the duration of the creep test. Any attempt to construct creep master curves using procedures based on time-temperature equivalence must take account of these transient changes in molecular mobility linked to physical ageing for predictions of long-term behaviour to have any validity.

The changes in creep behaviour with time shown in these figures following cooling or heating are associated with changes in the non-equilibrium structure of the amorphous phase established when the polymer is cooled below its glass-transition temperature. Similar effects are observed in the creep behaviour of semi-crystalline polymers even if the glass transition temperature is below ambient. These effects are believed to be caused by physical ageing in the amorphous phase associated with a relaxation process (the α -process) involving coupled motions of molecules spanning both the crystal and amorphous phases.

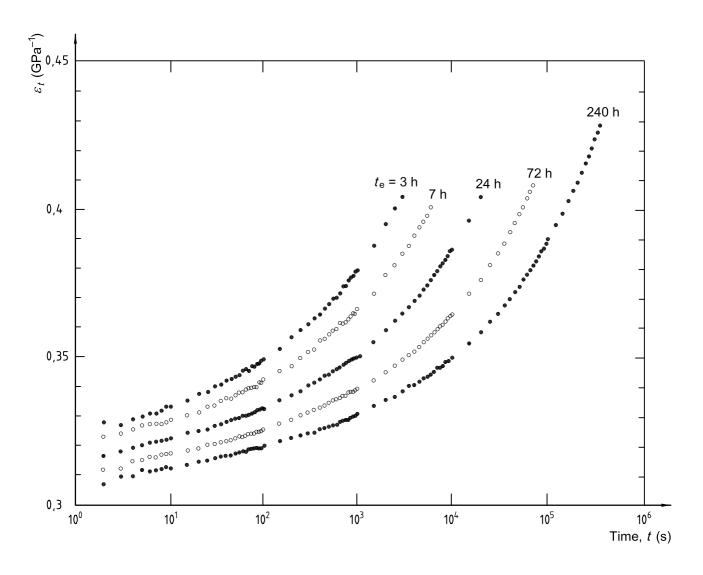


Figure A.1 — Creep curves for PVC at 23 °C obtained at different times $t_{\rm e}$ after rapid cooling of the specimen from 85 °C to 23 °C

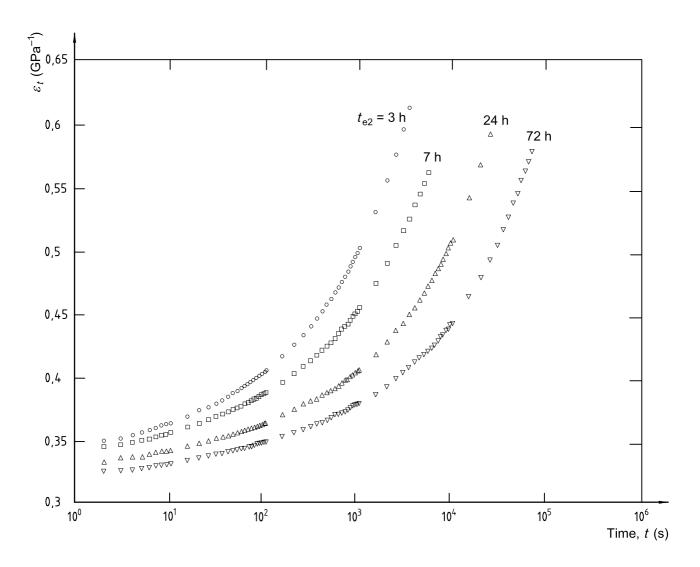


Figure A.2 — Creep curves for PVC at 44 °C obtained by application of the load at different times $t_{\rm e2}$ after heating from 23 °C (the specimen had been stored for 200 h at 23 °C prior to heating)

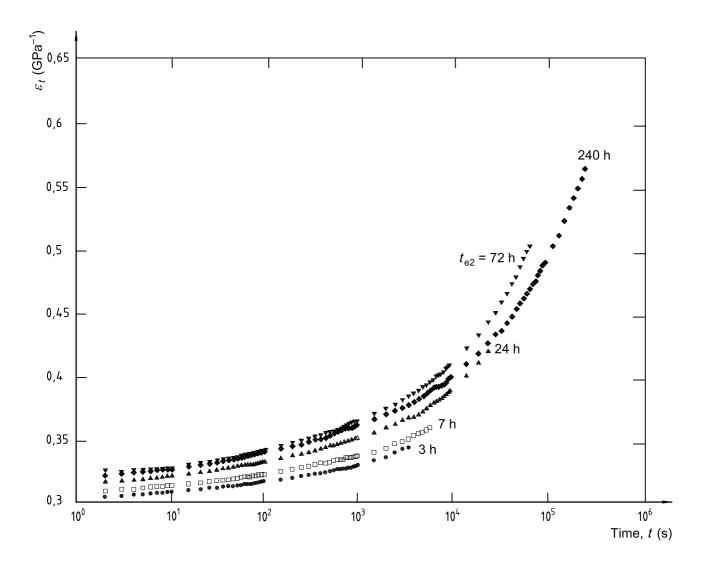


Figure A.3 — As for Figure A.2 but following storage for more than 1 year at 23 °C prior to heating

Bibliography

- [1] TURNER, S., Creep in thermoplastics Preliminary concepts and definitions, *British Plastics*, June (1964), pp. 322-324
- [2] LEHMANN, J., *Zeitstandverhalten von PMMA in der Freibewitterung und im Normklima*, V15 in 27. Jahrestagung der GUS, 18th March 1998, ISBN 3-9806167-0-3

Annex ZA (normative)

Normative references to international publications with their relevant European publications

This European Standard incorporates by dated or undated reference, provisions from other publications. These normative references are cited at the appropriate places in the text and the publications are listed hereafter. For dated references, subsequent amendments to or revisions of any of these publications apply to this European Standard only when incorporated in it by amendment or revision. For undated references the latest edition of the publication referred to applies (including amendments).

NOTE Where an International Publication has been modified by common modifications, indicated by (mod.), the relevant EN/HD applies.

Publication	<u>Year</u>	<u>Title</u>	<u>EN</u>	<u>Year</u>
ISO 62	1999	Plastics — Determination of water absorption	EN ISO 62	1999
ISO 291	1997	Plastics — Standard atmospheres for conditioning and testing	EN ISO 291	1997
ISO 472	1999	Plastics — Vocabulary	EN ISO 472	2001
ISO 527-1	1993	Plastics — Determination of tensile properties — Part 1: General principles	EN ISO 527-1	1996
ISO 527-2	1993	Plastics — Determination of tensile properties — Part 2: Test conditions for moulding and extrusion plastics	EN ISO 527-2	1996
ISO 10350-1	1998	Plastics — Acquisition and presentation of comparable single-point data — Part 1: Moulding materials	EN ISO 10350-1	2000
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