

SLOVENSKI STANDARD SIST-TS CEN/TS 839:2008

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Wood preservatives - Determination of the protective effectiveness against wood destroying basidiomycetes - Application by surface treatment

Holzschutzmittel - Bestimmung der vorbeugenden Wirksamkeit gegen holzzerstörende Basidiomyceten - Anwendung mit Oberflächenverfahren

Produits de préservation du bois - <u>Détermination de l'eff</u>icacité protectrice vis-a-vis des champignons basidiomycetes lighivores Application par traitement de surface 98eea81543c0/sist-ts-cen-ts-839-2008

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Wood preservatives - Determination of the protective effectiveness against wood destroying basidiomycetes -Application by surface treatment

Produits de préservation du bois - Détermination de l'efficacité protectrice vis-à-vis des champignons basidiomycètes lignivores - Application par traitement de surface Holzschutzmittel - Bestimmung der vorbeugenden Wirksamkeit gegen holzzerstörende Basidiomyceten -Anwendung mit Oberflächenverfahren

This Technical Specification (CEN/TS) was approved by CEN on 5 November 2007 for provisional application.

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Foreword

This document (CEN/TS 839:2008) has been prepared by Technical Committee CEN/TC 38 "Durability of wood and wood-based products", the secretariat of which is held by AFNOR.

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.

This document supersedes ENV 839:2002.

Significant technical differences between this standard and ENV 839:2002 are as follows:

- a) change of assessment from, visual examination for decay supplemented with culturing to assess colonisation of the interior, to determination of loss in mass due to fungal decay as well as visual examination for decay of the surface and the interior of the test specimens. The visual examination is now included as an optional means of assessment to determine colonisation and that this requires an additional series of test specimens (Annex C).
- b) addition in 7.5 of treated check test specimens for calculation of the correction value;
- c) taking into account of the terms given in EN 1001-1and the definitions of EN 1001-2;
- d) Introduction of an informative Annex E to take account of consideration for minimisation of environmental and health hazards caused by the use of this biological test.

This standard includes Annexes A, C, D and E that are informative and an Annex B that is normative. https://standards.iteh.ai/catalog/standards/sist/c7a12f40-4089-4bdd-b5b3-

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Introduction

This European Standard specifies a laboratory method of test which gives a basis for assessing the effectiveness of a wood preservative, when applied as a surface treatment, against wood destroying basidiomycetes. It tests whether the applied treatment is able to prevent the penetration of the fungi into the untreated interior of the test specimens under the conditions of test.

This laboratory method provides one criterion by which the effectiveness of a product can be assessed. In making this assessment, the methods by which the preservative may be applied should be taken into account. It is also recommended that results from this test should be supplemented by those from other relevant tests and above all by practical experience.

The procedures described in this standard method are intended to be carried out by suitably trained and/or supervised specialists.

Suitable precautions should include the use of separate rooms, areas within rooms, extraction facilities, conditioning chambers and special training for personnel. Also see Annex E for environmental, health and safety precautions.

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1 Scope

This European Standard specifies a method of test for the determination of the protective effectiveness of a wood preservative, applied to the surface of the wood, against wood destroying basidiomycetes cultured on an agar medium.

The method is applicable to all products which are to be applied by superficial application processes. This includes :

- organic solvent-based wood preservatives ; or
- organic water-dispersible formulations, as supplied or as prepared in the laboratory by dilution of concentrates; or
- water-soluble products ; or
- chemicals which are being studied as active ingredients for application by superficial processes.

NOTE This method may be used in conjunction with an ageing procedure, for example EN 73.

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.

EN ISO 3696, Water for analytical laboratory use - Specification and test methods (ISO 3696:1987).

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3 Terms and definitions 98eea81543c0/sist-ts-cen-ts-839-2008

For the purposes of this document, the following terms and definitions apply:

3.1

representative sample

sample having its physical and/or chemical characteristics identical to the volumetric average characteristics of the total volume being sampled.

[EN 1001-2,4.71]

3.2

supplier

sponsor of the test (person or company providing the sample of wood preservative to be tested). [Adapted from EN 1001-2,**4.83**]

3.3

superficial application process

process which does not include particular features or procedures intended to overcome the natural resistance of wood to penetration of a wood preservative in its ready to use form

[EN 1001-2,**4.82**]

4 Principle

Several series of test specimens of a susceptible wood species are end-sealed with a material to prevent penetration of the wood preservative under test into the end grain of the test specimens. The end-sealed test specimens are treated with the wood preservative under test using the process and application rate specified by the supplier.

NOTE Suitable application methods are brushing, pipetting and dipping.

The treated test specimens are exposed to attack by basidiomycetes in pure culture. The performance of the test product is assessed in terms of its ability to prevent decay as determined by the maximum acceptable loss in mass and the absence of visible decay of the surface and the untreated interior.

5 Test materials and apparatus

5.1 Biological material

The test fungi to be used are as follows:

5.1.1 Obligatory fungus in all cases

— Coniophora puteana (Schumacher ex Fries) Karsten (BAM Ebw. 15) on softwood.

Loss in mass of Scots pine sapwood in 16 weeks: a mass fraction of minimum 20 %.

5.1.2 Obligatory fungus for particular hazards

— Coriolus versicolor (Linnaeus) Quélet (CTB 863A) on hardwood and/or on softwood as appropriate.

Loss in mass of beech in 16 weeks: a mass fraction of minimum 20 %. https://standards.iten.avcatalog/standards/sist/c/a12i40-4089-4bdd-b5b3-Loss in mass of Scots pine sapwood in 16 weeks: a mass fraction of minimum 15 %.

5.1.3 Two species to be used compulsorily on the basis of the nature of the test product

For all products except creosote-type products:

— Poria placenta (Fries) Cooke sensu J. Eriksson (FPRL 280) on softwood.

Loss in mass of Scots pine sapwood in 16 weeks: a mass fraction of minimum 20 %;

— *Gloeophyllum trabeum* (Persoon ex Fries) Murrill (BAM Ebw. 109) on softwood.

Loss in mass of Scots pine sapwood in 16 weeks: a mass fraction of minimum 20 %.

For creosotes and similar products:

— Lentinus lepideus Fries ex Fries (BAM Ebw. 20) on softwood.

Loss in mass of Scots pine sapwood in 16 weeks: a mass fraction of minimum 20 %;

— Lentinus cyathiformis (Schaeffer ex Fries) Bresadola (CTB 67-02B) on hardwood.

Loss in mass of beech in 16 weeks: a mass fraction of minimum 20 %.

5.1.4 Optional fungi

For specific regional uses or conditions, it is also possible to select other fungi on an optional basis.

NOTE When optional fungi are used, information similar to that given in annex A for the obligatory fungi should be included in the test report.

5.1.5 Maintenance of strains

The strains shall be maintained and treated (frequency of subculturing, alternation of culture media, etc.) in accordance with the instructions of their laboratory of origin (see A.2). The parent strain shall be maintained in the laboratory of its origin so as to conserve and to assure its vigour.

If tests are not undertaken regularly or if a strain shows signs of degeneration a new standard culture of the strain should be obtained from the laboratory of its origin for each test (see A.2). When new strains are received, the virulence shall be tested to ensure the strain can achieve the minimum loss in mass (see 5.1.1, 5.1.2 and 5.1.3).

5.2 Products and reagents

5.2.1 Culture medium

The culture medium is a malt agar medium with the following composition:

- malt extract :
 - in concentrated form: (50 ± 0,5) g;
 - in powder form: (40 ± 0.5) g;
- agar causing no inhibition of growth of fungi:
 - $(20 \pm 0,5) g \text{ to } (30 \pm 0,5) g; TANDARD PREVIEW$
- water conforming to grade 3 of EN ISO 3696.
 standards.iteh.ai)
 - quantity to make up to 1000 ml.
 - SIST-TS CEN/TS 839:2008

Prepare this medium by/warming ithe mixture in a boiling water bath of steam 3 bath, stirring until completely dissolved. 98eea81543c0/sist-ts-cen-ts-839-2008

Place in each culture vessel (5.3.1) a sufficient quantity of the medium to provide a minimum depth of 3 mm to 4 mm when in its in-use position. Close the vessels as specified in 5.3.1 and sterilise in an autoclave at 121 °C for 20 min. Let the vessels cool in their in-use position.

5.2.2 Solvents and diluents

For water soluble or water dispersible preservatives:

— water conforming to grade 3 of EN ISO 3696.

For preservatives to be diluted or dissolved in an organic solvent:

- suitably volatile liquids that leave no residue in the wood that would have a toxic effect on the fungi at the end of the post-treatment conditioning period.
- NOTE Toluene and xylene of recognized analytical grade have been found suitable.

5.2.3 Fumigant (if necessary)

Xylene technical grade.

5.2.4 End-seal compound

A material resistant to the penetration of the wood preservative under test and the test fungi, or separate materials for each, and without any fungistatic or fungicidal activity within the test specimen.

NOTE Three brush coats of a 2-component epoxy lacquer, with drying between each application, have been found to be suitable.

5.3 Apparatus

5.3.1 *Culture vessels*, Kolle flasks or equivalent vessels with a capacity of between 400 ml and 650 ml, providing a flat surface area of between 85 cm^2 and 120 cm^2 for the medium.

NOTE 1 Examples of suitable vessels are given in EN 113.

NOTE 2 Kolle flasks are usually plugged with a wad of cotton wool. Other culture vessels are usually fitted with leak proof lids, the centres of which are pierced with a round hole of up to 15 mm diameter and plugged with a wad of cotton wool.

5.3.2 Drying oven, capable of being controlled at (103 ± 2) °C.

5.3.3 *Desiccators*, with efficient desiccant (silica gel for example).

5.3.4 Conditioning chamber, well ventilated and controlled at (20 ± 2) °C and (65 ± 5) % relative humidity.

5.3.5 *Drying supports*, which will give a minimum contact with the treated test specimens. The supports shall be of a material that does not react with the test solvent or test wood preservative, for example glass for organic products.

5.3.6 *Culture chamber*, (incubator or room), dark and controlled at (22 ± 2) °C and (70 ± 5) % relative humidity.

5.3.7 *Test specimen supports*, made of glass, stainless steel or any other inert material, that is to say, with no risk of having any effect on the culture medium, the fungus, the wood or the test wood preservative, or of being itself modified. Supports can be capable of holding either one or two test specimens. The supports are used to prevent direct contact of the test specimens with the culture medium, but shall not separate them from it by more than 3 mm. <u>98eea81543c0/sist-ts-cen-ts-839-2008</u>

NOTE If abnormally high moisture contents in the test specimens are experienced consistently, use of test specimen supports of approximately 5 mm thick can help to control the problem. If thicker test specimen supports are used, this should be recorded in the test report.

5.3.8 *Ordinary laboratory equipment*, including a balance capable of weighing to the nearest of 0,01 g and an autoclave.

6 Sampling of the preservative

The sample of the wood preservative shall be representative of the product to be tested. Samples shall be stored and handled in accordance with any written instructions from the supplier.

NOTE For the sampling of wood preservatives from bulk supplies, the procedure given in EN 212 should be used.

7 Test specimens

7.1 Species of wood

The species of wood to be used shall be susceptible to attack by fungi and shall be readily penetrated by liquids.

The reference species are Scots pine (*Pinus sylvestris* Linnaeus) representing softwoods and beech (*Fagus sylvatica* Linnaeus) representing hardwoods.

Additional tests may be undertaken using other species corresponding to the above characteristics, and of particular importance for certain countries, but if so this shall be stated in the test report.

7.2 Wood quality

The wood shall be free from cracks, stain, decay, insect damage or other defects. The wood shall not have been water-stored, floated, chemically treated or steamed.

NOTE Wood that has been kiln dried at temperatures below 60 °C can be used.

The Scots pine shall be exclusively sapwood containing little resin and having between 2,5 and 8 annual growth rings per 10 mm. The proportion of latewood in the annual rings shall not exceed 30 % of the whole.

The beech shall be even-grained, free from tyloses and discoloration. It shall have between 2 and 6 annual growth rings per 10 mm.

7.3 Provision of the test specimens

Prepare planed strips having a cross section of $(25 \pm 0.5) \text{ mm} \times (15 \pm 0.5) \text{ mm}$. The longitudinal faces shall be parallel to the direction of the grain. The annual rings shall have a contact angle of $(45 \pm 15)^{\circ}$ to the broad faces. Make transverse cuts, neatly to give sharp edges and a fine-sawn finish to the end-grain surfaces, to give test specimens $(50 \pm 0.5) \text{ mm}$ long.

For treatment, drying and ageing, the test specimens can be retained in planed strips of a length sufficient to provide one test specimen for exposure to each of the test fungi. Each strip should be end-sealed prior to treatment.

The specimens shall originate from a minimum of three trees or shall be taken at random from a stock originally of more than 500 test specimens and originating from at least five planks.

7.4 Dimensions and density of test specimens

The dimensions of each test specimen at a mass fraction of (12 ± 2) % moisture content shall be (50 ± 0.5) mm x (25 ± 0.5) mm x (15 ± 0.5) mm x (15 ± 0.5) mm.

NOTE A moisture meter of the two-pronged electrical conductivity type is suitable for assessing moisture content.

The total surface area of the faces to be treated is theoretically 40 cm² but an allowance shall be made for any encroachment of the sealing compound on to these faces.

In a batch of test specimens to be treated, the density of an individual is permitted to differ from the mean value of the batch by \pm 10 %. This tolerance is increased to \pm 20 % for the untreated test specimens. The mean density for the treated test specimens used for the test shall be recorded in the test report.

7.5 Number and distribution of test specimens

The test specimens are divided into:

- e₁ treated test specimens :
 - these are the treated test specimens subjected to attack by the wood destroying fungi. Use at least six test specimens for each combination of preservative, quantity to be applied, preservative concentration, test fungus and for each timber species.
 - In case of dipping select at least 6 test specimens within a range of 10 % from the target retention. Supplementary samples shall be treated, in order to have a sufficient number of correctly treated specimens to put in test.

NOTE The treated test specimens are assessed by visual examination for decay of their surfaces and/or interior by the test fungi. If optional tests for colonisation of the test fungi are required as an additional method of assessment, this should be carried out on a parallel series of treated test specimens (Annex C).