## INTERNATIONAL STANDARD

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## Corrosion of metals and alloys — Classification of low corrosivity of indoor atmospheres —

## Part 2: Determination of corrosion attack in indoor atmospheres

Corrosion des métaux et alliages — Classification de la corrosivité faible des atmosphères d'intérieur —

Partie 2: Détermination de l'attaque par corrosion dans les atmosphères d'intérieur

ISO 11844-2:2020

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## Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

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

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This document was prepared by Technical Committee ISO/TC 156, *Corrosion of metals and alloys*, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/ TC 262, *Metallic and other inorganic coatings, including for corrosion protection and corrosion testing of metals and alloys*, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

This second edition cancels and replaces the first edition (ISO 11844-2:2005), which has been technically revised. The main changes compared with the previous edition are as follows:

- lead has been included as a standard specimen with high sensitivity to vapour organic acids;
- <u>Annex D</u> has been added.

A list of all parts in the ISO 11844 series can be found on the ISO website.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at <u>www.iso.org/members.html</u>.

## Introduction

This document describes standard specimens, exposure and evaluation for the derivation of indoor corrosivity categories.

The determination of the corrosion attack is, at the present state of knowledge, the most reliable and, usually, also an economical way for evaluating corrosivity, taking into account all the main local environmental influences.

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# Corrosion of metals and alloys — Classification of low corrosivity of indoor atmospheres —

# Part 2: **Determination of corrosion attack in indoor atmospheres**

### 1 Scope

This document specifies methods for determining corrosion rates with standard specimens of metals in indoor atmospheres with low corrosivity. For this direct method of evaluation corrosivity, different sensitive methods can be applied using standard specimens of the following metals: copper, silver, zinc, steel and lead. The values obtained from the measurements are used as classification criteria for the determination of indoor atmospheric corrosivity.

### 2 Normative references

There are no normative references in this document.

### 3 Terms and definitions

No terms and definitions are listed in this document.

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

— ISO Online browsing platform: available at <u>https://www.iso.org/obp</u>

https:/—alIEC Electropedia: available at http://www.electropedia.org/ 92-76fa132cc181/iso-11844-2-2020

### 4 Principle

The corrosivity of the indoor location (e.g. control rooms, electric boxes, storage rooms, during transportation, in museums) is determined from the corrosion rate calculated from the mass change or resistance change per unit area of standard specimens of metals after exposure for a certain time period. Different materials are sensitive to different environmental parameters or their combinations.

### 5 Methods

The following methods, described in <u>Annexes A</u> and <u>B</u>, are available for the evaluation of the corrosion attack:

- determination of corrosion rate by mass change measurements (see <u>Annex A</u>);
- determination of corrosion rate by electrolytic cathodic reduction (see <u>Annex B</u>).

The methods described in <u>Annexes C</u> and <u>D</u> are suitable for continuous or periodic monitoring of the corrosion attack:

- determination of corrosion rate by resistance measurements (see <u>Annex C</u>);
- determination of corrosion rate by quartz crystal micro-balance methodology (see <u>Annex D</u>).

Special features of the methods, such as sensitivity, possibility for continuous or periodic assessment of corrosion attack, available space, etc., should be considered when choosing the most suitable methods. Examples of suitable racks for exposure of specimens are given in Figures 1 and 2.

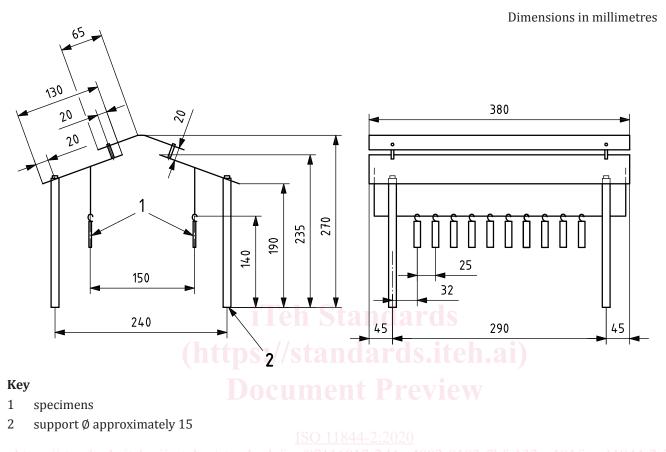


Figure 1 — Example of exposure racks for the sheltered exposure of specimens

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Dimensions in millimetres

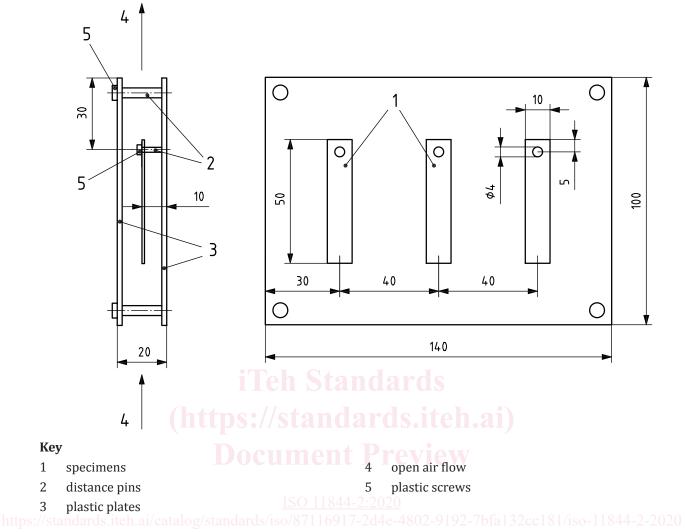


Figure 2 — A mounting plate for the unsheltered exposure of specimens

### Annex A

### (informative)

### **Determination of corrosion rate by mass change measurement**

### A.1 Principle

Mass increase measurements can be performed on all metals and comparatively large surfaces can be evaluated. The technique is relatively easy to operate.

The mass loss determination gives a best estimate of the corrosion effect. The method is not yet applicable to all metals. Both mass increase and mass loss determination using an ultra-micro-balance has a precision of about  $\pm 10 \text{ mg/m}^2$  with the method described below.

Due to the difficulty of distinguishing corrosion effects from other surface-related phenomena, such as sorption and contamination by particulate matter, the specimens should preferably be exposed under shelter.

### A.2 Specimens

It is preferable to use rectangular specimens in the form of flat sheets, as they can be readily weighed. A convenient specimen size is 10 mm  $\times$  50 mm. Specimens may be larger provided that they can be accurately weighed. The specimen thickness may preferably be 0,5 mm.

The materials used to prepare the specimens are of following quality:

Silver: 99,98 % min.

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Copper: standard 99,85 % min.og/standards/iso/87116917-2d4e-4802-9192-7bfa132cc181/iso-11844-2-2020

Zinc: 99,45 % min.

Carbon steel: CR 1, max. 0,15 % C, max. 0,04 % P, max. 0,05 % S, max. 0,6 % Mn

Lead: 99,97 % min.

The specimens should, before weighing, be prepared as follows.

- a) A hole with the diameter 4 mm is cut at the upper side of the specimen.
- b) Abrading:
  - 1) silver and copper with silicon carbide paper to 1 200 P (600 grit);
  - 2) zinc, carbon steel and lead to 500 P (320 grit).

To avoid risk of contamination, an abrading paper shall not be used for polishing specimens of different metals.

- c) Cleaning in de-ionised water.
- d) Degreasing in ethanol in an ultrasonic bath for 5 min.
- e) Drying.