

English Version

**Ambient air quality - Measurement of bioaerosols - Part 1:  
Determination of moulds using filter sampling systems and  
culture-based analyses**

Qualité de l'air ambiant - Mesurage de bioaérosols - Partie  
1: Dosage des moisissures à l'aide de systèmes de  
prélèvement sur filtres et d'analyses de cultures

Luftbeschaffenheit - Messen von Bioaerosolen - Teil 1:  
Bestimmung von Schimmelpilzen mittels Probenahme auf  
Filtern und kulturellem Nachweis

This Technical Specification (CEN/TS) was approved by CEN on 4 October 2010 for provisional application.

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## Foreword

This document (CEN/TS 16115-1:2011) has been prepared by Technical Committee CEN/TC 264 “Air quality”, the secretariat of which is held by DIN.

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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## Introduction

Airborne particles of biological origin are called bioaerosols. Depending on the emission source bioaerosols vary in composition; one component of ambient bioaerosols with possible ecological and health relevance can be moulds. Natural and anthropogenic sources for mould spores are widely distributed in the environment. Anthropogenic sources can for example be agriculture and construction activities or waste treatment.

Mould is a common name for filamentous fungi from different taxonomic groups (Zygomycetes, Ascomycetes, Deuteromycetes). They form a mycelium (hyphae) and spores – namely conidiospores (conidia), sporangiospores or ascospores – by which they become visible macroscopically. Most spores are in the size range of 2 µm to 10 µm, some up to 30 µm and only few up to 100 µm. Spores of some mould genera are small and become airborne very easily (e.g., *Aspergillus*, *Penicillium*) while others are bigger and/or embedded in a slime matrix (e.g., *Stachybotrys*, *Fusarium*) and less mobile.

The procedure described in this document is based on VDI 4252 Part 2 [1], VDI 4253 Part 2 [2] and is related to the ISO standards on indoor air ISO 16000-16 [3] and ISO 16000-17 [4].

## 1 Scope

This Technical Specification describes the measurement of moulds in ambient air in order to identify, quantify and characterize bioaerosol pollution in ambient air resulting from emissions from different sources.

The method described specifies the sampling of moulds as part of the suspended particulate matter (SPM, here particles with aerodynamic diameter up to ca. 30  $\mu\text{m}$ ) using a filter sampling system with gelatine/poly-carbonate filter combination followed by the culture-based analyses on DG18 agar. The sampling duration can be varied between 10 min to 24 h. The health effect of bioaerosols is not limited to any particle fraction, therefore, this document describes the sampling of moulds as part of the suspended particulate matter as a convention method.

**NOTE** The sampling method described in this document in principle is likely to be appropriate for the sampling of actinomycetes and other spore-forming bacteria (resistant to desiccation). For these species a special analytical procedure using different culture media should be applied, but this is not within the scope of this document.

The standard method set out in this Technical Specification is accepted by convention as reference method. The measured quantity, here the number of colony forming units per cubic meter ( $\text{CFU}/\text{m}^3$ ), is determined by the inlet design of the sampling head, the associated operational parameters and the analytical procedure.

Standardized methods for sampling, detection and enumeration of moulds including standards for sampling strategies are important for comparative assessment of moulds in ambient air. Before doing any measurements a plan for the measurement strategy is necessary (see CEN/TS 16115-2 [5]).

**WARNING** — The use of this Technical Specification may involve hazardous materials, operations and equipment. This Technical Specification does not purport to address all the safety problems associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.

## 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 8199:2007, *Water quality — General guidance on the enumeration of micro-organisms by culture (ISO 8199:2005)*

## 3 Terms and definitions

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

### 3.1

#### **aerodynamic diameter**

diameter of a sphere of density 1  $\text{g}/\text{cm}^3$  with the same terminal velocity due to gravitational force in calm air as the particle, under the prevailing conditions of temperature, pressure and relative humidity

[ISO 7708:1995, 2.2 [6]]

### 3.2

#### **ambient air**

outdoor air in the lower troposphere excluding workplace air

[EN 14907:2005, 3.1.1 [7]]

### 3.3

#### **analytical blank value**

value determined by a blank sample covering the analytical procedure to ensure that no significant contamination occurs during the complete analytical procedure including autoclaving, agar preparation, suspension and extraction of the filters, dilution, incubation, counting, etc.