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Rheological test methods - Fundamentals and interlaboratory comparisons - Part 2: Determination of the time-dependent structural change (thixotropy) (ISO/TR 20659-2:2024)

Rheologische Prüfverfahren - Grundlagen und Ringversuch - Teil 2: Thixotropie (ISO/TR 20659-2:2024)

Méthodes d'essai rhéologiques - Principes fondamentaux et comparaisons interlaboratoires - Partie 2: Détermination de la variation structurelle dans le temps (thixotropie) (ISO/TR 20659-2:2024)

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European foreword

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Foreword

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Rheological test methods — Fundamentals and interlaboratory comparisons —

Part 2: Determination of the time-dependent structural change (thixotropy)

1 Scope

This document gives information on an interlaboratory comparison for the determination of the timedependent structural change (thixotropy) using rheological test methods. Thixotropy is the reversible, timedependent decrease of shear viscosity η at a constant shear rate $\dot{\gamma}$ or shear stress τ .

This document provides examples of fields of application, in which important material properties can be characterized by the thixotropy. These fields of application include:

- effectiveness of rheological additives and thixotropic agents, respectively;
- stability of the structure at rest (e.g. behaviour when starting to pump);
- wet film thickness after processing;
- levelling and sagging behaviour (e.g. without brushmarks or sag formation);
- orientation of effect pigments.

2 Normative references SIST-TP CEN ISO/TR 20659-2:2025

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements 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 3219-1, Rheology — Part 1: Vocabulary and symbols for rotational and oscillatory rheometry

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO 3219-1 apply.

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

- ISO Online browsing platform: available at https://www.iso.org/obp
- IEC Electropedia: available at <u>https://www.electropedia.org/</u>

4 Measuring technique for the determination of thixotropy

4.1 Conditions for the measuring technique

<u>Clause 4</u> briefly describes methods that are currently in use. In principle, the thixotropy depends on the temperature, the pressure, and the thermal and mechanical history of the material. A detailed specification of the measuring profile is therefore a precondition for reproducible measurements and comparable

evaluation; this applies especially for the level of shear load (shear rates, shear stresses, shear strain, oscillation frequencies), the duration of the individual measuring segments and the number of measuring points.

Thixotropy can be determined by rotational as well as by oscillatory tests. Measuring devices equipped with a mechanical bearing or air bearing are suitable for rotational tests. For oscillatory tests, a rheometer with air bearing is used. It is essential to ensure that the measuring device is used in combination with the suitable measuring geometry, i.e. in accordance with the torque range, the torque resolution and the rotational speed range. Typically, rotational viscometers and rheometers that are subject to test equipment monitoring and are regularly calibrated and verified (if necessary), are used. Measuring results that are independent of geometry can only be obtained by using absolute measurement geometries according to ISO 3219-2.

If, independently of the measuring device and the measuring method used, no time-stable measuring results are obtained during the measurement of Newtonian standard samples, then the measuring device, the measuring geometry or the measuring method is unsuitable. If the functional course of these time-stable measuring device, the measuring geometry and the measuring method are suitable for the investigation of the sample. Typically, this inspection is carried out under isothermal conditions in the expected viscosity range of the sample by using several Newtonian standard samples. If preliminary tests reveal that the viscosity of the measuring sample varies over three decades, then the verification of the measuring device and measuring geometry are performed with three Newtonian standard samples. This is carried out for all measuring temperatures. The influences due to application, e.g. sample filling, evaporation, shear heating, wrong choice of method and the sample material coming out of the gap, are mostly discernible and detected by this kind of verification.

Upon measurement, the possibility of evaporation is considered. A reduction of this influence can be reached, e.g. by using a sample area coverage. All boundary conditions of the measurement are documented in the record, especially the usage of a coverage, the kind of sample trimming and the adjustment of the gap distance. According to the specifications of the measuring methods described in Clause 4, the methods are changed if influences on the measuring results occur. Another parameter is checking whether the duration of the load is shorter than the medium time scale of the structural changes of the sample. This is determined for each measuring method and its specifications by preliminary tests. However, the duration of the load is selected in consideration of situations where conditions of application of the sample are longer. If this duration is longer than the time scale of structural change of the sample, then a thixotropic behaviour will not be detected; nevertheless, the sample can be still thixotropic. In order to determine the thixotropy in a correct and reproducible manner, when filling the measuring geometry, the influence that the time between filling and the start of the measurement has on the measuring result is taken into consideration. This time is distinctly longer than the time scale of the structural change. This can be determined by preliminary measurements in which the waiting time is varied. The waiting time is sufficiently high if the thixotropy of the sample is comparable for two measurements running after a fresh filling within the limits of the requested precision in accordance with the measuring method chosen.

Measuring points can only be recorded if each single measuring point is controlled by the instrument according to the specification. At every change of the specified value, a transient process of the entire measuring equipment occurs towards the new specified value. This transient process can be different (see Figure 1).