INTERNATIONAL STANDARD

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Coal — Determination of plastic properties — Constant-torque Gieseler plastometer method

Charbon — Détermination des propriétés plastiques — Méthode du plastomètre Gieseler à couple constant

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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 www.iso.org/directives).

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. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see www.iso.org/patents).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation on the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT) see the following URL: www.iso.org/iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 27, *Solid mineral fuels*, Subcommittee SC 5, *Methods of analysis*.

This second edition cancels and replaces the first edition (ISO 10329:2009), which has been technically revised.

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Coal — Determination of plastic properties — Constanttorque Gieseler plastometer method

1 Scope

This document specifies a method for obtaining a relative measure of the plastic behaviour of coal when heated under prescribed conditions. The method is used to obtain values of the plastic properties of coals and blends used in carbonization and in other situations where determination of plastic behaviour of coals is of practical importance.

NOTE The empirical nature of this test requires proper equipment calibration to produce fluidity readings which are a true indication of the relative plastic behaviour of the coal.

2 Normative references

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 18283, Hard coal and coke — Manual sampling

ISO 13909-1, Hard coal and coke — Mechanical sampling — Part 1: General introduction

ISO 13909-2, Hard coal and coke — Mechanical sampling — Part 2: Coal — Sampling from moving streams

ISO 13909-3, Hard coal and coke — Mechanical sampling — Part 3: Coal — Sampling from stationary lots

3 Terms and definitions

<u>ISO 10329:2017</u>

tps://standards.iteh.ai/catalog/standards/iso/66751511-bfaf-4ba4-a24e-c941ed11882e/iso-10329-2017 For the purposes of this document, the following terms and definitions apply.

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

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

3.1

dial division per minute

measure of stirrer rotation rate, as used in the constant-torque Gieseler plastometer method

Note 1 to entry: There are 100 dial divisions for each full 360° rotation of the stirrer. The fluidity result is expressed as total dial divisions turned by the stirrer in a 1 min time period, i.e. dd/min.

3.2

initial softening temperature

temperature at which dial movement or electronic readout indicates a stirring shaft movement of one *dial division per minute* (3.1), with continued indication of movement of at least 1 dd/min thereafter

3.3

maximum fluidity temperature

temperature at which stirring shaft rotation reaches the maximum rate

3.4

plastic range

difference between the initial softening temperature and the solidification temperature

3.5

final fluidity temperature

temperature at which the last 1 dd/min stirrer rotation rate is reached

3.6

solidification temperature

temperature at which the stirring shaft stops

3.7

maximum fluidity

maximum rate of rotation for the stirring shaft in *dial divisions per minute* (3.1)

3.8

jamming

swelling up of coal into the retort tube during the test, which may produce a lower fluidity result than expected and can only be noted after visual inspection of the disassembled crucible and retort at the conclusion of the test

3.9

breaking

free spinning behaviour of coal, either by rotating at maximum motor speed or by abrupt changes in rotation, which occurs as a result of a molten ball of coal forming around the base of the stirrer, and which makes reporting of the true *maximum fluidity* (3.7) of the coal difficult

4 Principle

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Measurements of the plastic properties of coals are made by applying a constant torque to a stirrer placed in a crucible into which the coal is charged. The crucible is immersed in a bath and the temperature increased uniformly. The rotation of the stirrer is recorded in relation to increase in temperature.

5 Apparatus

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https://standards.iteh.ai/catalog/standards/iso/66751511-bfaf-4ba4-a24e-c941ed11882e/iso-10329-2017 **5.1** Gieseler plastometer retort, composed of the following component parts (see Figure 1).

5.1.1 Retort crucible, cylindrical, with $(21,4 \pm 0,1)$ mm inside diameter, and $(35,0 \pm 0,3)$ mm in depth with exterior threads for joining the crucible to the barrel.

The crucible shall have a $(2,38 \pm 0,02)$ mm diameter notch with an included angle of 70° in the centre of its inside base to serve as a seat for the stirrer.

5.1.2 Retort crucible cover, with interior threads for joining the crucible cover to the crucible and exterior threads for joining the crucible cover to the barrel.

The inside diameter of the hole which accommodates the stirrer shall be $(9,5 \pm 0,1)$ mm.

5.1.3 Guide sleeve, provided near the upper end of the stirrer to guide the latter within the barrel with a clearance of between 0,05 mm and 0,10 mm.

5.1.4 Gas exit hole, provided on the barrel to afford an exit for the volatile products during a test, placed, for example, at the midpoint of the barrel; as an option, a tube may be fitted if desired.

5.1.5 Barrel, (121,0 ± 2,5) mm long, having an inside diameter of (9,5 ± 0,1) mm.

The top end of the barrel shall be 12,7 mm in inside diameter to a depth sufficient to allow the fitting of a guide sleeve through which the axle of the stirrer passes when the apparatus is assembled.