

SLOVENSKI STANDARD oSIST prEN ISO 21656:2020

01-marec-2020

Trdna alternativna goriva - Določevanje vsebnosti pepela (ISO/DIS 21656:2019)

Solid recovered fuels - Determination of ash content (ISO/DIS 21656:2019)

Feste Sekundärbrennstoffe - Bestimmung des Aschegehaltes (ISO/DIS 21656:2019)

Combustibles solides de récupération - Détermination de la teneur en cendres (ISO/DIS 21656:2019)

Ta slovenski standard je istoveten z: prEN ISO 21656

ICS: Trda goriva 75.160.10 Solid fuels

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en,fr,de

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DRAFT INTERNATIONAL STANDARD ISO/DIS 21656

ISO/TC 300

Voting begins on: **2019-12-19**

Secretariat: SFS

Voting terminates on: 2020-03-12

Solid recovered fuels — Determination of ash content

Combustibles solides de récupération — Détermination de la teneur en cendre

ICS: 75.160.10

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Reference number ISO/DIS 21656:2019(E)

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Published in Switzerland

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Foreword

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For an explanation of 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 <u>www.iso.org/</u> iso/foreword.html.

This document was prepared by Technical Committee ISO/TC 300, Solid recovered fuels.

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>.

ST EN ISO 21656:20

Introduction

This document covers the determination of ash content of solid recovered fuels. It is primarily geared toward laboratories, producers, suppliers and purchasers of solid recovered fuels but is also useful for the authorities and inspection organizations.

The method specified in this document is based on EN 15403.

For information about environmental aspect, see <u>Annex B</u>.

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DRAFT INTERNATIONAL STANDARD

Solid recovered fuels — Determination of ash content

1 Scope

This document specifies methods for the determination of ash content of all solid recovered fuels.

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.

EN 15442,¹⁾Solid recovered fuels — Methods for sampling

EN 15443,²)*Solid recovered fuels* — *Methods for the preparation of the laboratory sample*

ISO/DIS 21637, Solid recovered fuels — Terminology, definitions and descriptions

ISO/DIS 21660-3, Solid recovered fuels — Determination of moisture content using the oven dry method — Part 3: Moisture in general analysis sample

3 Terms and definitions

For the purposes of this document, the terms and definitions given in ISO/DIS 21637 and the following apply.

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>

standards.iteh.ai/catalog/standards/sist/49d4536a-b80444991-833d-c68aa8e87ae1/sist-en-iso-21656-2021 — IEC Electropedia: available at <u>http://www.electropedia.org/</u>

3.1

ash content on dry basis

mass of inorganic residue remaining after ignition of a fuel under specified conditions, expressed as mass fraction in percent of the dry matter in the fuel

Note 1 to entry: For the characterisation of solid recovered fuels (SRF) it is necessary to analyse the content of the volatile alkaline and therefore the temperature of 550°C is used for biomass SRF.

3.2

total ash content on dry basis

mass of inorganic residue remaining after ignition of a fuel under specified conditions, expressed as mass fraction in percent of the dry matter in the fuel, which also includes removed ash contributors

¹⁾ ISO/CD 21645:2019 "Solid recovered fuels — Method for sampling" is currently being processed for the preparation of the "DIS"-enquiry.

²⁾ ISO/CD 21646:2019 "Solid recovered fuels — Sample preparation" is currently being processed for the preparation of the "DIS"-enquiry.

3.3

removed ash contributors

rac

coarse inert material (i.e. metals, glass, stones, tiles etc.) removed from the pre-dried sample before preparation, in order to avoid damage to the preparation equipment,

Note 1 to entry: Removed ash contributors are included in the ash content calculations.

3.4

total organic matter

combustible part of solid recovered fuels, which consists of the sum of volatile matter and fixed carbon

Note 1 to entry: It is calculated as: 100 - moisture content - ash content.

Note 2 to entry: It is the mass fraction of the matter lost by ignition, also known as "Loss of Ignition" (LOI).

3.5

volatile matter

relative part of the analysed sample, after moisture removal, that is lost when material is heated up under specific conditions of temperature, time and in a reduced atmosphere (anoxic conditions)

3.6

fixed carbon

relative part of carbon contained in a material that can only be degraded in oxic conditions and high temperature

Note 1 to entry: It is calculated as: 100 - moisture content - volatile matter content - ash content.

4 Principle

The sample is heated in air atmosphere up to a temperature of (550 ± 10) °C for Method A or (815 ± 10) °C for Method B under rigidly controlled conditions of time, sample mass and equipment specifications. The ash content is determined by calculation from the mass of the residue remaining after heating.

NOTE tan Difference in the ash content if determined at a higher temperature, 815 °C, according to Reference^[1] as 56-2021 compared to 550 °C (major and minor elemental determination), is explained by the decomposition of carbonates forming CO2, by losses of volatile inorganic compounds and further oxidation of inorganic compounds (to higher oxidation states).

Automatic equipment (such as thermogravimetric analysers) may be used as long as the equipment is validated by parallel measurements to the reference method. The automatic equipment shall fulfil all the requirements regarding sample size, heating procedure, temperature, atmosphere and weighing accuracy. Deviations from this paragraph shall be reported and justified.

5 Apparatus

5.1 Dish, consisting of inert material such as porcelain, silica or platinum, with a depth from 10 mm to 20 mm and such a size that the sample loading does not exceed 0,1 g/cm² bottom area.

5.2 Furnace, capable of maintaining a zone of uniform temperature at the levels required in <u>Clause 7</u> and to reach these levels in the specified heating rates. The ventilation rate through the furnace should be such that no lack of oxygen arises during the heating procedure.

NOTE A ventilation rate from 5 air changes/min to 10 air changes/min should be suitable.

5.3 Balance, capable of weighing the dish containing the sample to the nearest 0,1 mg.

5.4 Desiccator, without desiccant.