
**Plastics — Determination of aerobic
biodegradation of non-floating plastic
materials in a seawater/sediment
interface — Method by analysis of
evolved carbon dioxide**

*Plastiques — Détermination de la biodégradation aérobie des
matières plastiques non-flottantes dans une interface eau de mer/
sédiments — Méthode par analyse du dioxyde de carbone libéré*





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ISO copyright office
CP 401 • Ch. de Blandonnet 8
CH-1214 Vernier, Geneva
Phone: +41 22 749 01 11
Fax: +41 22 749 09 47
Email: copyright@iso.org
Website: www.iso.org

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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 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 www.iso.org/iso/foreword.html.

The committee responsible for this document is ISO/TC 61, *Plastics*, Subcommittee SC 14, *Environmental aspects*, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 249, *Plastics*, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

This second edition cancels and replaces the first edition (ISO 19679:2016), which has been technically revised.

The main changes compared to the previous edition are as follows:

- in [Annex A](#): Density of O₂ in air at 1 atm, 28 °C and a relative humidity of 100 % has been corrected and the subsequent calculations have been adapted accordingly.

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 www.iso.org/members.html.

Introduction

Products made with biodegradable plastics are designed to be recovered by means of organic recycling in composting plants or in anaerobic digesters. The uncontrolled dispersion of biodegradable plastics in natural environments is not desirable. The biodegradability of products cannot be considered as an excuse to spread wastes that should be recovered and recycled. However, test methods to measure rate and level of biodegradation in natural environments (such as soil or the marine environment) are of interest in order to better characterize the behaviour of plastics in these very particular environments. As a matter of fact, some plastics are used in products that are applied in the sea (e.g. fishing gear) and sometimes they can get lost or put willingly in the marine environment. The characterization of biodegradable plastic materials can be enlarged by applying specific test methods that enable the quantitative assessment of biodegradation of plastics exposed to marine sediment and seawater. Plastic products are directly littered or arrive with fresh waters in the pelagic zone (free water). From there, and depending on density, tides, currents, and marine fouling plastics can sink to the sublittoral, and reach the seafloor surface. Many biodegradable plastics have a density higher than 1 and therefore tend to sink. The sediment passes from aerobic to anoxic and finally anaerobic conditions going from the surface (the interface with seawater) into deeper layers, displaying a very steep oxygen gradient.

Plastics — Determination of aerobic biodegradation of non-floating plastic materials in a seawater/sediment interface — Method by analysis of evolved carbon dioxide

1 Scope

This document specifies a test method to determine the degree and rate of aerobic biodegradation of plastic materials when settled on marine sandy sediment at the interface between seawater and the seafloor, by measuring the evolved carbon dioxide (CO₂). This test method can also be applied to other solid materials.

This test method is a simulation under laboratory conditions of the habitat found in different seawater/sediment-areas in the sea, e.g. in a benthic zone where sunlight reaches the ocean floor (photic zone) that, in marine science, is called sublittoral zone

The determination of biodegradation of plastic materials and other solid materials buried in marine sediment is outside the scope of this document.

NOTE Measurement of aerobic biodegradation can also be obtained by monitoring the oxygen consumption, as described in ISO 18830.

The conditions described in this document do not always correspond to the optimum conditions for the maximum degree of biodegradation to occur.

2 Normative references

There are no normative references in this document.

3 Terms and definitions

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:

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

3.1

theoretical amount of evolved carbon dioxide

ThCO₂

maximum theoretical amount of carbon dioxide evolved after completely oxidising a chemical compound, calculated from the molecular formula or from determination of *total organic carbon (TOC)* (3.2)

Note 1 to entry: It is expressed as mg of carbon dioxide evolved per mg or g of test compound.

3.2

total organic carbon

TOC

amount of carbon bound in an organic compound

Note 1 to entry: Total organic carbon is expressed as mg of carbon per 100 mg of the compound.

3.3
dissolved organic carbon
DOC

part of the organic carbon in water which cannot be removed by specified phase separation methods, for example by centrifugation at 40 000 ms⁻² for 15 min or by membranes with pores of 0,2 µm to 0,45 µm diameter

3.4
pre-conditioning phase
pre-incubation of an inoculum under the conditions of the subsequent test in the absence of test material, with the aim to consume potential organic matter present in excess that could disturb biodegradation measurement and to improve the acclimatization of the microorganisms to the test conditions

4 Principle

This test method is based on the determination of evolved CO₂ and derives from ISO 14852. The testing medium is based on a solid phase and a liquid phase. The solid phase is a sandy marine sediment laid in the bottom of a closed flask; the liquid phase is a column of natural or artificial sea water, poured on the sediment. The test material is preferably in the form of a film to be laid down on top of the sediment, at the interface between the solid phase and the liquid phase. This is a simulation of an object that has sunk and finally reached the sea floor. The system is contained in a closed flask.

The CO₂ evolved during the microbial degradation is determined by a suitable analytical method. The level of biodegradation is determined by comparing the amount of CO₂ evolved with the theoretical amount (ThCO₂) and expressed in percentage. The test result is the maximum level of biodegradation, determined from the plateau phase of the biodegradation curve. The principle of a system for measuring evolved CO₂ is given in ISO 14852:2018, Annex A.

The details of interlaboratory testing based on the test method specified in this document are available in Reference [6].

5 Test environment

Incubation shall take place in the dark or in diffuse light in an enclosure which is free from vapours inhibitory to microorganisms and which is maintained at a constant temperature, preferably between 15 °C to 25 °C, but not exceeding 28 °C, to an accuracy of ± 2 °C. Any change in temperature shall be justified and clearly indicated in the test report.

NOTE Temperatures applied in the test can be different from those found in marine environments

6 Reagents

6.1 **Distilled or deionized water**, free of toxic substances (copper in particular) and containing less than 2 mg/l of DOC.

6.2 **Artificial seawater.**

Dissolve:

Sodium chloride (NaCl)	22 g
Magnesium chloride hexahydrate (MgCl ₂ · 6H ₂ O)	9,7 g
Sodium sulfate (Na ₂ SO ₄)	3,7 g

Calcium chloride (CaCl_2)	1 g
Potassium chloride (KCl)	0,65 g
Sodium hydrogen carbonate (NaHCO_3)	0,20 g
in water (6.1) and make up to 1 000 ml.	

6.3 Natural seawater/sediment.

Take a sample of a sandy sediment and seawater with a shovel beneath the low-water line into a bucket. Transfer the wet sediment together with seawater into sealed containers for transport and fast deliver it to the laboratory. After delivery, conserve the sediment at low temperature (approximately 4 °C) until use. The seawater/sediment sample should be preferably used within 4 weeks after sampling. Record storage time and conditions.

NOTE Seawater and sediment can also be sampled from large, well-running public marine aquaria.

Measure the TOC, pH and nitrogen content of the sediment and of the natural seawater if used instead of artificial seawater. The carbon content of sediment should be in the range of 0,1 % to 2 %.

A preliminary oxidation can be applied to the sediment in order to decrease the organic matter content and the background respiration. Sediment and seawater are fluxed with air and gently stirred (max. 20 r/min to 30 r/min) in a large container for the desired period of time. Include this pre-treatment process in the test report.

7 Apparatus

7.1 Test flasks.

Biometer flasks of the volume of about 250 ml are appropriate. Reactors with higher volumes can be used, if test conditions are not affected. The vessels shall be located in a constant-temperature room or in a thermostatic apparatus (e.g. water-bath). Stirring can be applied on seawater on condition that it does not disturb the sediment/seawater interface.

NOTE A suitable apparatus is shown in [Figure A.1](#). An example of a stirred apparatus is given in OECD TG 308: 2002, Annex 4^[2].

7.2 Container for the CO_2 absorber.

A glass beaker to be located in the headspace of the reactor and filled with 10 ml of $\text{Ba}(\text{OH})_2$ 0,0125 mol/l or with 3 ml of KOH 0,5 mol/l.

7.3 Analytical balance, shall have a sensitivity of at least 0,1 mg.

7.4 pH meter.

8 Procedure

8.1 Test material

The test material should be in film or sheet form. Cut samples of the test material in the shape of a disk. Disks shall have a smaller diameter than the glass flasks, so that the disks can be easily laid on the bottom of the glass flask.

The sample shall be of known mass and contain sufficient carbon to yield CO_2 that can be adequately measured by the system used.

Use a test material concentration of at least 100 mg/l of seawater plus sediment. This mass of the sample should correspond to a TOC of about 60 mg/l. The maximum mass of sample per flask is limited by the oxygen supply to the glass flask. A test material concentration of 150 mg/l to 300 mg/l of seawater plus sediment is recommended.

Calculate the TOC from the chemical formula or determine it by a suitable analytical technique (e.g. elemental analysis or measurement in accordance with ISO 8245) and calculate the ThCO_2 .

The form and shape of the test material may influence its biodegradation. Similar shapes and thicknesses should preferably be used if different kinds of plastic materials are to be compared.

NOTE When the test material in form of film is laid down on the surface of the sediment, it can limit the gas exchange between the water body and the sediment, promoting the formation of anaerobic zones under the test material. In order to reduce this effect, it is possible to perforate the film sample homogeneously over the entire surface.

8.2 Reference material

Use ashless cellulose filters as a reference material¹⁾. If possible, the TOC, form, and size should be comparable to that of the test material. As a negative control, a non-biodegradable polymer (such as polyethylene) in the same form as the test material shall be used.

8.3 Preparation of the sediment

Filter the sediment in a funnel with a coarse filter paper to eliminate excess seawater. Sediment is ready for testing when dripping of sea water is ended. Sediment after filtering is named "wet sediment" hereafter.

8.4 Test setup

Provide several flasks, so that the test includes at least the following:

- a) three flasks for the test material (symbol F_T);
- b) three flasks for the blank (symbol F_B);
- c) three flasks for reference material (symbol F_C);
- d) three flasks for negative control (symbol F_N).

Two flasks for test material, blank, reference material, and negative control may be used instead of three for screening purposes.

8.5 Pre-conditioning phase

In a typical case, use a test flask with a volume of 250 ml. Put 30 g of the wet sediment on the bottom of the flask. Carefully pour 70 ml of natural or artificial seawater. Reactors with higher volumes can be used, if test conditions are not affected. The test should be performed with a water/sediment volume ratio between 3:1 and 5:1 and a sediment layer of about 0,3 cm to 0,5 cm, depending on the granulometry of the sediment.

When using very coarse-grained sediment, the layer can be increased up to 1,5 cm.

Add CO_2 absorber to the absorber compartments of the test flask in a typical case 3 ml of KOH 0,5 mol/l or 10 ml of $\text{Ba}(\text{OH})_2$ 0,0125 mol/l. Place the flasks in a constant-temperature environment and allow all vessels to reach the desired temperature. Take the necessary readings and monitor the CO_2 evolution.

1) Laboratory filter paper Whatman n° 42 has been found satisfactory for this purpose and is an example of a suitable product available commercially. This information is given for the convenience of users of this document and does not constitute an endorsement by ISO of this product.

This phase is carried out in order to verify that the endogenous respiration is similar in the different vessels and also to obtain a preliminary oxidation of excess organic matter, in order to start the test with a lower endogenous respiration. The inoculum can be gently stirred in order to accelerate the biodegradation of excess organic matter.

This phase is generally protracted for a week but is possible to extend this time if a high amount of CO₂ evolved is measured.

In case the CO₂ evolution of a vessel is different, reject the diverging vessel or in case of multiple anomalies, start again using new sediment.

8.6 Start of the test

Dunk the plastic film sample, cut as described in 8.1, on the sediment of each vessel. Mass of samples (test and reference material) should be about 20 mg each when using a flask with a volume of 250 ml corresponding to an initial test item concentration specified in 8.1. In order to ensure a homogeneous contact between sample and sediment, it is recommended to cover the sample with a suitable cover slip. The cover slips shall also be introduced in blank vessels, for assuring similar conditions.

NOTE A suitable cover slip can be made using a common non-biodegradable vinyl-coated fibreglass mosquito net with a fibre diameter of about 280 µm and a 1,8 mm × 1,6 mm mesh.

For an example, see [Annex A](#).

Repeat the procedure for the reference material and the material for the negative control to the respective flasks. Record the mass of the sediment, the sample and the volume of seawater introduced in each vessel.

Nutrients may be supplemented as needed to support microbial diversity and to maintain the capacity to biodegrade the test material. The need and timing of additional nutrients or other appropriate measures may be judged by observation of the temporal course of the biodegradation of the reference substance cellulose. Any addition and the applied method shall be reported in the test report.

8.7 Carbon dioxide measurement

8.7.1 The CO₂ reacts with Ba(OH)₂ and precipitates as BaCO₃. The amount of CO₂ produced is determined by titrating the remaining Ba(OH)₂ with 0,05 mol/l HCl to a phenolphthalein end-point or by automatic titrator. Because of the static incubation, the BaCO₃ builds up on the surface of the liquid and shall be broken up periodically by shaking the container gently to ensure continued absorption of the evolved CO₂. This problem can be avoided by using KOH instead of Ba(OH)₂, which does not form a precipitate.

NOTE A discussion on the use of KOH in place of Ba(OH)₂ is reported in Reference [5].

8.7.2 The containers for the CO₂ absorber shall be removed and titrated before their capacity is exceeded. The period of time will vary with sediments and test materials and increases slowly as the carbon content of the sediment is reduced (a recommended frequency of every 3 to 4 days for the first 2 to 3 weeks and every 1 to 3 weeks, thereafter). At the time of removal of the containers, the reactor should be allowed to sit open so that the air is refreshed before replacing 10 ml of fresh Ba(OH)₂ and resealing the reactor. The reactors should remain open approximately 15 min.

8.7.3 The CO₂ evolution rate may reach a plateau when all of the accessible carbon has been oxidized. The test may be terminated at this point or earlier, at the discretion of the user. If possible, the residual test material may be extracted from the sediment with an appropriate method and quantified (optional).

NOTE The evolved CO₂ can be quantitatively measured also using other suitable methods such as those based on infrared CO₂-analysers or those based on TOC analysers equipped with an infrared photometer or on gravimetric analysis.

8.8 End of the test

When a constant level of CO₂ evolution is attained (plateau phase reached) and no further biodegradation is expected, the test is considered to be completed. The maximum test period is 24 months. In the case of long test durations, special attention shall be paid to the technical system (e.g. tightness of the test vessels and connections). Any special measures taken, for example, to ensure microbial diversity or to provide sufficient nutrients shall be detailed in the test report. On the last day of the test, measure the pH, acidify all the bottles with 1 ml of concentrated HCl in order to decompose the carbonates and bicarbonates, continue the test for 24 h and finally measure the amount of CO₂ evolved in each of the series of flasks.

9 Calculation and expression of results

9.1 Calculation

9.1.1 Amount of CO₂ produced

9.1.1.1 Net CO₂ produced

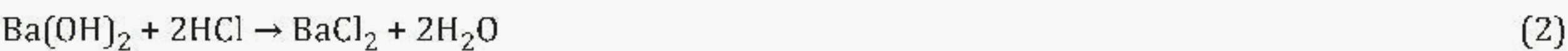
The first step in calculating the amount of CO₂ produced is to correct the test material reactors for endogenous CO₂ production. The control reactor serves as a blank to correct for CO₂ which may be produced through endogenous respiration of the microorganisms. The amount of CO₂ produced by a test material is determined by the difference (in ml of titrant) between the experimental and blank containers. The next step is to convert ml of HCl titrated into mg of CO₂ produced.

9.1.1.2 Ba(OH)₂ used as CO₂ absorber

When CO₂ enters the absorber containers, it reacts in the following manner:



The BaCO₃ formed is insoluble and precipitates. The amount of Ba(OH)₂ remaining in the solution is determined by titration of the 10 ml of the CO₂ absorber with HCl according to the following chemical reaction:



The amount of the remaining Ba(OH)₂ is determined with the [Formula \(3\)](#):

$$R_t = \frac{\text{mol HCl}}{2} \tag{3}$$

where R_t is the remaining Ba(OH)₂.

The amount of the reacted Ba(OH)₂ is determined as the difference between the amount originally present in the absorber and the amount remaining after reaction with CO₂, see [Formula \(4\)](#).

$$R_r = R_0 - R_t \tag{4}$$

where

- R_r is the amount of reacted Ba(OH)₂;
- R_0 is the amount of Ba(OH)₂ originally present in the absorber;
- R_t is the amount of Ba(OH)₂ remaining after the reaction with CO₂.

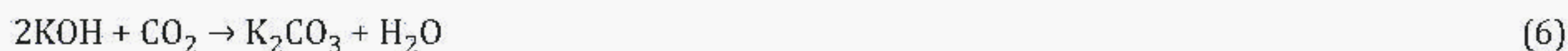
This means that the number of mol of CO₂ produced is derived using [Formula \(5\)](#):

$$\text{mol CO}_2 = R_r \quad (5)$$

where R_r is the amount of reacted Ba(OH)₂.

9.1.1.3 KOH used as CO₂ absorber

The evolved CO₂ reacts with KOH in the following manner:



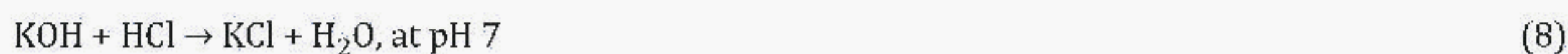
K₂CO₃, the product of [Formula \(6\)](#), is soluble and does not precipitate.

The fresh KOH solution, where no CO₂ has been absorbed, can be titrated with HCl as:



The KOH solutions used as CO₂ absorbers will have both unreacted KOH and K₂CO₃ as per [Formula \(6\)](#).

During titration, both chemical species will react with HCl, as follows:



The pH shifts in [Formulae \(6\)](#) and [\(7\)](#) are superimposed and cannot be distinguished. Only a single end point in the range of pH between 7 and 8, corresponding to the two reactions, can be identified by using a suitable indicator.

The adsorbed CO₂ can be determined by subtracting from the H⁺ equivalents needed to neutralize the original KOH solution and the H⁺ equivalents needed to neutralize the reactions represented by [Formulae \(8\)](#) and [\(9\)](#), as shown in [Formula \(10\)](#). In practice:

$$\text{mmol CO}_2 = (V_{\text{HCl}(7)} - V_{\text{HCl}(8+9)}) \times [\text{HCl}] \quad (10)$$

where

$V_{\text{HCl}(7)}$ is the volume of HCl (expressed in ml) consumed in [Formula \(7\)](#);

$V_{\text{HCl}(8+9)}$ is the volume of HCl (expressed in ml) consumed in [Formulae \(8\)](#) and [\(9\)](#);

$[\text{HCl}]$ is the concentration of the HCl solution (0,05 mol/l).

If an end point titrator is available, the mmol of CO₂ can be determined, without using an indicator, with a further reaction. A further addition of HCl makes HCl react with KHCO₃, produced with [Formula \(9\)](#):



The number of equivalent consumed in [Formula \(11\)](#), and therefore in [Formula \(9\)](#), corresponds to the K₂CO₃ produced during [Formula \(6\)](#), that in turn corresponds to the absorbed CO₂.

Consequently, 1 mole of KHCO₃ corresponds to 1 mole of CO₂ reacted in [Formula \(6\)](#), and thus the mmol CO₂ are equivalent to the mmol HCl consumed in [Formula \(11\)](#) end point

Therefore, the amount of CO₂ can be calculated using [Formula \(12\)](#):

$$mmol\ CO_2 = (V_{HCl(11)}) \times [HCl] \tag{12}$$

where

$V_{HCl(11)}$ is the volume of HCl (expressed in ml) consumed in [Formula \(11\)](#);

$[HCl]$ is the concentration of the HCl solution (0,05 mol/l).

The amount of CO₂ expressed in mg is finally obtained using [Formula \(13\)](#):

$$mg\ CO_2 = mmol\ CO_2 \times 44 \tag{13}$$

where 44 is the molecular weight (g/mol) of CO₂.

9.1.2 Percentage of biodegradation

The percentage of biodegradation is the ration between the evolved CO₂ and theoretical CO₂ (ThCO₂). The ThCO₂ is shown in [Formula \(14\)](#) and percentage of biodegradation in [Formula \(15\)](#):

$$ThCO_2 = S \times TOC(\%) \times \frac{44}{12} \tag{14}$$

where

S is the amount of specimen (mg)

$TOC(\%)$ is the TOC of the plastic material (or reference material) divided by 100;

44 is the molecular weight (g/mol) of CO₂;

12 is the molecular weight (g/mol) of C.

Therefore:

$$\%B = \frac{CO_2}{ThCO_2} \times 100 \tag{15}$$

where

$\%B$ is the percentage of biodegradation

CO₂ is the CO₂ produced, expressed in mg;

ThCO₂ is the theoretical amount of evolved CO₂.

9.2 Visual inspection

At the end of the test, check the condition of the samples. If still present, samples can be retrieved for mass determination, other analysis, and photographs.

9.3 Expression and interpretation of results

Compile a table of the CO₂ values measured and the percentages of biodegradation for each measurement interval and each test flask. For each vessel, plot an evolved CO₂ cumulative curve and a biodegradation curve in percentage as a function of time.

A curve of mean biodegradation values may be plotted.

The maximum level of biodegradation determined as the mean value of the plateau phase of the biodegradation curve or the highest value, e.g. when the curve decreases or, further on, slowly increases in the plateau phase, characterizes the degree of biodegradation of the test material.

The wettability and the shape of the test material may influence the result obtained, and hence the test procedure may be limited to comparing plastic materials of similar chemical structure.

Information on the toxicity of the test material may be useful in the interpretation of test results showing a low biodegradability.

10 Validity of results

The test is considered valid if,

- a) the degree of biodegradation of the reference material (F_C) is $> 60\%$ after 180 days,
- b) the evolved CO_2 of the blank F_B at the end of the test does not exceed $3,5 \text{ mg CO}_2/\text{g wet sediment}$ (see 8.3) after 6 months;

NOTE This value has been determined in an interlaboratory test.

- c) the amount of CO_2 evolved from the three blank F_B are within 20% of the mean at the plateau phase or at the end of the test;
- d) the difference between the percentage biodegradation of the reference material in the different vessels is less than 20% of the mean at the end of the test;
- e) the percentage of biodegradation of the negative control (flasks F_N) is below 10% at the end of the test.

If these criteria are not fulfilled, repeat the test using another sediment.

11 Test report

The test report shall contain at least the following information:

- a) a reference to this document, i.e. ISO 19679:2020;
- b) all information necessary to identify the test and reference materials, including their TOC, ThCO_2 , chemical;
- c) the main test parameters, including test volume, test medium used, incubation temperature and final pH;
- d) the source and amount of the marine sediment used;
- e) any deviations from the procedure;
- f) any unusual features observed;
- g) the date of the test;
- h) the analytical techniques used, including the principle of the respirometer and the TOC;
- i) all the test results obtained for the test and reference materials (in tabular and graphical form), including the evolved CO_2 , the percentage biodegradation values;
- j) the duration of the lag phase, biodegradation phase and maximum level of degradation, as well as the total test duration; and, optionally, if run or determined, the negative control F_N ;

- k) any other relevant data (e.g. result of the visual final inspection and analysis of final samples, if still retrievable; photos of the final samples);
- l) details of the methods used during the test period in order to support microbial diversity or to avoid nutrient deficiency.

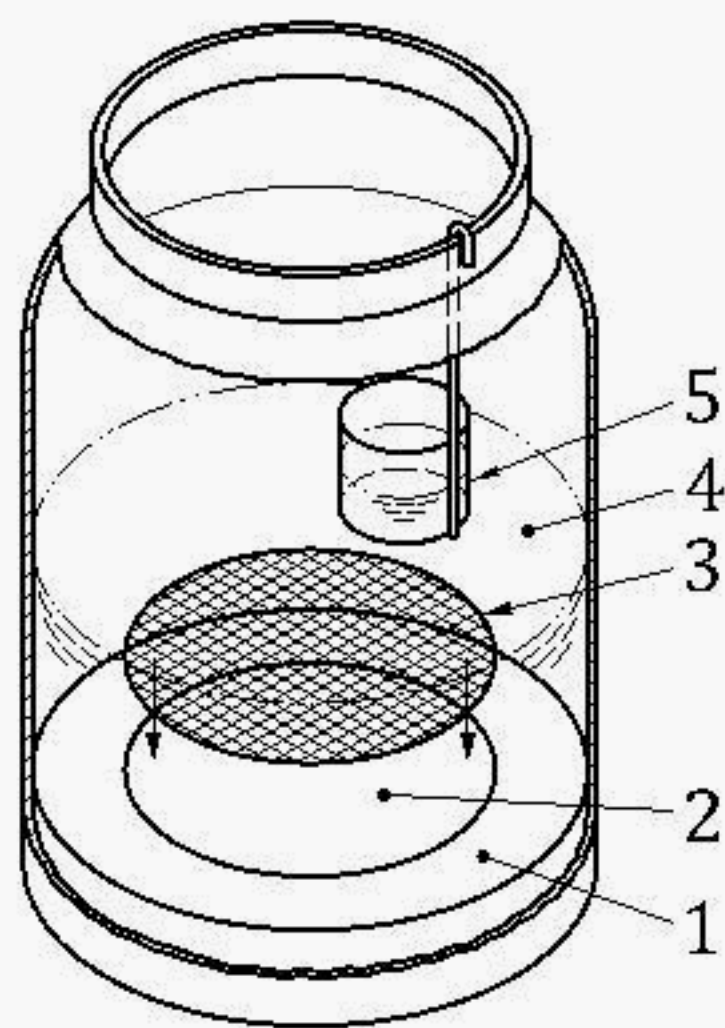
Annex A
(informative)

Example of respirometric system based on CO₂ measurement

The measurement of the evolved CO₂ can be obtained by trapping the evolved CO₂ in a closed system and then by quantifying it by suitable titration systems. Microorganisms in the vessel consume the oxygen and form CO₂. This is absorbed by a CO₂ absorber (generally NaOH) that can then be titrated to determine the amount of the absorbed CO₂.

In a typical case, a 250 ml vessel is used. The sediment occupies about 20 ml, the seawater 70 ml and the headspace 160 ml. The O₂ present in air at 1 atm and 28 °C and a relative humidity of 100 % is about 0,261 mg/ml. This means that the O₂ available at the beginning is 0,261 mg/ml × 160 ml = 41,76 mg (1,305 mmol). The amount dissolved in the seawater can be neglected. This amount of O₂ is sufficient to oxidize to CO₂ and H₂O an amount of biodegradable organic carbon equal to 15,66 mg and producing 1,305 × 44 = 57,42 mg of CO₂.

The system is opened in order to refresh the headspace when the O₂ concentration reaches 25 % of the original oxygen concentration.



- Key**
- 1 sediment
 - 2 sample
 - 3 cover slip
 - 4 liquid medium
 - 5 container for the CO₂ absorber

Figure A.1 — Test flask

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