

Photography — Processing waste — Determination of ammoniacal nitrogen (microdiffusion method)

ICS 13.030.20; 37.040.30

National foreword

This British Standard reproduces verbatim ISO 6853:2001 and implements it as the UK national standard.

The UK participation in its preparation was entrusted to Technical Committee CPW/42, Photography, which has the responsibility to:

- aid enquirers to understand the text;
- present to the responsible international/European committee any enquiries on the interpretation, or proposals for change, and keep the UK interests informed;
- monitor related international and European developments and promulgate them in the UK.

A list of organizations represented on this committee can be obtained on request to its secretary.

Cross-references

The British Standards which implement international publications referred to in this document may be found in the BSI Standards Catalogue under the section entitled "International Standards Correspondence Index", or by using the "Find" facility of the BSI Standards Electronic Catalogue.

A British Standard does not purport to include all the necessary provisions of a contract. Users of British Standards are responsible for their correct application.

Compliance with a British Standard does not of itself confer immunity from legal obligations.

Summary of pages

This document comprises a front cover, an inside front cover, the ISO title page, pages ii to v, a blank page, pages 1 to 8, an inside back cover and a back cover.

The BSI copyright date displayed in this document indicates when the document was last issued.

This British Standard, having been prepared under the direction of the Consumer Products and Services Policy and Strategy Committee, was published under the authority of the Standards Policy and Strategy Committee on 18 January 2002

© BSI 18 January 2002

ISBN 0 580 37505 6

Amendments issued since publication

Amd. No.	Date	Comments

INTERNATIONAL STANDARD

**ISO
6853**

Second edition
2001-12-15

Photography — Processing waste — Determination of ammoniacal nitrogen (microdiffusion method)

*Photographie — Effluents de traitement — Détermination de l'azote
ammoniacal (méthode par microdiffusion)*



Reference number
ISO 6853:2001(E)

Contents		Page
Foreword		iv
Introduction		v
1	Scope	1
2	Normative references	1
3	Principle	1
4	Reaction	2
5	Safety and operational precautions	2
6	Materials and reagents	3
7	Apparatus	5
8	Sampling and sample preparation	5
9	Procedure	6
10	Expression of results	7
11	Test report	7
Bibliography		8

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.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 3.

Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

Attention is drawn to the possibility that some of the elements of this International Standard may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

International Standard ISO 6853 was prepared by Technical Committee ISO/TC 42, *Photography*.

This second edition cancels and replaces the first edition (ISO 6853:1987), of which it constitutes a technical revision.

Introduction

This International Standard is one of a series devoted to the analysis of photographic wastes; it encompasses the field of analysis of the ammoniacal nitrogen content in a photographic effluent.

This International Standard is intended for use by individuals with a working knowledge of analytical techniques. Some of the procedures use caustic, toxic or otherwise hazardous chemicals. Safe laboratory practice for the handling of chemicals requires the use of safety glasses or goggles and, in some cases, other protective apparel such as rubber gloves, face masks or aprons. Normal precautions for the safe performance of any chemical procedure must be exercised at all times, but specific details have been provided for hazardous materials. Hazard warnings are designated by a letter enclosed in angle brackets " $\langle \rangle$." These are defined in clause 5 and then used throughout the text. More detailed information on hazards, handling and use of these chemicals may be available from the manufacturer.

Photographic laboratories can establish conformity to effluent regulations only by chemical analysis. If this cannot be done in-house, an outside laboratory should be used.

Photography — Processing waste — Determination of ammoniacal nitrogen (microdiffusion method)

1 Scope

This International Standard specifies a method for the determination of ammonia and other volatile amines that can be liberated from photographic processing wastes by strong alkali, the results being expressed in terms of nitrogen.

The method is applicable for the determination of the ammonia concentration of typical photoprocessing wastes in the range of 10 mg/l to 200 mg/l of ammonia or 8 mg/l to 160 mg/l of nitrogen. Other volatile amines are determined as ammonia, but their concentrations in photoprocessing wastes are usually very low.

2 Normative references

The following normative documents contain provisions which, through reference in this text, constitute provisions of this International Standard. For dated references, subsequent amendments to, or revisions of, any of these publications do not apply. However, parties to agreements based on this International Standard are encouraged to investigate the possibility of applying the most recent editions of the normative documents indicated below. For undated references, the latest edition of the normative document referred to applies. Members of ISO and IEC maintain registers of currently valid International Standards.

ISO 648:1977, *Laboratory glassware — One-mark pipettes*.

ISO 5667-1:1980, *Water quality — Sampling — Part 1: Guidance on the design of sampling programmes*.

ISO 5667-2:1991, *Water quality — Sampling — Part 2: Guidance on sampling techniques*.

ISO 5667-3:1994, *Water quality — Sampling — Part 3: Guidance on the preservation and handling of samples*.

ISO 6353-1:1982, *Reagents for chemical analysis — Part 1: General test methods*.

ISO 6353-2:1983, *Reagents for chemical analysis — Part 2: Specifications — First series*.

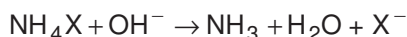
ISO 6353-3:1987, *Reagents for chemical analysis — Part 3: Specifications — Second series*.

ISO 10349-1:1992, *Photography — Photographic-grade chemicals — Test methods — Part 1: General*.

3 Principle

Ammonia is liberated from the sample by treatment with potassium metaborate and is absorbed into boric acid. The absorbed ammonia is then determined by titration with standard sulfuric acid. The liberation and absorption are carried out in a microdiffusion cell, which is a small covered dish with concentric chambers for sample, sealant and absorbing solution. When the sample and the metaborate are mixed in the sample chamber, ammonia is evolved and absorbed into the boric acid chamber. The gaseous free path from sample to absorbent is short, in order to ensure a room temperature distillation in a reasonable time. Scale-up of this method or alteration of the sample to buffer ratio should be avoided, as these modifications change the rate of ammonia diffusion.

4 Reaction



5 Safety and operational precautions

5.1 Hazard warnings

Some of the chemicals specified in the test procedures are caustic, toxic or otherwise hazardous. Safe laboratory practice for the handling of chemicals requires the use of safety glasses or goggles, and in some cases other protective apparel such as rubber gloves, face masks and aprons. Specific danger notices are given in the text for particularly dangerous materials, but normal precautions are required during the performance of any chemical procedure at all times.

The first time that a hazardous material is noted in the procedures, the hazard will be indicated by the word "**DANGER**" followed by a symbol consisting of angle brackets "< >" containing a letter that designates the specific hazard. A double bracket "<< >>" will be used for particularly perilous situations. In subsequent statements involving handling of these hazardous materials, only the hazard symbol consisting of the brackets and letter(s) will be displayed. Furthermore, for a given material, the hazard symbols will be used only once in a single paragraph.

Hazard warning symbols will not be used for common organic solvents when used in quantities of less than 1 litre, unless they are particularly hazardous.

Detailed warnings for handling chemicals and their diluted solutions are beyond the scope of this International Standard.

Employers shall provide training and health and safety information in conformance with legal requirements.

The hazard code system used in this International Standard is intended to provide information to the users and is not meant for compliance with any legal requirements for labelling, as these vary from country to country.

It is strongly recommended that anyone using these chemicals obtain pertinent information from the manufacturer about the hazards, handling, use and disposal of these chemicals.

5.2 Hazard information code system

- Harmful if inhaled. Avoid breathing dust, vapour, mist or gas. Use only with adequate ventilation.
- <C> Harmful if contact occurs. Avoid contact with eyes, skin or clothing. Wash thoroughly after handling.
- <F> Will burn. Keep away from heat, sparks and open flame. Use with adequate ventilation.
- <O> Oxidizer. Contact with other material may cause fire. Do not store near combustible materials.
- <S> Harmful if swallowed. Wash thoroughly after handling. If swallowed, obtain medical attention immediately.
- <<S>> May be fatal if swallowed. If swallowed, obtain medical attention immediately.

5.3 Safety precautions

ALL PIPETTE OPERATIONS SHALL BE PERFORMED WITH A PIPETTE BULB OR PLUNGER PIPETTE. THIS IS A CRITICAL SAFETY WARNING!

Safety glasses shall be worn for all laboratory work.

6 Materials and reagents

6.1 General

6.1.1 Handling and labelling

Reagents shall be handled in conformity with health and safety precautions as shown on containers, or as given in other sources of such information. Proper labelling of prepared reagents includes the chemical name, date of preparation, expiration date, restandardization date, name of preparer, and adequate health and safety precautions. The discharge of reagents shall conform to applicable environmental regulations.

6.1.2 Purity

Reagents used in the test procedures shall be certified reagent-grade chemicals and shall meet appropriate standards, or be chemicals of a purity acceptable for the analysis. For details, see ISO 6353-1, ISO 6353-2 and ISO 6353-3.

6.1.3 Water

Whenever water is specified without other qualifiers in the test procedures, only distilled water or water of equivalent purity shall be used.

6.1.4 Strength of solutions

6.1.4.1 Acids and ammonium hydroxide are full strength unless otherwise specified.

6.1.4.2 When a standardized solution is required, its amount-of-substance-concentration is expressed in moles per litre. The number of significant figures to which the molarity is known shall be sufficient to ensure that the reagent does not limit the reliability of the test method.

6.1.4.3 When a standardized solution is not required, its concentration is expressed in grams per litre (g/l) to the appropriate number of significant figures.

6.1.4.4 When a solution is to be diluted, its dilution is indicated by $(X + Y)$, meaning that X volumes of reagent, or concentrated solution, are to be diluted with Y volumes of water (6.1.3).

6.2 Reagents

6.2.1 Potassium tetraborate solution, 514 g/l.

Weigh $673 \text{ g} \pm 0,1 \text{ g}$ of potassium tetraborate, $\text{K}_2\text{B}_4\text{O}_7 \cdot 4\text{H}_2\text{O}$, and dissolve it in 550 ml of water in a 1 litre beaker. Then weigh $247 \text{ g} \pm 0,1 \text{ g}$ of potassium hydroxide, KOH (DANGER: <<C>>), and dissolve it in the tetraborate solution. Boil on a hotplate for 5 min, cool and add 5 ml of a 10 % aqueous solution of nonylphenoxypoly (6–10) ethylene oxide, NPPO, or similar wetting agent¹⁾. Transfer to a 1 litre volumetric flask, rinsing the beaker into the flask several times. When cool, dilute to volume and mix well. Note that the wetting agent will separate out on standing; therefore, the flask must be shaken vigorously before each use.

6.2.2 Boric acid absorbent solution

Add about 800 ml of water to a 1 litre volumetric flask. Stir, using a magnetic stirrer, and add 2 mg to 3 mg of xylene cyanole FF, weighed to the nearest 1 mg, followed by 0,5 ml of NPPO, followed by 5,0 ml of methyl red indicator solution prepared by dissolving 0,125 g of methyl red in 250 ml of methanol (DANGER: <F><S>). Add $6 \text{ g} \pm 0,1 \text{ g}$ of boric acid, H_3BO_3 , keeping the contents of the flask stirred until all the constituents are dissolved.

1) Non-ionic detergent with a hydrophilic lipophilic balance in the range of 13 to 14.

ISO 6853:2001(E)

Dilute to within about 15 ml of the mark and mix. Place 1,5 ml of this solution in the centre of a microdiffusion cell and observe the colour.

If the colour in the cell is pink, add just sufficient 0,1 mol/l sodium hydroxide (6.2.7) to the solution in the 1 litre flask to obtain a neutral colour when 1 ml is viewed in the microdiffusion cell.

Check that excess sodium hydroxide is absent by adding 0,10 ml of 0,002 50 mol/l sulfuric acid (6.2.5) to 1 ml of the solution, at which point a pink colour should be produced. Note that the solution in the 1 litre flask will appear red, even when the 1 ml in the microdiffusion cell looks neutral.

6.2.3 Sulfuric acid solution, $c(\text{H}_2\text{SO}_4) \approx 0,5 \text{ mol/l}$.

Carefully add 28,3 ml of sulfuric acid (DANGER: <<C>>), $\rho \approx 1,84 \text{ g/ml}$, to 500 ml of water in a 1 litre volumetric flask while mixing and cooling. Dilute to the mark with water.

6.2.4 Sulfuric acid solution, $c(\text{H}_2\text{SO}_4) \approx 0,05 \text{ mol/l}$.

Pipette 100,0 ml of 0,5 mol/l sulfuric acid (6.2.3) into a 1 litre volumetric flask and dilute to the mark with water.

NOTE This solution is not the standard volumetric solution that is required in 6.2.5.

6.2.5 Sulfuric acid solution, $c(\text{H}_2\text{SO}_4) = 0,002 50 \text{ mol/l}$.

Pipette 50,0 ml of standard volumetric 0,050 00 mol/l sulfuric acid into a 1 litre volumetric flask and dilute to the mark with water.

The solution prepared in 6.2.4 is not a standard volumetric solution. A commercial volumetric standard solution is recommended. If a volumetric standard solution needs to be prepared, see an analytical chemistry text.

6.2.6 Sodium hydroxide solution, $c(\text{NaOH}) \approx 1 \text{ mol/l}$.

Dissolve $40 \text{ g} \pm 0,5 \text{ g}$ of sodium hydroxide (DANGER: <<C>>) in 500 ml of water in a 1 litre volumetric flask. Mix, cool, and dilute to the mark with water.

6.2.7 Sodium hydroxide solution, $c(\text{NaOH}) \approx 0,1 \text{ mol/l}$.

Dissolve $4,0 \text{ g} \pm 0,1 \text{ g}$ of sodium hydroxide (<<<C>>) in 500 ml of water in a 1 litre volumetric flask. Mix, cool, and dilute to the mark with water.

6.2.8 Cleaning solutions for microdiffusion cells

6.2.8.1 Cleaning solution A

From a graduated measuring cylinder, add about 750 ml of water and about 750 ml of 0,5 mol/l sulfuric acid (6.2.3) to a 2 litre beaker and stir. Continue stirring, and add 2 ml to 3 ml of household liquid dishwashing detergent and 1 ml of methyl red indicator solution.

NOTE Avoid the use of detergents containing ammonia.

6.2.8.2 Cleaning solution B

Add about 1,5 litres of water and 2 ml to 3 ml of the detergent to a 2 litre beaker and mix. Add 10 ml of 1 mol/l sodium hydroxide (6.2.6), and then add enough methyl red indicator solution, while stirring, to produce a yellow colour.

6.2.8.3 Cleaning solution C

Add about 1,5 litres of water and about 5 ml of the detergent to a 2 litre beaker and stir. Continue to stir, and add about 10 ml of 0,05 mol/l sulfuric acid (6.2.4) and enough methyl red indicator to produce a pink colour.

6.2.9 Standard nitrogen samples, to check the ammonia liberation technique.

6.2.9.1 Nitrogen stock solution, $\rho_{\text{N}} = 1\,000\text{ mg/l}$ ($c_{\text{N}} = 0,071\,4\text{ mol/l}$).

Dry ammonium chloride for 2 h in an oven at 100 °C and allow it to cool in a desiccator before weighing. Weigh $3,819\text{ g} \pm 0,001\text{ g}$ of this ammonium chloride and quantitatively transfer it to a 1 litre volumetric flask. Dissolve it in water and dilute to volume. This stock solution is stable for at least 3 months.

6.2.9.2 Standard nitrogen solution, $\rho_{\text{N}} = 100\text{ mg/l}$ ($c_{\text{N}} = 0,007\,14\text{ mol/l}$).

Pipette 10 ml of the nitrogen stock solution (6.2.9.1) into a 100 ml volumetric flask and dilute to the mark with water.

6.2.9.3 Standard nitrogen solution, $\rho_{\text{N}} = 20\text{ mg/l}$ ($c_{\text{N}} = 0,001\,428\text{ mol/l}$).

Pipette 2 ml of the nitrogen stock solution (6.2.9.1) into a 100 ml volumetric flask and dilute to the mark with water.

7 Apparatus

7.1 General

All glassware subject to heating shall be of heat-resistant borosilicate glass.²⁾

Pipettes and other volumetric glassware shall meet the requirements specified in ISO 10349-1.

7.2 Microdiffusion cell, 83 mm, Obrick microdiffusion.

7.3 Micropipette, of capacity 0,500 ml and conforming to ISO 648.

7.4 Syringe pipette, of capacity 1,00 ml.

7.5 Syringe microburette.

7.6 Syringe, capable of delivering 1,50 ml.

7.7 Polytetrafluoroethylene-coated magnetic bar, 7 mm × 2 mm.

8 Sampling and sample preparation

It is necessary that the analysis be carried out on a representative sample. The sampling of a process effluent or a plant effluent can encompass many difficulties and due care shall be exercised. See ISO 5667-1, ISO 5667-2 and ISO 5667-3.

Sampling shall be carried out in conformance with regulatory requirements. Sampling should be done under typical operating conditions and normally should be representative of the overall plant effluent. Daily samples that are truly representative of the effluents require sampling over 24 h and sampling that is proportional to the flow rate.

Samples taken during a sudden discharge or during another non-routine operation will not yield results representative of the normal operation.

2) Pyrex® is an example of suitable glassware available commercially. This information is given for the convenience of the users of this International Standard and does not constitute an endorsement by ISO of this product.

9 Procedure

9.1 Cleaning microdiffusion cells and covers

Soak the cells and covers in cleaning solution A (6.2.8.1) for 1 h; remove and soak in cleaning solution B (6.2.8.2) for 1 h; then remove and soak in cleaning solution C (6.2.8.3) for 1 h.

Remove the cells and covers, shake them as dry as possible and invert them to dry on a clean towel. Do not touch the insides of the clean cells and covers.

If the cleaned cells and covers have not been washed on the day that they are to be used, rinse them with water and place them on a clean towel to dry before using.

9.2 Liberation of ammonia from samples

Using the syringe (7.6), add 1,50 ml of the boric acid absorbent solution (6.2.2) to the centre of the microdiffusion cell (7.2). Using the 1,00 ml syringe pipette (7.4), add 1,00 ml of the potassium tetraborate solution (6.2.1) to the outer sealing chamber and 1,00 ml to the sample chamber, i.e. the second chamber from the outside.

The tetraborate solution should be vigorously shaken before sampling in order to distribute the NPPO. Great care shall be taken not to splash any tetraborate solution into the centre chamber. If this occurs, a green coloration will be produced and the sample shall be discarded. The tetraborate solution should be deposited on only one part of the sample chamber. Leave enough space for the sample to be added without mixing until the cell is sealed.

Using the calibrated 0,500 ml micropipette (7.3), carefully add 0,500 ml of the sample to the empty side of the sample chamber. Cover the cell immediately and rotate the cover to spread the tetraborate solution and form a good seal. Using a rotary tipping motion, mix the sample with the tetraborate solution for 30 s, making certain that the sample and absorbent chambers are fully covered and that the contents of the separate chambers do not mix. Let the cell stand for 2 h. Longer standing times will not introduce any error.

9.3 Titration

Remove the lid carefully from the cell and place the cell on a magnetic stirrer. Place the magnetic bar (7.7) in the centre chamber without splashing. Using the syringe microburette (7.5), titrate the solution in the centre chamber with 0,002 50 mol/l sulfuric acid (6.2.5) to the first pink colour. Keep the burette tip immersed while titrating. Stir for about 15 s. If the colour fades to light green, continue to titrate until the pink colour remains for at least 15 s. Identify the volume used for titration as V_1 .

9.4 Blank reading and a calibration curve check

Concurrently with the sample to be analysed, run procedures 9.1, 9.2 and 9.3 on 0,500 ml of water, on 0,500 ml of the 100 mg/l standard nitrogen solution (6.2.9.2), and on the 20 mg/l standard nitrogen solution (6.2.9.3). Identify the volume used for the titration on the blank as V_2 ; on the 100 mg/l standard nitrogen solution as V_3 ; and on the 20 g/l standard nitrogen solution as V_4 .

10 Expression of results

10.1 Method of calculation

The total nitrogen concentration ρ_{N} , in milligrams per litre, is given by the formula

$$\rho_{\text{N}} = \frac{72,6(V_1 - V_2)}{V_{\text{s}}}$$

where

V_1 is the volume, in millilitres, used for the titration of the sample;

V_2 is the volume, in millilitres, used for the titration of the water blank;

V_{s} is the volume, in millilitres, of the sample taken for diffusion.

If V_{s} is 0,500 ml, the formula becomes

$$\rho_{\text{N}} = 145(V_1 - V_2)$$

NOTE The above constants (72,6 and 145) are experimental values and slightly less than theoretical. They were obtained by a linear regression analysis of a calibration curve and take into account a small loss of amino nitrogen in the diffusion step. Laboratories may wish to generate their own constants for more accurate results.

As a check on the precision, perform similar calculations on the results ($V_3 - V_2$) and ($V_4 - V_2$) for the two standard nitrogen samples. These should be carried out for each set of effluent samples or at least daily. The agreement of these results indicates the probable reliability of the results on unknown samples.

10.2 Precision

The 95 % confidence limits for a single determination are expected to be between 4 mg and 8 mg of nitrogen per litre. The stoichiometric yield by the microdiffusion method is about 96 %.

11 Test report

The test report shall include the following information:

- a reference to this International Standard, i.e. the reference of the method used;
- an identification of the sample;
- the results and the method of expression used;
- any unusual features noted during the determination;
- any operation not included in this International Standard, or regarded as optional.

Bibliography

- [1] ISO 385-1:1984, *Laboratory glassware — Burettes — Part 1: General requirements.*
- [2] ISO 385-2:1984, *Laboratory glassware — Burettes — Part 2: Burettes for which no waiting time is specified.*
- [3] ISO 385-3:1984, *Laboratory glassware — Burettes — Part 3: Burettes for which a waiting time of 30 s is specified.*
- [4] ISO 835-1:1981, *Laboratory glassware — Graduated pipettes — Part 1: General requirements.*
- [5] ISO 835-2:1981, *Laboratory glassware — Graduated pipettes — Part 2: Pipettes for which no waiting time is specified.*
- [6] ISO 835-3:1981, *Laboratory glassware — Graduated pipettes — Part 3: Pipettes for which a waiting time of 15 s is specified.*
- [7] ISO 835-4:1981, *Laboratory glassware — Graduated pipettes — Part 4: Blow-out pipettes.*
- [8] ISO 1042:1998, *Laboratory glassware — One-mark volumetric flasks.*
- [9] ISO 4788:1980, *Laboratory glassware — Graduated measuring cylinders.*
- [10] ISO 5725-1:1994, *Accuracy (trueness and precision) of measurement methods and results — Part 1: General principles and definitions.*
- [11] ISO 6851:2001, *Photography — Processing waste — Determination of total amino nitrogen (microdiffusion Kjeldahl method).*

BSI — British Standards Institution

BSI is the independent national body responsible for preparing British Standards. It presents the UK view on standards in Europe and at the international level. It is incorporated by Royal Charter.

Revisions

British Standards are updated by amendment or revision. Users of British Standards should make sure that they possess the latest amendments or editions.

It is the constant aim of BSI to improve the quality of our products and services. We would be grateful if anyone finding an inaccuracy or ambiguity while using this British Standard would inform the Secretary of the technical committee responsible, the identity of which can be found on the inside front cover.
Tel: +44 (0)20 8996 9000. Fax: +44 (0)20 8996 7400.

BSI offers members an individual updating service called PLUS which ensures that subscribers automatically receive the latest editions of standards.

Buying standards

Orders for all BSI, international and foreign standards publications should be addressed to Customer Services. Tel: +44 (0)20 8996 9001.
Fax: +44 (0)20 8996 7001. Email: orders@bsi-global.com. Standards are also available from the BSI website at <http://www.bsi-global.com>.

In response to orders for international standards, it is BSI policy to supply the BSI implementation of those that have been published as British Standards, unless otherwise requested.

Information on standards

BSI provides a wide range of information on national, European and international standards through its Library and its Technical Help to Exporters Service. Various BSI electronic information services are also available which give details on all its products and services. Contact the Information Centre.
Tel: +44 (0)20 8996 7111. Fax: +44 (0)20 8996 7048. Email: info@bsi-global.com.

Subscribing members of BSI are kept up to date with standards developments and receive substantial discounts on the purchase price of standards. For details of these and other benefits contact Membership Administration.
Tel: +44 (0)20 8996 7002. Fax: +44 (0)20 8996 7001.
Email: membership@bsi-global.com.

Information regarding online access to British Standards via British Standards Online can be found at <http://www.bsi-global.com/bsonline>.

Further information about BSI is available on the BSI website at <http://www.bsi-global.com>.

Copyright

Copyright subsists in all BSI publications. BSI also holds the copyright, in the UK, of the publications of the international standardization bodies. Except as permitted under the Copyright, Designs and Patents Act 1988 no extract may be reproduced, stored in a retrieval system or transmitted in any form or by any means – electronic, photocopying, recording or otherwise – without prior written permission from BSI.

This does not preclude the free use, in the course of implementing the standard, of necessary details such as symbols, and size, type or grade designations. If these details are to be used for any other purpose than implementation then the prior written permission of BSI must be obtained.

Details and advice can be obtained from the Copyright & Licensing Manager.
Tel: +44 (0)20 8996 7070. Fax: +44 (0)20 8996 7553.
Email: copyright@bsi-global.com.