Draft for Public Comment

Form 36

DPC: 24/30463968 DC

BSI Group Headquarters

389 Chiswick High Road London W4 4AL

Tel: +44 (0)20 8996 9000
Fax: +44 (0)20 8996 7400
www.bsigroup.com

Date: 10 October 2024
Origin: International

Latest date for receipt of comments: 03 December 2024

Project No. 2022/02324

Responsible committee: CH/304 Healthcare Organization Management

Interested committees:

Title: Draft BS ISO 18706 Healthcare organization management. Pandemic response (respiratory). Functions and quality evaluation of test booth for specimen collection.

Please notify admin.start@bsigroup.com if you are aware of any keywords that might assist in clarifying or identifying the standard or of the content of this standard:

i) has any issues related to 3rd party IPR, patent or copyright
ii) affects other national standard(s)
iii) requires additional national guidance or information

WARNING: THIS IS A DRAFT AND MUST NOT BE REGARDED OR USED AS A BRITISH STANDARD. THIS DRAFT IS NOT CURRENT BEYOND 03 December 2024

This draft is issued to allow comments from interested parties; all comments will be given consideration prior to publication. No acknowledgement will normally be sent. **See overleaf for information on the submission of comments.**

No copying is allowed, in any form, without prior written permission from BSI except as permitted under the Copyright, Designs and Patent Act 1988 or for circulation within a nominating organization for briefing purposes. Electronic circulation is limited to dissemination by e-mail within such an organization by committee members.

Further copies of this draft may be purchased from BSI Shop http://shop.bsigroup.com or from BSI Customer Services, Tel: +44(0) 20 8996 9001 or email cservices@bsigroup.com. British, International and foreign standards are also available from BSI Customer Services.

Information on the co-operating organizations represented on the committees referenced above may be obtained from https://standardsdevelopment.bsigroup.com/.

Documents
Document Sharing Hub

Full document details: 24/30463968 DC

Introduction

This draft standard is based on national and international discussions. Your comments on this draft are invited and will assist in the preparation of the consequent standard.

For international standards, comments will be reviewed by the relevant UK national committee before sending the consensus UK vote and comments to the international committee, which will then decide appropriate action. If the international standard is approved, it is usual for the text to be published as a British Standard.

For national standards, comments will be reviewed by the relevant UK national committee and the resulting standards published as a British Standard.

UK Vote

Please indicate whether you consider the UK should submit a negative (with supporting technical reasons) or positive vote on this draft. Please indicate if you are aware of any reason why this draft standard should not be published as a British Standard.

Submission of Comments

- **Annotated drafts are not acceptable and will be rejected.**
- All comments should be submitted online at **https://standardsdevelopment.bsigroup.com/**. You will need to register in order to comment.

Template for comments and secretariat observations

Date: xx/xx/20xx	Document: ISO/DIS xxxx

1	2	(3)	4	5	(6)	7
MB	Clause No./ Subclause No./Annex **(e.g. 3.1)**	Paragraph/ Figure/Table/Note	**Type of comment**	**Commend (justification for change) by the MB**	**Proposed change by the MB**	**Secretariat observations on each comment submitted**
	3.1	Definition 1	ed	Definition is ambiguous and needs clarifying.	Amend to read '...so that the mains connector to which no connection...'	
	6.4	Paragraph 2	te	The use of the UV photometer as an alternative cannot be supported as serious problems have been encountered in its use in the UK.	Delete reference to UV photometer.	

Documents
Document Sharing Hub

Full document details: 24/30463968 DC

DRAFT International Standard

ISO/DIS 18706

Healthcare organization management — Pandemic response (respiratory) — Functions and quality evaluation of test booth for specimen collection

ICS: 11.020.01

ISO/TC **304**

Secretariat: **ANSI**

Voting begins on:
2024-10-10

Voting terminates on:
2025-01-02

THIS DOCUMENT IS A DRAFT CIRCULATED FOR COMMENTS AND APPROVAL. IT IS THEREFORE SUBJECT TO CHANGE AND MAY NOT BE REFERRED TO AS AN INTERNATIONAL STANDARD UNTIL PUBLISHED AS SUCH.

IN ADDITION TO THEIR EVALUATION AS BEING ACCEPTABLE FOR INDUSTRIAL, TECHNOLOGICAL, COMMERCIAL AND USER PURPOSES, DRAFT INTERNATIONAL STANDARDS MAY ON OCCASION HAVE TO BE CONSIDERED IN THE LIGHT OF THEIR POTENTIAL TO BECOME STANDARDS TO WHICH REFERENCE MAY BE MADE IN NATIONAL REGULATIONS.

RECIPIENTS OF THIS DRAFT ARE INVITED TO SUBMIT, WITH THEIR COMMENTS, NOTIFICATION OF ANY RELEVANT PATENT RIGHTS OF WHICH THEY ARE AWARE AND TO PROVIDE SUPPORTING DOCUMENTATION.

This document is circulated as received from the committee secretariat.

Reference number
ISO/DIS 18706:2024(en)

© ISO 2024

AVA Documents
Document Sharing Hub

Full document details: 24/30463968 DC

COPYRIGHT PROTECTED DOCUMENT

© ISO 2024

All rights reserved. Unless otherwise specified, or required in the context of its implementation, no part of this publication may be reproduced or utilized otherwise in any form or by any means, electronic or mechanical, including photocopying, or posting on the internet or an intranet, without prior written permission. Permission can be requested from either ISO at the address below or ISO's member body in the country of the requester.

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

Published in Switzerland

Documents
Document Sharing Hub
Full document details: 24/30463968 DC

Contents

Page

Foreword ... **iv**
Introduction ... **v**
1 Scope ... **1**
2 Normative references ... **1**
3 Terms and definitions ... **1**
4 Negative pressure booth ... **3**
4.1 Principle ... 3
4.2 Structure ... 3
4.2.1 General ... 3
4.2.2 Main body ... 3
4.2.3 Filtration ... 4
4.2.4 Glove and work shelf ... 4
4.2.5 Control unit ... 5
4.2.6 Option for outdoor usage ... 5
4.3 Function ... 5
4.3.1 General ... 5
4.3.2 HEPA filter leak ... 5
4.3.3 Noise level ... 5
4.3.4 Light intensity ... 5
4.3.5 Differential pressure ... 5
4.3.6 Air velocity ... 6
5 Positive pressure booth ... **6**
5.1 Principle ... 6
5.2 Structure ... 6
5.2.1 General ... 6
5.2.2 Main body ... 7
5.2.3 Filtration ... 7
5.2.4 Glove and work shelf ... 8
5.2.5 Control unit ... 8
5.2.6 Option for outdoor usage ... 8
5.3 Function ... 8
5.3.1 General ... 8
5.3.2 HEPA filter leak ... 8
5.3.3 Noise level ... 8
5.3.4 Lighting intensity ... 9
5.3.5 Differential pressure ... 9
5.3.6 Air velocity ... 9
Annex A (normative) **Performance Evaluation** ... **10**
Annex B (informative) **Size information considering ergonomic factors** ... **16**
Bibliography ... **17**

© ISO 2024 – All rights reserved

Documents
Document Sharing Hub
Full document details: 24/30463968 DC

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).

ISO draws attention to the possibility that the implementation of this document may involve the use of (a) patent(s). ISO takes no position concerning the evidence, validity or applicability of any claimed patent rights in respect thereof. As of the date of publication of this document, ISO had not received notice of (a) patent(s) which may be required to implement this document. However, implementers are cautioned that this may not represent the latest information, which may be obtained from the patent database available at www.iso.org/patents. ISO shall not be held responsible for identifying any or all such patent rights.

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.

This document was prepared by Technical Committee ISO/TC 304, Healthcare organization management.

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.

© ISO 2024 – All rights reserved

Documents
Document Sharing Hub
Full document details: 24/30463968 DC

Introduction

Globally, as the movement of population have been active, infectious diseases that have occurred locally have spread throughout the world and are now becoming a pandemic.

Over the past 20 years, infectious diseases such as Severe Acute Respiratory Syndrome (SARS, in 2002), H1N1 Influenza (H1N1, in 2009), and Middle East Respiratory Syndrome (MERS, in 2012) appear to be surfacing with increased intensity and shorter cycle times. Experts are concerned that occurrence of disease continues to increase and the cycle tends to be shorter.

When an infectious disease spreads on a large scale such as pandemic, if screening treatment is carried out inside a hospital, hospital workers, treatment, inpatients, and patient caregivers are exposed to the risk of infection. To reduce the risk of such cross-infection, sample collection methods such as drive-through screening station (DTSS) and walk-through screening station (WTSS) are being developed.

The specimen collection booth is a facility that physically separates medical staff and walking visitors in WTSS to reduce the risk of cross-infection and enable quick testing.

This standard considered structural elements, functions, and evaluation methods to be equipped for the safe use of a specimen collection booth to quickly collect samples from suspected respiratory diseases.

© ISO 2024 – All rights reserved

Documents
Document Sharing Hub
Full document details: 24/30463968 DC

Documents
Document Sharing Hub