

SLOVENSKI STANDARD oSIST prEN ISO 23402-1:2020

01-marec-2020

Zobozdravstvo - Prenosna dentalna oprema za zdravstveno oskrbo - 1. del: Splošne zahteve (ISO/DIS 23402-1:2020)

Dentistry - Portable dental equipment for use in non permanent healthcare environment - Part 1: General requirements (ISO/DIS 23402-1:2020)

Zahnheilkunde - Bewegliche dentale Ausrüstung zur Anwendung in nicht-permanenten Gesundheitseinrichtungen - Teil 1: Allgemeine Anforderungen (ISO/DIS 23402-1:2020)

Médecine bucco-dentaire - Matériel dentaire portatif utilisable dans des environnements de soins de santé non permanents - Partie 1: Exigences générales (ISO/DIS 23402-1:2020)

Ta slovenski standard je istoveten z: prEN ISO 23402-1

ICS:

11.060.20 Zobotehnična oprema Dental equipment

oSIST prEN ISO 23402-1:2020 en

oSIST prEN ISO 23402-1:2020

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 23402-1:2020

https://standards.iteh.ai/catalog/standards/sist/1b12ecab-1a4a-49d9-95aa-0f2409390467/sist-en-iso-23402-1-2020

DRAFT INTERNATIONAL STANDARD ISO/DIS 23402-1

ISO/TC **106**/SC **6**

Secretariat: DIN

Voting begins on: **2020-01-16**

Voting terminates on:

2020-04-09

Dentistry — Portable dental equipment for use in nonpermanent healthcare environment —

Part 1:

General requirements

ICS: 11.060.20

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 23402-1:2020 https://standards.iteh.ai/catalog/standards/sist/1b12ecab-1a4a-49d9-95aa

THIS DOCUMENT IS A DRAFT CIRCULATED FOR COMMENT 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.

ISO/CEN PARALLEL PROCESSING



Reference number ISO/DIS 23402-1:2020(E)

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 23402-1:2020 https://standards.iteh.ai/catalog/standards/sist/1b12ecab-1a4a-49d9-95aa-0f2409390467/sist-en-iso-23402-1-2020



COPYRIGHT PROTECTED DOCUMENT

© ISO 2020

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 Fax: +41 22 749 09 47 Email: copyright@iso.org Website: www.iso.org Published in Switzerland

ii

Contents			Page
Fore	eword		iv
Intr	oductio	on	v
1		e	
2	-	native references	
3		ns and definitions	
4	Classification		
	4.1	General	
	4.2	For electrically operated devices	
	4.3	According to intended use environment	
	4.4	According to supply sources	
	4.5	According to transport conditions	
	4.6	According to human portability	
5	Requirements		
	5.1	General	
	5.2	Transport requirements	
		5.2.1 General	
		5.2.2 Maximum mass	
		5.2.3 Maximum dimensions	4
		5.2.4 Environmental exposure	4
		5.2.5 Impact	4
		5.2.7 Vibration	
		5.2.8 Particulate and liquid ingress during transport	
		5.2.9 Flammability	
	5.3		5 5
	5.4 h	Assembly and disassembly requirements Utility requirements	5
	5.5	Operational requirements 7/sist-en-iso-23402-1-2020	5
		5.5.1 Ambient operating conditions	
		5.5.2 Usability	
		5.5.3 Applied parts not intended to supply heat to a patient	6
		5.5.4 Cleaning and disinfection	6
		5.5.5 Noise	6
6	Sam	pling	6
7	Mea	surement and test methods	6
,	7.1	Tests for transport requirements	
	7.2	Environmental exposure	
0		ufacturer's instructions	
8	8.1	General	
	8.2	Instructions for use	
	8.3	Technical description	
	•		
9	Marking		
	9.1	Marking on the equipment	
	9.2	Marking of packaging	
10	Pack	aging	8
Ann	ex A (in	formative) Applicable testing methodology from IEC 60601-1:2005	9

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.

This document was prepared by Technical Committee ISO/TC 106, *Dentistry*, Subcommittee SC 6, *Dental equipment*.

ISO 23402-1 consists of the following parts, under the general title *Dentistry — Portable dental* equipment for use in non-permanent healthcare environments: 23402-1-2020

— Part 1: General requirements

The following parts are planned or in preparation:

- Part 2: Portable dental units
- Part 3: Portable suction equipment
- Part 4: Portable patient chair

A list of all parts in the ISO 23402 series can be found on the ISO website.

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

Transportable dental equipment is used by dental professionals to provide care to patients in a variety of settings. Because the intended use applications and intended means for transporting such equipment vary considerably, a wide variety of transportable dental equipment is commercially available. For example, certain transportable equipment is designed and constructed to be carried or rolled on its own wheels between rooms within a healthcare facility, while other transportable dental equipment is made to be folded and packed to carry over terrain which may be rugged and used in transient dental care settings which may have only limited shelter and utility services.

Transportable equipment that may be moved from one location to another while being carried by one or more persons is referred to as portable equipment. The term portable equipment applies to equipment that may be carried from room to room in a given facility or to remote parts of the world. This International Standard focuses on portable dental equipment which is specifically designed and constructed to be transported between non-clinical environments and used by dental professionals to provide dental care in such settings, including temporary field clinics.

Such portable dental equipment for use in non-permanent healthcare environments enables dental professionals to provide a high standard of care to patients who do not have access to or are not able to travel to traditional health care facilities. Settings in which this equipment is commonly used include military field environments, humanitarian aid field clinics, public health outreach clinics, patient residences, long-term care facilities, prisons, schools and workplaces.

A number of trends in health care have driven increased utilization of portable dental equipment in non-permanent healthcare environments. Military forces use portable dental equipment in support of mobilized forces or for humanitarian outreach. A variety of government and non-government organizations are increasingly providing humanitarian dental care to underserved populations and populations affected by disasters. Civilian health care workers are also increasingly providing dental services to a growing population who are simply unable to visit traditional dental clinics due to age, disability, or income. Academic and research bodies regularly conduct dental education programs, particularly at external / off-site locations (including dentistry, dental hygiene, dental assisting, etc.).

The transport and end-use conditions for portable dental equipment used in non-permanent healthcare environments drive certain unique requirements which generally do not apply to portable, mobile or stationary dental equipment used in traditional dental clinics or hospitals. Because portable equipment used in non-permanent healthcare environments is intended to be moved between venues, in some cases carried over rugged terrain or in inclement conditions, it must be designed and constructed to be safely transported by humans without damage, be efficiently assembled and disassembled, and deliver reliable service at the point of use. Special consideration must be given to the austerity of the environment in which the equipment may be used and the availability and quality of utility supplies (electrical power, water, compressed air, etc.). In order for the equipment to be sufficiently portable and capable of operating in extreme conditions, certain requirements for dental equipment intended for use in traditional clinical settings may not be practical and must be reconsidered for portable dental equipment for use in non-permanent healthcare environments. There can also be unique safety and infection control concerns to consider.

This document is one in a series of standards (planned or published), with the objective of standardizing requirements for portable dental equipment for use in non-permanent healthcare environments.

oSIST prEN ISO 23402-1:2020

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 23402-1:2020

https://standards.iteh.ai/catalog/standards/sist/1b12ecab-1a4a-49d9-95aa-0f2409390467/sist-en-iso-23402-1-2020

Dentistry — Portable dental equipment for use in nonpermanent healthcare environment —

Part 1:

General requirements

1 Scope

This document specifies general requirements and test methods for portable dental equipment for use in non-permanent healthcare environments.

Portable dental equipment within the scope of this document includes portable dental units, portable patient chairs, portable operator's stools, portable operating lights, portable suction source equipment, portable air compressors, and other portable dental equipment in instances where these devices are designed and constructed to be transported for use in non-permanent healthcare environments.

Particular requirements for specific types of portable dental equipment for use in non-permanent healthcare environments will be specified in subsequent parts of this document.

This document does not apply to stationary dental equipment, wearable equipment (such as headlamps and loupes), mobile dental equipment or portable dental equipment that is not intended to be used in non-permanent healthcare environments or not designed to be disassembled, folded or packed for human transport between non-permanent healthcare environments. Also, requirements for stationary dental equipment that may be installed in a dental mobile medical facility (e.g., vehicular or containerized mobile dental clinic) are not considered in this document.

Normative references 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 1942, Dentistry — Vocabulary

ISO 21530, Dentistry — Materials used for dental equipment surfaces — Determination of resistance to *chemical disinfectants*

ISO 4180:2009, Packaging — Complete, filled transport packages — General rules for the compilation of performance test schedules

IEC 60529:1989+AMD1:1999, Degrees of protection provided by enclosures (IP Code)

IEC 60601-1:2005+AMD1:2012, Medical electrical equipment — Part 1: General requirements for basic safety and essential performance

IEC 62366-1, Medical devices — Part 1: Application of usability engineering to medical devices

IEC 80601-2-60, Medical electrical equipment — Part 2-60: Particular requirements for the basic safety and essential performance of dental equipment

Terms and definitions 3

For the purposes of this document, the terms and definitions given in IEC 60601-1, ISO 1942 and the following apply.

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

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

3.1

dental equipment

device or combination of devices for use in providing dental treatment and/or associated procedures, including machines, apparatus, furniture and accessories

3.2

transportable dental equipment

dental equipment that is intended, once installed and placed into service, to be moved from one place to another whether or not connected to a supply and without an appreciable restriction of range

Mobile dental equipment and portable dental equipment. **EXAMPLE**

3.3

portable dental equipment

transportable dental equipment intended to be moved from one location to another while being carried by one or more persons 1611 STANDARD FREVIL

mobile dental equipment

transportable dental equipment intended to be moved from one location to another while supported by its own wheels or equivalent means

3.5

$\textbf{non-permanent healthcare environment} \ ^{0.467/sist-en-iso-23402-1-2020}$

temporary setting for providing dental or medical treatment

3.6

transport case

packaging container intended to be used for transport and for protection of the portable dental equipment

Classification

4.1 General

The following classifications pertain to electrical characteristics, end-use facility provisions, supply provisions and portability considerations applicable to portable dental equipment for use in nonpermanent healthcare environments.

4.2 For electrically operated devices

If the equipment is electrically operated, classification according to IEC 60601-1 and 80601-2-60 applies.