## **BUREAU OF INDIAN STANDARDS**

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# भारतीय मानक मसौदा दंत चिकित्सा - मल्टिफंक्शन हैंडपीस

# Draft Indian Standard Dentistry — Multifunction Handpieces

#### ICS 11.060.25

Dentistry Sectional Committee, MHD 08

Last date for comments: 05 April 2024

#### NATIONAL FOREWORD

(Adoption clause will be added later)

The text of ISO Standard has been approved as suitable for publication as an Indian Standard without deviations. Certain conventions are however not identical to those used in Indian Standards. Attention is particularly drawn to the following:

- a) Wherever the words 'International Standard' appear referring to this standard, they should be read as 'Indian Standard'
- b) Comma (,) has been used as a decimal marker while in Indian Standards, the current practice is to use a point (.) as the decimal marker.

In this adopted standard, reference appears to certain International Standards for which Indian Standards also exist. The corresponding Indian Standards which are to be substituted in their respective places are listed below along with their degree of equivalence for the editions indicated:

International Standard	Corresponding Indian Standard	Degree of Equivalence
ISO 1942, Dentistry — Vocabulary	IS 17895: 2023/ISO 1942: 2020, Dentistry Vocabulary	Identical
Stationary dental units and	IS 6116 (Part 1) : 2021/ISO 7494-1:2018, Dentistry Stationary dental units and dental patient chairs Part 1: General requirements ( <i>Second Revision</i> )	Identical

ISO 10993-1, Biological evaluation of medical devices — Part 1: Evaluation and testing within a risk management process	IS 17932 (Part 1) : 2023, Biological evaluation of medical devices Part 1: Evaluation and testing within a risk management process (ISO 10993-1 : 2018, MOD)	Modified/Technically Equivalent
ISO 15223-1, Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied — Part 1: General requirements	IS 18105 (Part 1) : 2023/ ISO 15223-1:2021, Medical devices Symbols to be used with information to be supplied by the manufacturer Part 1: General requirements ( <i>Third Revision</i> )	Identical
ISO 17664-1, Processing of health care products — Information to be provided by the medical device manufacturer for the processing of medical devices	MHD/12/20835, Processing of health care products — Information to be provided by the medical device manufacturer for the processing of medical devices Part 1: Critical and semi critical medical devices	Identical
IEC 60601-1, Medical electrical equipment — Part 1: General requirements for basic safety and essential performance	MHD/15/22648, Medical electrical equipment : Part 1 General requirements for basic safety and essential performance ( <i>Third Revision</i> )	Modified/Technically Equivalent
IEC 62366-1, Medical devices — Part 1: Application of usability engineering to medical devices	IS 17922 (Part 1) : 2023/ IEC 62366-1: 2015 + AMD 1 : 2020, Medical Devices Part 1: Application of Usability Engineering ( <i>First Revision</i> )	Identical

The technical committee has reviewed the provisions of the following International Standards referred in this adopted standard and has decided that they are acceptable for use in conjunction with this standard:

International Standard ISO 9687	<i>Title</i> Dentistry — Graphical symbols for dental equipment	
ISO 21530	Dentistry — Materials used for dental equipment surfaces — Determination of resistance to chemical disinfectants	
ISO 21531	Dentistry — Graphical symbols for dental instruments	
IEC 80601-2-60	Medical electrical equipment — Part 2-60: Particular requirements for the basic safety and essential performance of dental equipment	

This standard alters the reference from ISO 17664 to ISO 17664-1, wherever ISO 17664 appears in this standard it shall be read as ISO 17664-1.

For the purpose of deciding whether a particular requirement of this standard is complied with, the final value, observed or calculated, expressing the result of a test shall be rounded off in accordance with IS 2: 2022 'Rules for rounding off numerical values (*second revision*)'.

This standard also makes a reference to the BIS Certification Marking of the product. Details of which is given in National Annex A.

#### NATIONAL ANNEX A

(National Foreword)

### A-1 BIS CERTIFICATION MARKING

The product(s) conforming to the requirements of this standard may be certified as per the conformity assessment schemes under the provisions of the *Bureau of Indian Standards Act*, 2016 and the Rules and Regulations framed thereunder, and the product(s) may be marked with the Standard Mark.

#### Scope

This document specifies requirements, test methods, instructions for use and marking for multifunction handpieces (colloquially called "syringes") intended to be used in the oral cavity of the patient.

This document does not apply to dental handpieces and motors, intraoral cameras, dental polymerisation lamps, powered scalers, powder jet handpieces, prophy handpieces, suction cannulas and saliva ejectors.

The technical content of the document has not been enclosed as it is identical with the corresponding ISO standard. For details, please refer to ISO 22569: 2020 or kindly contact:

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