



DPC: 24/30482502 DC

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Project No. 2023/02644

Responsible committee: CH/194 Biological evaluation of medical devices

Interested committees:

Title: Draft BS EN ISO 10993-17 :2023/Amd 1 Biological evaluation of medical devices. Part 17: Toxicological risk assessment of medical device constituents — Amendment.

Please notify the secretary if you are aware of any keywords that might assist in classifying or identifying the standard or if the content of this standard

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- ii) affects other national standard(s)
- iii) requires additional national guidance or information

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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 Please indicate if you are aware of any reason why this draft standard should not be published as a British Standard.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
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1	2	(3)	4	5	(6)	7
MB	Clause No./ Subclause No./Annex (e.g. 3.1)	Paragraph/ Figure/Table/Note	Type of comment	Comment (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.	



DRAFT Amendment

ISO 10993-17:2023/ DAM 1

Biological evaluation of medical devices —

Part 17: Toxicological risk assessment of medical device constituents

AMENDMENT 1

Évaluation biologique des dispositifs médicaux —

*Partie 17: Appréciation du risque toxicologique des constituants
des dispositifs médicaux*

AMENDEMENT 1

ICS: 11.100.20

ISO/TC 194

Secretariat: **DIN**

Voting begins on:
2024-07-31

Voting terminates on:
2024-10-23

This document is circulated as received from the committee secretariat.

ISO/CEN PARALLEL PROCESSING

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