



SLOVENSKI STANDARD

SIST EN ISO 10993-17:2024/A1:2026

01-februar-2026

Biološko ovrednotenje medicinskih pripomočkov - 17. del: Toksikološka ocena tveganja sestavin medicinskih pripomočkov - Dopolnilo A1 (ISO 10993-17:2023/Amd 1:2025)

Biological evaluation of medical devices - Part 17: Toxicological risk assessment of medical device constituents - Amendment 1 (ISO 10993-17:2023/Amd 1:2025)

Biologische Beurteilung von Medizinprodukten - Teil 17: Toxikologische Risikobewertung von Medizinproduktbestandteilen - Änderung 1 (ISO 10993-17:2023/Amd 1:2025)

Évaluation biologique des dispositifs médicaux - Partie 17: Appréciation du risque toxicologique des constituants des dispositifs médicaux - Amendement 1 (ISO 10993-17:2023/Amd 1:2025)

<https://standards.iteh.ai>
Ta slovenski standard je istoveten z: EN ISO 10993-17:2023/A1:2025

ICS:

11.100.20	Biološko ovrednotenje medicinskih pripomočkov	Biological evaluation of medical devices
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SIST EN ISO 10993-17:2024/A1:2026 en,fr,de

EUROPEAN STANDARD
NORME EUROPÉENNE
EUROPÄISCHE NORM

**EN ISO 10993-
17:2023/A1**

December 2025

ICS 11.100.20

English Version

**Biological evaluation of medical devices - Part 17:
Toxicological risk assessment of medical device
constituents - Amendment 1 (ISO 10993-17:2023/Amd
1:2025)**

Évaluation biologique des dispositifs médicaux - Partie
17: Appréciation du risque toxicologique des
constituants des dispositifs médicaux - Amendement 1
(ISO 10993-17:2023/Amd 1:2025)

Biologische Beurteilung von Medizinprodukten - Teil
17: Toxikologische Risikobewertung von
Medizinproduktbestandteilen - Änderung 1 (ISO
10993-17:2023/Amd 1:2025)

This amendment A1 modifies the European Standard EN ISO 10993-17:2023; it was approved by CEN on 2 November 2025.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for inclusion of this amendment into the relevant national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the CEN-CENELEC Management Centre or to any CEN member.

This amendment exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the CEN-CENELEC Management Centre has the same status as the official versions.

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