SLOVENSKI STANDARD
SIST EN ISO 26825:2022

01-april-2022

Anestezijska in dihalna oprema - Nalepke za injekcijske brizge z zdravili, ki se uporabljajo pri anesteziji - Barve, oblika in lastnosti (ISO 26825:2020)

Anaesthetic and respiratory equipment - User-applied labels for syringes containing drugs used during anaesthesia - Colours, design and performance (ISO 26825:2020)

Anästhesie und Beatmungsgeräte - Aufkleber für Spritzen mit Arzneimitteln zur Anwendung bei der Anästhesie, die vom Anwender angebracht werden - Farben, Design und Leistung (ISO 26825:2020)

Matériel d'anesthésie et de réanimation respiratoire - Étiquettes apposées par l'utilisateur sur les seringues contenant des médicaments utilisés pendant l'anesthésie - Couleurs, aspect et propriétés (ISO 26825:2020)

Ta slovenski standard je istoveten z: **EN ISO 26825:2022**

ICS:

11.040.10	Anestezijska, respiratorna in reanimacijska oprema	Anaesthetic, respiratory and reanimation equipment
11.040.25	Injekcijske brizge, igle in katetri	Syringes, needles an catheters

SIST EN ISO 26825:2022 **en,fr,de**

EUROPEAN STANDARD

NORME EUROPÉENNE

EUROPÄISCHE NORM

EN ISO 26825

February 2022

ICS 11.040.10; 11.040.25

English Version

Anaesthetic and respiratory equipment - User-applied labels for syringes containing drugs used during anaesthesia - Colours, design and performance (ISO 26825:2020)

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This European Standard 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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EUROPEAN COMMITTEE FOR STANDARDIZATION
COMITÉ EUROPÉEN DE NORMALISATION
EUROPÄISCHES KOMITEE FÜR NORMUNG

CEN-CENELEC Management Centre: Rue de la Science 23, B-1040 Brussels

Contents

Page

European foreword

The text of ISO 26825:2020 has been prepared by Technical Committee ISO/TC 121 "Anaesthetic and respiratory equipment" of the International Organization for Standardization (ISO) and has been taken over as EN ISO 26825:2022 by Technical Committee CEN/TC 215 "Respiratory and anaesthetic equipment" the secretariat of which is held by BSI.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by August 2022, and conflicting national standards shall be withdrawn at the latest by August 2022.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. CEN shall not be held responsible for identifying any or all such patent rights.

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Endorsement notice

The text of ISO 26825:2020 has been approved by CEN as EN ISO 26825:2022 without any modification.

INTERNATIONAL STANDARD

ISO 26825

Second edition
2020-10

Anaesthetic and respiratory equipment — User-applied labels for syringes containing drugs used during anaesthesia — Colours, design and performance

Matériel d'anesthésie et de réanimation respiratoire — Étiquettes apposées par l'utilisateur sur les seringues contenant des médicaments utilisés pendant l'anesthésie — Couleurs, aspect et propriétés

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Contents

Page

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).

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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 121, *Anaesthetic and respiratory equipment*, Subcommittee SC 1, *Breathing attachments and anaesthetic machines*.

This second edition cancels and replaces the first edition (ISO 26825:2008), which has been technically revised. The main changes compared to the previous edition are as follows:

— change of the former requirement on the drug name into a recommendation in 5.4.1;

— revision of the labels for benzodiazepines, suxamethonium, muscle relaxant reversal drugs and adrenaline;

— addition of a requirement on the size of diagonal stripes on the label in 5.4.4;

— revision of the indication of the concentration of the drug on the label;

— addition of recommendations on labelling of ready mixed drugs;

— deletion of the colour fluorescent red;

— revision of Table 1 on background colour coding, Table 2 on representation of colours and Table A.1 on examples of alternative colour designations, and merging of the relevant information into one table (Table 1).

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.

