

SLOVENSKI STANDARD SIST EN ISO 3826-1:2019/oprA1:2022

01-september-2022

Plastični zložljivi vsebniki za človeško kri in krvne komponente - 1. del: Običajni vsebniki - Dopolnilo 1 (ISO 3826-1:2019/DAM 1:2022)

Plastics collapsible containers for human blood and blood components - Part 1: Conventional containers - Amendment 1 (ISO 3826-1:2019/DAM 1:2022)

Kunststoffbeutel für menschliches Blut und Blutbestandteile - Teil 1: Konventionelle Beutel - Änderung 1 (ISO 3826-1:2019/DAM 1:2022)

Poches en plastique souple pour le sang et les composants du sang - Partie 1: Poches conventionnelles - Amendement 1 (ISO 3826-1:2019/DAM 1:2022)

1b915e0f6/sist-en-iso-3826-1-2019-opra1-2022

Ta slovenski standard je istoveten z: EN ISO 3826-1:2019/prA1

ICS:

11.040.20 Transfuzijska, infuzijska in Transfusion, infusion and injekcijska oprema injection equipment

SIST EN ISO 3826-1:2019/oprA1:2022 en,fr,de

SIST EN ISO 3826-1:2019/oprA1:2022

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 3826-1:2019/oprA1:2022 https://standards.iteh.ai/catalog/standards/sist/859db9cf-c6e5-48f3-84db-2e41b915e0f6/sist-en-iso-3826-1-2019-opra1-2022

DRAFT AMENDMENT ISO 3826-1:2019/DAM 1

ISO/TC 76

Secretariat: DIN

Voting begins on: **2022-06-16**

Voting terminates on: 2022-09-08

Plastics collapsible containers for human blood and blood components —

Part 1: **Conventional containers** AMENDMENT 1

Poches en plastique souple pour le sang et les composants du sang — Partie 1: Poches conventionnelles AMENDEMENT 1

(standards.iteh.ai)

ICS: 11.040.20

<u>SISTEN ISO 3826-1:2019/oprA1:2022</u> https://standards.iteh.ai/catalog/standards/sist/859db9cf-c6e5-48f3-84db-2e41b915e0f6/sist-en-iso-3826-1-2019-opra1-2022

This document is circulated as received from the committee secretariat.

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.

ISO/CEN PARALLEL PROCESSING



Reference number ISO 3826-1:2019/DAM 1:2022(E) ISO 3826-1:2019/DAM 1:2022(E)

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 3826-1:2019/oprA1:2022 https://standards.iteh.ai/catalog/standards/sist/859db9cf-c6e5-48f3-84db-2e41b915e0f6/sist-en-iso-3826-1-2019-opra1-2022



COPYRIGHT PROTECTED DOCUMENT

© ISO 2022

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 Email: copyright@iso.org Website: www.iso.org

Published in Switzerland

ISO 3826-1:2019/DAM 1:2022(E)

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 76, *Transfusion, infusion and injection, and blood processing equipment for medical and pharmaceutical use*.

A list of all parts in the ISO 3826 series can be found on the ISO website.5-4863-84db-

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 <u>www.iso.org/members.html</u>.

SIST EN ISO 3826-1:2019/oprA1:2022

iTeh STANDARD PREVIEW (standards.iteh.ai)

SIST EN ISO 3826-1:2019/oprA1:2022 https://standards.iteh.ai/catalog/standards/sist/859db9cf-c6e5-48f3-84db-2e41b915e0f6/sist-en-iso-3826-1-2019-opra1-2022