



**SLOVENSKI STANDARD  
SIST EN ISO 10477:2020**

**01-december-2020**

**Nadomešča:  
SIST EN ISO 10477:2018**

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**Zobozdravstvo - Polimerni materiali za prevleke in mostičke (ISO 10477:2020)**

Dentistry - Polymer-based crown and veneering materials (ISO 10477:2020)

Zahnheilkunde - Polymerbasierte Kronen- und Verblendwerkstoffe (ISO 10477:2020)

Médecine bucco-dentaire - Produits à base de polymères pour couronnes et facettes (ISO 10477:2020)

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**Ta slovenski standard je istoveten z: EN ISO 10477:2020**  
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**ICS:**

11.060.10      Zobotehnični materiali      Dental materials

**SIST EN ISO 10477:2020**      **en**

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EUROPEAN STANDARD

EN ISO 10477

NORME EUROPÉENNE

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English Version

## Dentistry - Polymer-based crown and veneering materials (ISO 10477:2020)

Médecine bucco-dentaire - Produits à base de polymères pour couronnes et facettes (ISO 10477:2020)

Zahnheilkunde - Polymerbasierte Kronen- und Verblendwerkstoffe (ISO 10477:2020)

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## European foreword

This document (EN ISO 10477:2020) has been prepared by Technical Committee ISO/TC 106 "Dentistry" in collaboration with Technical Committee CEN/TC 55 "Dentistry" the secretariat of which is held by DIN.

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of an identical text or by endorsement, at the latest by May 2021, and conflicting national standards shall be withdrawn at the latest by May 2021.

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2020-10

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**Dentistry — Polymer-based crown and  
veneering materials**

*Médecine bucco-dentaire — Produits à base de polymères pour  
couronnes et facettes*

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CH-1214 Vernier, Geneva  
Phone: +41 22 749 01 11  
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## 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](http://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](http://www.iso.org/patents)).

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For an explanation on 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 the following URL: [www.iso.org/iso/foreword.html](http://www.iso.org/iso/foreword.html).

This document was prepared by Technical Committee ISO/TC 106, *Dentistry*, Subcommittee SC 2, *Prosthetic materials*, in collaboration with the European Committee for Standardization (CEN) Technical Committee CEN/TC 55, *Dentistry*, in accordance with the Agreement on technical cooperation between ISO and CEN (Vienna Agreement).

This fourth edition cancels and replaces the third edition (ISO 10477:2018), which has been technically revised.

The main changes compared to the previous edition are as follows:

- missing exemption condition for opaque resin test specimens in line 6 of [Table 1](#) has been added;
- required number of specimen in [7.8](#) (Shade consistency and colour stability) has been corrected;
- several editorial changes have been made to describe the procedures more precisely;
- appropriate intervals for the storage times described in the procedures have been added.

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](http://www.iso.org/members.html).

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## Introduction

Specific qualitative and quantitative test methods for demonstrating freedom from unacceptable biological hazards are not included in this document, but it is recommended that, for the assessment of possible biological hazards, reference should be made to ISO 10993-1 and ISO 7405.

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