Speedex light body

Carefully read the instructions prior to use.

Product description

Speedex® is a two-component impression material based on condensation-type polysiloxanes. Speedex is intended for the re ing of the current physical situation in a patient's mouth ("dental impression material") for the purpose of repairing, reshaping or replacing the patient's teeth. This includes base materials (Speedex® putty soft; Speedex® putty; Speedex® light body; Speedex® medium) and the universal activator (Speedex® Universal Activator) After manual mixing of base and activator, the Speedex materials form pastes which are used individually or in combination as dental impression material, usually together with a standard commercial or individual impression tray and using conventional impression

Composition Speedex light body

Condensation-type silicone elastomer, free-flowing consistency.

- Shade: dark blue

Indications

- Correction material for the correction impression technique
- Syringe material for the two-phase impression technique
- Impression material for relining

Contraindication

- Do not use in the case of known allergies to ingredients of the Speedex impression materials
- Loose teeth can be further loosened or extracted by taking an
- Material combinations with impression materials of other manufacturers are not permitted

Safety instructions

- Eyes, respiratory organs and skin can be irritated by Speedex Universal Activator
- Impressions that come into contact with the oral mucosa may be contaminated. Observe disinfection instructions
- Only supplied to dentists and dental laboratories or upon their
- Carefully read the Speedex instructions for use before using

NOTICE

Contains nano materials (bonded particles)

Side effects / Interactions

No harmful reactions or secondary effects on patients and / or dental personnel are known.

Users / Patient target groups

The use of Speedex impression materials in the patient's mouth may only be undertaken by specialist personnel such as dentists and dental assistants with special qualifications. Suitable for all patient groups

PREPARATION

All impression trays that have been designed for dental use, either prefabricated or individual, can be used for impression taking. For perfect adhesion, we recommend applying a thin layer of COLTENE® Adhesive to the trays before use. This prevents detachment of the impression material during removal from the oral cavity. Detailed information on the application can be found in the COLTENE Adhesive Instructions for Use.

PROPER USE

Dosage

Dispense Speedex Universal Activator and Speedex light body on a mixing block in two equal strands. Make sure that the tubes are closed immediately after use.

Take the following safety precautions when working with Speedex Universal Activator:

- Protective glasses are recommended. In case of contact with eyes, rinse immediately with water and consult an ophthalmologist
- Avoid direct skin contact
- Always wear protective gloves when working (vinyl gloves are recommended)
- Do not swallow or ingest

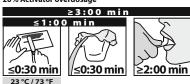
Application time Normal dosage



Intensive mixing, high temperatures and humidity accelerate the curing of the impression materials. Low temperatures slow down the curing process.

By over- and underdosing the Speedex Universal Activator the processing time can be accelerated or slowed down.

20% Activator overdosage



20 % Activator underdosage



Mixing

Take up one component (Universal Activator or light body) with spatula and spread evenly on the other component. Then pick up both components with the spatula and spread out on the mixing block. Repeat this procedure until the mixture is homogeneous. Please observe the specified processing times.

Depending on the impression method, apply the impression material in the patient's mouth or to the impression tray. Insert the impression tray into the oral cavity and press for 2-3 seconds and keep

Test methods for correct application

Material residue needs considerably more time to set at room temperature than in the patient's mouth. You should therefore check intraorally if the material is set before removing from the mouth. If the manual pressure test does not leave a pressure mark in the impression material after the time specified in the chapter on "Dosage", the impression can be removed from the patient's mouth. However, if the test leaves a visible pressure mark in the material, the material is not yet fully cured.

CLEANING & DISINFECTION

Cleaning of the impression

After cleaning, the final impressions can be disinfected with commercially available disinfectant solutions (e.g. Dürr Dental SE, MD 520 impression disinfection) specifically designed for dental impression materials and dried using air pressure. If hydrogen peroxide is used for disinfection, it is recommended to thoroughly rinse with lukewarm water afterwards in order to avoid bubble formation.

The impression can be removed with a blunt instrument. Adhesive residues can be dissolved by placing in commercially available universal solvents or benzine.

Manufacture of models

The ideal time lies between 30 min and 72 h after taking the impression. The surface tension will be reduced and pouring will be facilitated if the impression is briefly washed out with a detergent and rinsed thoroughly in lukewarm clear water afterwards. All industry-standard dental stone model materials (e.g. Fujirock Dental Stone, Hard Rock Dental Stone) can be used

Shelf life and storage

- Expiry date: see primary packaging
- Storage temperature: 15-23°C / 59-73°F
- Relative humidity: $50\% \pm 10\%$
- Shelf life after first opening: 3 months

Protect against exposure to heat and sun. Avoid extreme temperature fluctuations. Close container and tubes immediately after use.

Ora

Speedex impression materials can be disposed in small quantities in household waste. Special country-specific regulations may apply.

Technical data according to ISO 4823

Measurements were taken at 23°C / 73°F room temperature and 50% relative humidity.

king time:	00:30 min
rking time:	≤1:30 min
al setting time:	3:00 min
king ratio Putty/Activator:	5 ml/1 ml
	5 g/0.88 g

Reporting obligation

All serious incidents occurring in conjunction with this product must be reported immediately to the manufacturer as well as to the

SAFETY DATA SHEET







Consult instructions for use



Keep away from sun light



Temperature limitation Marking of Conformity Europe



Identification for Ukraine



Legal Manufacturer



Expiry Date



Batch Code



RX only



Medical Device

μЛ

Manufacturing Date

UDI

Unique Device Identifier

EC REP

European Authorized Representative

REF Reference Number

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