Prent Des

VAH listed

Instrument disinfection for dental impression materials

- ready for use
- does not influence dimensional accuracy
- no corrosion of surfaces

Product type

Ready-to-use solution for the instrument disinfection of dental impression materials made of alginate, A-silicone, C-silicone and polyether, as well as prosthetic and orthodontic workpieces. For professional use only.

Composition

100 g solution contain:
0.48 g dimethylbenzylalkylammonium chloride
0.20 g ethandial (glyoxal)
0.01 g pentandial (glutaral)

Chemical-physical data

Appearance:	light blue, clear solution
pH value:	5.5 +/- 1.0
Density (20 °C):	1.000 +/- 0.003

Range of application

Instrument disinfection for dental impression materials.

Application

Thoroughly rinse the impression material or dental technical workpiece under running water of drinking water quality immediately after removing it from the mouth. Briefly immerse dental impressions made from alginate, A-silicone, C-silicone and polyether in the disinfection solution and let them dry for 15 minutes. Immerse other workpieces (e.g. prosthetic or orthodontic workpieces) in the solution according to the contact time. Rinse with drinking water afterwards. The manufacturer's specifications for the materials must be observed.

Delivery forms

Single pack	Sales unit	Ref. No.
5 litre canister	1 x 5 L	09983

Efficacy and contact times

Application recommendation for the disinfection of dental impressions acc. to EN 14885 (undiluted, tested at 20 °C, dirty conditions):

bactericidal, yeasticidal (EN 13727, EN 14561, EN 13624, EN 14562)	5 min.
virucidal against enveloped viruses (EN 14476, EN 17111)	15 min.

Application notes

Change the working solution immediately if there is any visible contamination, but at least once every working day. The working solution must not be mixed with other process chemicals.

Storage

Store at 5 °C to 25 °C.

Important information

Contains glyoxal, glutaral. May produce an allergic reaction. Safety data sheet available on request. According to the EU Medical Device Regulation, users/patients are obligated to report any serious incident that has occurred in relation to the device to the manufacturer and the competent authority of the EU Member State in which the user/patient is established.

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