

EU - Declaration of Conformity for Medical Devices

according to Annex IX Chapter I of Regulation (EU) 2017/745 on medical devices

Registration number (SRN):

Manufacturer:

DETAX GmbH
Street:

Carl-Zeiss-Straße 4
City:

Country:

Germany

We hereby declare under our sole responsibility that the products in the Annex with the Basic UDI-DI

++EDET0255N

and the classification according to Annex VIII of Regulation (EU) 2017/745

Class IIa, rule 5, sentence 2

and the intended use:

Resin material for the production of earmolds and ITE shells in 3D printing processes

comply with the relevant provisions of the following EU legislation:

Regulation (EU) 2017/745 on medical devices

as well as the standards and common specifications:

DIN EN ISO 15223-1:2022 DIN EN ISO 20417:2022 DIN EN ISO 10993-1:2021

DIN EN 62366-1:2021 DIN EN ISO 14971:2022

with the participation of the following management systems

DIN EN ISO 13485:2021

and the conformity assessment procedure referred to in the header

by the notified body: 0483

mdc medical device certification GmbH

Kriegerstr. 6

D-70191 STUTTGART

This declaration is valid until a significant change in the specification or manufacture of the product or until a change in the above-mentioned standards, common specifications, management systems and legal regulations, maximally until the expiry of the certificate with the No.:

D1002700058 in force until 31.07.2028

Ettlingen, 18.07.2024

Thorsten Preiss Managing Director Dr. Sönke Rössing Managing Director

Jak Rossing



Annex to

EU-Declaration of Conformity for Medical Devices

Date: 18.07.24

Page:

Basic UDI-DI: ++EDET0255N

Distributed by: Egger Otoplastik +, DEUTSCHLAND

UDI-DI Article

+EDET034181 e3D.M01 blue, 1000 g

Egger Art.-Nr. 43014

