

Registration number (SRN): **DE-MF-000022666**
Manufacturer: **DETAX GmbH**
Street: **Carl-Zeiss-Straße 4**
City: **76275 Ettlingen**
Country: **Germany**

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

++EDET0815Y

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

Class IIa, rule 19, sentence 1, 3rd indent

and the intended use:

Plastic material for the production of 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

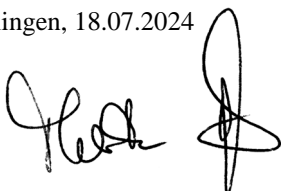
and the conformity assessment procedure referred to in the header
by the notified body: 0483

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

Basic UDI-DI: ++EDET0815Y*Distributed by: Egger Otoplastik +, DEUTSCHLAND*

UDI-DI	Article
+EDET044541	e3D.S01 white, 1000 g Egger Art.-Nr. 43028

