

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

**++EDET0245L**

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

**Class IIa, rule 5, sentence 2**

and the intended use:

**Earmold silicone**

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**

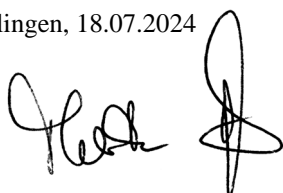
and the conformity assessment procedure referred to in the header

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: ++EDET0245L***Distributed by: Egger Otoplastik +, DEUTSCHLAND*

<b>UDI-DI</b>	<b>Article</b>
+EDET029161	egger flex/AB 25 t farbl.-transp. 8x50ml Egger Nr. 20735

