

## EC-Declaration of Conformity for Medical Devices

(in accordance with annex II of the Council Directive for Medical Devices 93/42/EEC)

**Name of manufacturer:** VOCO GmbH

**Address:** Anton-Flettner-Straße 1 - 3  
27472 Cuxhaven  
Germany

We declare that the product

**Article:** Meron Plus QM

**Type:** Resin reinforced glass ionomer luting cement

**Item Number:** see annex

**Class:** IIa

**Rule:** 8

corresponds to the regulations of the following Council Directive :  
**93/42/EEC for Medical Devices**  
and that we take full responsibility for issuing this declaration.

**Name and address of notified body**

**MedCert GmbH**  
**Pilatuspool 2**  
**20355 Hamburg**  
**Germany**  
**Identification number : 0482**

**Valid as long as the certificate in accordance with 93/42 EEC is valid**

City: Cuxhaven

Date: February 18, 2021

*Dr. A. Less*

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Regulatory Affairs

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(in accordance with annex II of the Council Directive for Medical Devices 93/42/EEC)

**Name of manufacturer:** VOCO GmbH

### ANNEX

**Article:** Meron Plus QM

REF #	Description
1727	QuickMix Syringe 8.5 g
1728	QuickMix Syringe 3 x 8.5 g

City: Cuxhaven

Date: February 18, 2021

*Dr. A. Lenz*

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Regulatory Affairs