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: Rebilda DC

Type: Dual-curing flowable core build-up and post luting system

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 16, 2021

Dr. A. Lenz

Regulatory Affairs

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

ANNEX

Article: Rebilda DC

REF #	Description
1395	Set cartridge 50 g dentine, Futurabond DC bottle each of 4 ml liquid 1 and 2, dispenser type 3
1396	Cartridge 50 g dentine
1397	Cartridge 50 g blue
1398	Cartridge 50 g white
1402	Set QuickMix Syringe 10 g dentine, Futurabond DC bottle each of 4 ml liquid 1 and 2
1403	QuickMix Syringe 10 g dentine
1404	QuickMix Syringe 10 g blue
1405	QuickMix Syringe 10 g white

City: Cuxhaven

Date: February 16, 2021

Dr. A. Lenz

Regulatory Affairs