EC-Declaration of Conformity for Medical Devices

(in accordance with annex IX chapter I of the Regulation for Medical Devices (EU) 2017/745)

Name of manufacturer:	VOCO GmbH
Address:	Anton-Flettner-Straße 1 - 3 27472 Cuxhaven Germany
SRN:	DE-MF-000008601

We declare that the product:

Article:	Ufi Gel P
Basic UDI-DI:	++E2210475000000000HX
Risk class acc. to Annex VIII:	IIa
Applied rule:	5
Item Number:	see item list
Type:	Permanently soft, cold-curing relining material on silicone base

corresponds to the regulations of the following Council Regulation:

(EU) 2017 / 745 for medical devices

and that we take full responsibility for issuing this declaration.

Name and identification number Notified Body:

DNV MedCert GmbH
NB number: 0482
Certificate No.: 0675GB448260109

Declaration valid until: January 21, 2031

Digitally signed document - see signature page

EC-Declaration of Conformity for Medical Devices

(in accordance with annex IX chapter I of the Regulation for Medical Devices (EU) 2017/745)

Name of manufacturer: VOCO GmbH

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

ITEM LIST

Article: Ufi Gel P

REF #	Description
2070	Tube base 50 ml, catalyst 50 ml, glazing base 10 ml, glazing catalyst 10 ml, adhesive bottle 10 ml
2071	Tube 2 × 50 ml
2076	Bottle 10 ml adhesive
2077	Bottle 2 × 10 ml Glazing

Signatures

This document has been electronically signed by:

Andreas Lawerenz

PRRC Technical Documentation

Cuxhaven, 14.01.2026

QES (EU) - Signed via Certifaction

Verify signatures

The signatures can be verified in Adobe Acrobat Reader.

Powered by Certifaction

Learn more about privacy-first eSigning at **Certifaction.com**