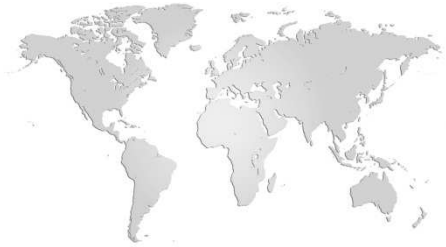
EC CERTIFICATE

for the Quality Assurance System



according the Directive 93/42/EEC, Annex II excluding section (4)

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the company

bonyf AG
Heiligkreuz 16, 9490 Vaduz, Liechtenstein

applies a quality assurance system according to the Directive 93/42/EEC Annex II for the medical devices listed in the annex. The approval is based on the result of the re-certification audit report no. 50537-Z5-00, the decision dated 2018-02-05 and is only valid in connection with the successful performance of the annual surveillance audits.

This certificate is valid from 2020-07-27 to 2023-02-06

Registration No.: 50537-16-06

DEKRA Certification GmbH Stuttgart, Handwerkstraße 15, D-70565

Ruth Delbeck-Bayer
DEKRA Certification GmbH Stuttgart; 2020-07-27
Notified Body ID-number: 0124

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-295.10.02

Annex to the EC Certificate No. 50537-16-06

Valid from 2020-07-27 to 2023-02-06

Revision status of the annex: 0 dated 2020-07-27

Devices/device categories included in the certificate:

Class II a:

- MD 0402
  - Home use denture care products
    - Denture repair system

Klasse II b:

- MD 0108
  - Non-active medical devices for disinfecting, cleaning, rinsing
    - Disinfecting tablets for removable dental appliances
      - NitrAdine Desinfecting Tablets

DEKRA Certification GmbH • D-70565 Stuttgart, Handwerkstraße 15 • DEKRA Certification

Ruth Delbeck-Bayer
DEKRA Certification GmbH, Stuttgart, 2020-07-27
Notified Body ID-number: 0124