

EC Declaration of Conformity

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DC-067

Rev.No

7

Manufacturer:

DiaDent Group International 16, Osongsaengmyeong 4-ro Osong-eup Heungdeok-gu, Cheongju-si, Chungcheongbuk-do,

28161, Republic of Korea

We, the manufacturer, herewith declare that the products

Authorized Representative:

DiaDent Europe B.V.

Antennestraat 70, 1322AS Almere, the Netherlands

INSTRUMENT TRAY

(Divided Tray, Stainless Steel Tray, Instrument Tray Type A, B(Perforated, Non-Perforated))

12143

GMDN :

meet the provisions of The Medical Devices Directive 93/42/EEC amended by 2007/47/EC which apply to them.

The medical device has been assigned to ClassI(Rule 1) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC.

The product concerned has been designed and manufactured under a quality management system according to Annex VII of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC.

following the procedure relating to the EC Declaration of Conformity set out in Annex VII of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC

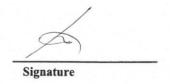
The above mentioned declaration of conformity is exclusively under the responsibility of

DiaDent Group International

16, Osongsaengmyeong 4-ro Osong-eup Heungdeok-gu, Cheongju-si Chungcheongbuk-do, 28161, Republic of Korea

2020-06-01

Date





DiaDent Group International

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