DiaDent® Serving 122 countries around the world

EC Declaration of Conformity

Doc No : DC-023
Rev.No : -5-

Manufacturer:
DiaDent Group International
16, Osongsaengmyeong 4-ro Osong-eup Heungdeok-gu, Cheongju-si Chungcheongbuk-do, 28161, Republic of Korea

Authorized Representative:
DiaDent Europe B.V.
Antennestraat 70, 1322AS Almere, the Netherlands

We, the manufacturer, herewith declare that the products

DIA-SPIRAL FILLER

meet the provisions of Directive 93/42/EEC and 2007/47/EEC which apply to them.

The medical device has been assigned to class I (Rule 5) according to Annex IX of the Directive 93/42/EEC. It bears the mark

The product concerned has been designed and manufactured under a quality management system according to Annex VII of Directive 93/42/EEC.

following the procedure relating to the EC Declaration of Conformity set out in Annex VII of Directive 93/42/EEC.

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

2016.03.22

Place , date

signature,