

EC Declaration of Conformity

Doc No : DC-046

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

NI-TI ARCH WIRES

(Super Elastic, Thermal Activated)

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


The medical device has been assigned to class II a (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,,,