

**EC Declaration of Conformity**

Doc No : DC-078

Rev.No : -3-

*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**DiaDent**<sup>®</sup>

We, the manufacturer, herewith declare that the products

**Dia-Cem<sup>TM</sup>TempNE**

UMDNS-Code: 16-711

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

The medical device has been assigned to class IIa (Rule 7) according to Annex IX of the Directive 93/42/EEC. It bears the mark

**CE 0197**

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

**DiaDent**<sup>®</sup>

Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

**TÜV Rheinland LGA Products GmbH**  
**Tillystraße 2, 90431, Nürnberg, Germany**

Certificate No.: HD 60102396 0001

Issue date: 2016-01-15

Expiry date: 2020-06-01

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

**DiaDent**<sup>®</sup>

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

2016.01.15

Place, date

signature,