

**EC Declaration of Conformity**

Doc No : DC-013

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 Plus G7**

(including system components and accessories)

UMDNS-Code: **16-388**

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 8) 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 II of Directive 93/42/EEC.



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**

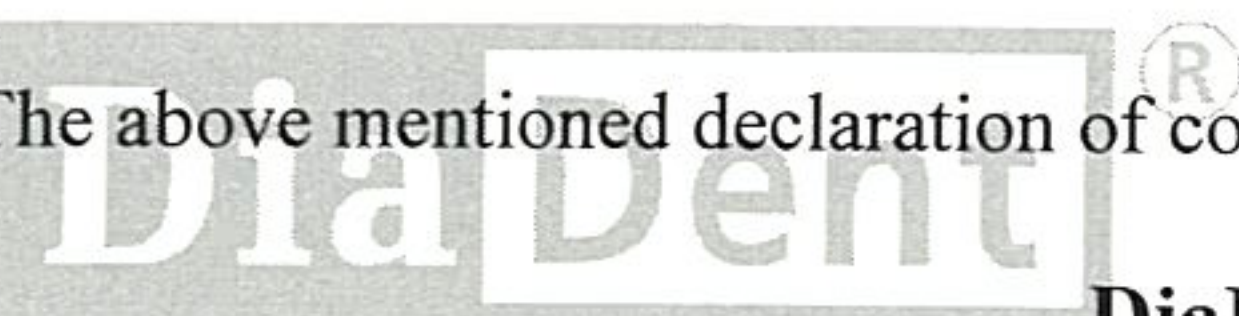
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following the procedure relating to the EC Declaration of Conformity set out in Annex II of Directive 93/42/EEC.

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

2016.01.15

Place, date

signature,



EC Declaration of Conformity ISO 9001 ISO 13485

**DiaDent Group International**

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