

Doc No : DC-001

Rev.No:

#### 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

## **Gutta Percha Points**

**GMDN** 31872

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 Class IIa(Rule 8) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No.

DD 60149569 0001

Issue date

2020-05-25

Expiry date

2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex V 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

DiaDent Group International

Date Chunacheona buk-do, 28161, Korea

Signature

C Certified (50 13485



**DiaDent Group International** 

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

Tel: 82-43-266-2315 http://www.diadent.co.kr



DC-016 Doc No : Rev.No: 6

#### 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

# Sterile Paper Point

**GMDN** 38777

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 Class IIa(Rule 6) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No DD 60149569 0001

Issue date 2020-05-25 2024-05-26 Expiry date

following the procedure relating to the EC Declaration of Conformity set out in Annex Vof 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

DiaDent Group International

Date

Signature





DiaDent Group International

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

Tel: 82-43-266-2315 Fax: 82-43-262-8658 http://www.diadent.co.kr



Doc No :

DC-076

Rev.No :

6

#### 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

# e-Temp

**GMDN** 

31783

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 Class IIa(Rule 7) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

C € 0197

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No

HD 60149568 0001

Issue date

2020-05-25

Expiry date

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following the procedure relating to the EC Declaration of Conformity set out in Annex II 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-cup Heungdeok-gu, Cheongju-si Chungcheongbuk-do, 28161, Republic of Korea

2020-06-01

Date

DiaDent Group International

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

Tel: 82-43-266-2315

Fax: 82-43-262-8658



www.diadent.co.kr



Doc No : DC-103

Rev.No : 2

#### 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

# DiaPrep Pro

(including system components and accessories)

GMDN

45500

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 Class IIa(Rule 6) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

C € 0197

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No : HD 60149568 0001

Issue date : 2020-05-25 Expiry date : 2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex II 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 Dia Deri Group Inte

Date 16. Osongsaengmyeong 4-ro. Osong-eup, Heuro

Cheong, u-si. Chungcheong buk-do, 28161, Korea

DiaDent Group International

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

Tel: 82-43-266-2315 http://www.diadent.co.kr





Doc No : DC-105 Rev.No : 1

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

Dia Temp

GMDN

31783

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 Class IIa(Rule 7) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

C € 0197

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No

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Issue date

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

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**DiaDent Group International** 

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

Dent

2020-06-01

Diabent Group International

Date

. Chungcheong buk-do, 28161. Korea **Signature** 

**DiaDent Group International** 

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

Tel: 82-43-266-2315 http://www.diadent.co.kr





Doc No : DC-074

7

Rev.No :

#### 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

## STERI-IRRIGATION TIPS SIDE OPEN TYPE

**GMDN** 64403

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 Class IIa(Rule 6) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

# TÜV Rheinland LGA Products GmbH

Tillystraße 2, 90431, Nürnberg, Germany

Certificate No DD 60149569 0001

Issue date 2020-05-25

Expiry date 2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex Vof 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

nuhacheona buk-do ax: 82-43-**DiaDent Group International** 

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

Tel: 82-43-266-2315 http://www.diadent.co.kr





DC-102 Doc No 2 Rev.No

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

Dia-X File

**GMDN** 

40529

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 Class IIa(Rule 6) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No

DD 60149569 0001

Issue date

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following the procedure relating to the EC Declaration of Conformity set out in Annex Vof 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 Internationalent Group international 16, Osongsaengmyeong 4-ro, Osong-eup, Heungdeok-gu, Cheongju-si, Chungcheongbuk-do, 28161, Korea

Tel: 82-43-266-2315 www.diadent.co.kr





Doc No : DC-109

Rev.No : 2

#### 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

#### DiaFil

(including system components and accessories)

GMDN : 35870

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 Class IIa(Rule 8) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

( € 0197

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No : HD 60149568 0001

Issue date : 2020-05-25 Expiry date : 2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex II 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

16, Osongsaengmyeong 4-ro, Osong-eup, Heungdeok-gu, Cheongju-si, Chungcheongbuk-do, 28161, Korea
Tel: 82-43-266-2315
www.diadent.co.kr
Fax: 82-43-262-8658
E-mail: diadent@diadent.co.kr
Fax: 82-43-252-8658

Certified (50 13485)



Doc No

DC-110

Rev.No:

2

#### Manufacturer:

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

**Authorized Representative:** DiaDent Europe B.V.

Antennestraat 70, 1322AS Almere, the Netherlands

28161, Republic of Korea

We, the manufacturer, herewith declare that the products

## DiaFil Flow

(including system components and accessories)

**GMDN** 

35870

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 Class IIa(Rule 8) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No

HD 60149568 0001

Issue date

2020-05-25

Expiry date

2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex II 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

BiaDent

Signature

**DiaDent Group International** 

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

Fax: 82-43-262-8658



Tel: 82-43-266-2315 http://www.diadent.co.kr



Doc No DC-034 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

### FILE MEASURE SET

(Plastic)

**GMDN** 

64813

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. It bears the mark

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No

N/A N/A

Issue date

Expiry date

N/A

following the procedure relating to the EC Declaration of Conformity set out in Annex VIIof 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

**Group International** 

Date

mreeng 4-ro. Osong-eup, Heungdeok**-g⁄Signature** Cheong,u-si. Chungcheong buk-da, 28161. Kore

DiaDent Group International -8658

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

Tel: 82-43-266-2315

Fax: 82-43-262-8658

http://www.diadent.co.kr





Doc No : DC-103 2 Rev.No :

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

## DiaPrep Pro

(including system components and accessories)

**GMDN** 

45500

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 Class IIa(Rule 6) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

Certificate No HD 60149568 0001

2020-05-25 Issue date

2024-05-26 Expiry date

following the procedure relating to the EC Declaration of Conformity set out in Annex II 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

C Certified (50 13485)

DiaDent Group International

16, Osongsaengmyeong 4-ro, Osong-eup<sub>0</sub> Heungdeok-gu, Cheongju-si, Chungcheongbuk-do, 28161, Korea

Tel: 82-43-266-2315

Fax: 82-43-262-8658-si, Chungcheong buk-do, 28161, Korea E-mail: dladent@diadent@co.kr23'5 Fax: 82-43-262-8658

www.diadent.co.kr



DC-110 Doc No : 2 Rev.No:

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

## DiaFil Flow

(including system components and accessories)

**GMDN** 

35870

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 Class IIa(Rule 8) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

C E 0197

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

Compliance of the designated product with The Medical Devices Directive 93/42/EEC amended by 2007/47/EC has been assessed and certified by the Notified Body

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

HD 60149568 0001 Certificate No

Issue date 2020-05-25 Expiry date 2024-05-26

following the procedure relating to the EC Declaration of Conformity set out in Annex II 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-cup Heungdeok-gu, Cheongju-si Chungeheongbuk-do, 28161. Republic of Korea

2020-06-01

Date

DiaDent Group International 16, Osongsaengmyeong 4-ro, Osong-eup, Heunggeok-gu, Cheongjussi, Chungcheongkuk-do, 28161, Korea Tel: 82-43-266-2315 Fax: 82-43-262-86582-43-265-23-5

www.diadent.co.kr





Doc No :

DC-049

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

### MEDICAMENT BOTTLE

(Round Type. Square Type) **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

DiaDent Group International

16. Osong Elegmyeong in ro. Osong-eup. Heurgaeok-nu. Cheongrums, Chungs eong buk-do. 28161, Korës Tel: 82-48-266-2315 Fax: 82-43-262-8658



Tel: 82-43-266-2315 www.diadent.co.kr

Fax: 82-43-262-8658



DC-002 Doc No : 9 Rev.No:

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

# Gutta Percha Obturators (incl. Gutta Percha Cartridge)

(including system components and accessories)

**GMDN** 

31872

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 Class IIa(Rule 8) according to Annex IX of the Medical Devices Directive 93/42/EEC amended by 2007/47/EC. It bears the mark

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

Certificate No

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### 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 InternationaPiaDent Stone rovernational

16, Osongsaengmyeong 4-ro, Osong-eup, Heungdeok-gu, Cheongju-si, Chungcheongbuk-do, 28161, Korea Tel: 82-43-266-2315 Fax: 82-43-262-8658 -17-262-8658

www.diadent.co.kr

