EC Certificate Full Quality Assurance System: Certificate KR12/01766

The management system of

MEDICLUS Co., Ltd.

No. 1210, 134, Gongdan-ro, Heungdeok-gu, Cheongju-si,
Chungcheongbuk-do, Korea

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

**Dental light curing temporary filling material (Once-Fil Flow),
Dental etchant (Any-Etch),
Dental composite resin for anterior and posterior restoration (Any-Com),
Dental root canal cleanser (Endo-Prep EDTA Cream)**

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 15 January 2018 until 13 January 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 29 December 2017
Issue 8. Certified since 13 January 2012

Certification is based on reports numbered KR/SEL Y-PC/11284

Authorised by

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