

The management system of

META SYSTEMS CO., LTD.

also doing business as S-Denti Co., Ltd.

#1214-18, Sicox tower 12F, 484, Dunchon-daero, Jungwon-gu,
Seongnam-si, Gyeonggi-do, 13229, 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

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 23 November 2020 until 09 April 2024
and remains valid subject to satisfactory surveillance audits.

Issue 5. Certified since 19 April 2005
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered WW/PCI 212119

Authorised by

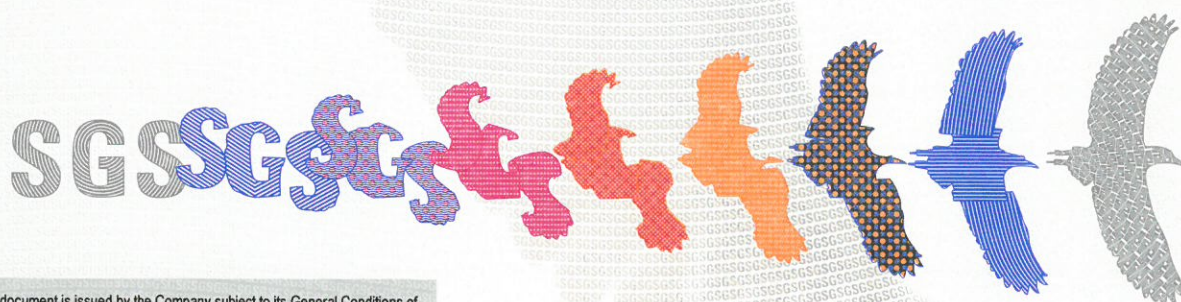


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LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

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Directive 93/42/EEC on medical devices, Annex II (excluding section 4)

Issue 5

Detailed scope

**Electronic Apex Locator for measuring the working length of the tooth
in endodontic treatment (Model: i-ROOT, i-ROOT 100,
ROOTOR and Apex ID, EQ-PEX);**

**Dental Endo Motor including Contra angle handpiece
(model : ES-100, EQ-M);**

**Dental Root Canal Obturating Materials
(model: GUTTA PERCHA CARTRIDGE);**

**Endodontic Obturation System including Pack, Fill,
Pack tip and Fill needle (model:EQ-V and elementsIC);**

Sonic Powered Endo Irrigation(model: EQ-S)

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