

The management system of

Aceonedent Korea Ind Co.

103-606, Bucheon Techno-Park, 22, Samjak-ro,
Ojeong-Gu, Bucheon-si, Gyeonggi-do,, Korea

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC on medical devices, Annex V

For the following products

**Gutta Percha Points;
Sterile Absorbent Paper Points.**

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

This certificate is valid from 16 December 2019 until 15 June 2023
and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 15 June 2015
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered KR/SEL/Y-PC/15405

Authorised by



SGS Belgium NV, Notified Body 1639

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