EC Certificate Full Quality Assurance System: KR01/53133

The management system of

Vericom Co., Ltd

48, Toegyegongdan 1-gil, Chuncheon-si, Gangwon-do 200-944,
Republic of 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 30 May 2017 until 17 September 2021 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 7 September 2019
Issue 22. Certified since 17 September 2001

Certification is based on reports numbered WW/PCI 201952)

Authorised by

SGS United Kingdom Ltd, Notified Body 0120
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SGS CE 02 0315 M2

Vericom Co., Ltd

Directive 93/42/EEC

on medical devices, Annex II (excluding section 4)

Issue 22

Detailed scope

Light-cured composite resin ;

Pit and fissure sealant ;

Bonding agents ;

Light-cured radiopaque flowable composite resin ;

Polymer-based filling, restorative material ;

Gutta Percha Points and Sterile Absorbent Paper Point ;

Etching agents ;

Temporary filling materials ;

Root canal cleanser ;

Root canal filling materials ;

Dual – cured composite resin ;

Dental ceramic block ;

Self-adhesive resin cement ;

Sterile single use dental needles ;

Dental ceramic resin ;

Dental hybrid ceramic for the restoration of carious teeth ;

Dental root canal sealing material;

Dental Glass ceramic.

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