

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)

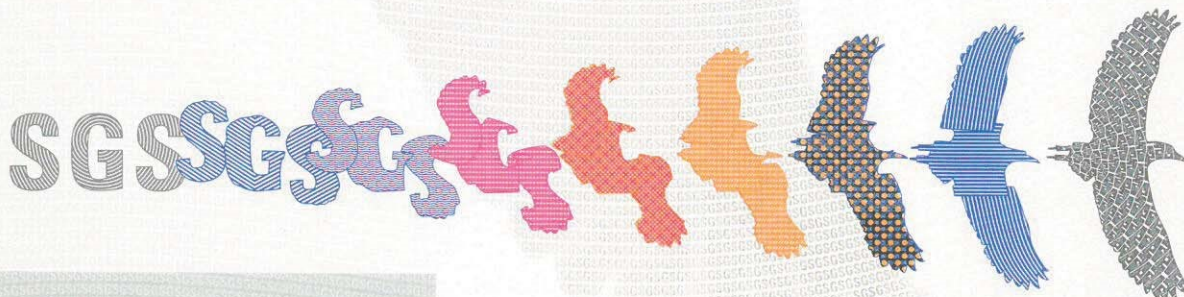
Authorised by

### SGS United Kingdom Ltd, Notified Body 0120

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# 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