

The management system of

Radiant Innovation Inc.

1F, No.3, Industrial E. 9th Rd., Science-Based Industrial Park
HsinChu, 300, Taiwan

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

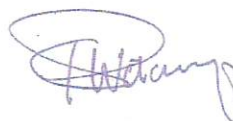
The scope of registration appears on page 2 of this certificate.

This certificate is valid from 16 December 2019 until 16 February 2022
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 16 February 2011
and first certified by SGS Belgium NV since 16 December 2019

This is a multi-site certification.
Additional site details are listed on subsequent pages

Certification is based on reports numbered TW/TPE K603835

Authorised by



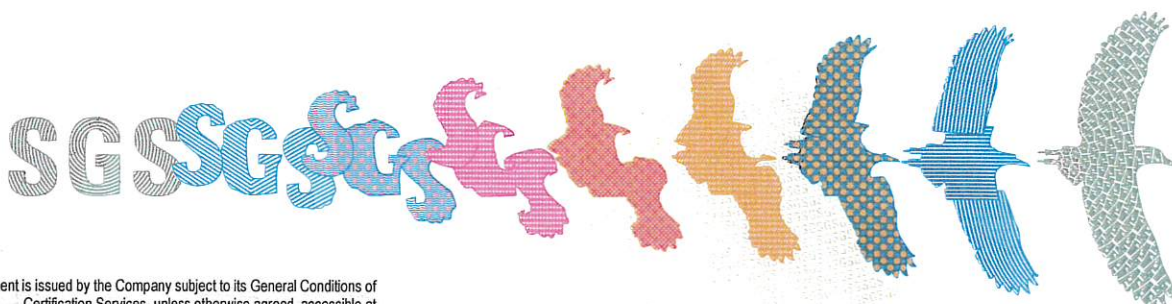
Pieter Weterings
Certification Manager

SGS Belgium NV, Notified Body 1639

SGS House Noorderlaan 87 2030 Antwerp Belgium
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LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

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Radiant Innovation Inc.

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

Issue 1

Detailed scope

**Infrared Ear Thermometers with & without Probe Cover and Infrared Forehead
Thermometers for measuring human body temperature.**

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..

Additional facilities

KunShan Radiant Innovation Co., Ltd. ,

**No. 20, TaiHong Road, WuSongJiang Development Zone, YuShan Town
KunShan City, JiangSu, China**