

The management system of

IMPLANTS DIFFUSION INTERNATIONAL s.a.r.l.

23-25 rue Emile Zola, 93100 Montreuil, France

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

**Sterile dental implants,
Sterile prosthetic components (closing cap, healing cap),
Non-sterile prosthetic components (attachments, abutments),
Non-sterile instruments with dental shank (drills, thread taps,
trephines, ridge spreaders, bladed wheels, instrument extensions,
screwdrivers with dental shank)**

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.

This certificate is valid from 16 December 2019 until 04 June 2022 and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 01 March 2012 and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered FR/MD 217462

Authorised by

SGS Belgium NV, Notified Body 1639

SGS House Noorderlaan 87 2030 Antwerp Belgium
t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com

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