

EC Certificate Full Quality Assurance System: Certificate FR12/00908

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 for implant setting
(drills, thread taps, trephines).**

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 04 July 2016 until 04 June 2019
and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 14 March 2017
Issue 5. Certified since 01 March 2012

Certification is based on reports numbered FR/MD 217462

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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