

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

LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

Page 1 of 1





This document is issued by the Company subject to its General Conditions of Certification Services, unless otherwise agreed, accessible at www.sgs.com/tems_and_conditions.htm. Attention is drawn to the limitations of liability, indemnification and jurisdictional issues established therein. The authenticity of this document may be verified at https://www.sgs.com/en/certified-clients-and-products/certified-client-directory. Any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.