

EC Certificate Full Quality Assurance System: Certificate  
Certificat CE Système complet d'assurance de qualité  
FR19/81843446

The management system of / Le système de management de

**MDJ sas**

ZI Racine Palladuc, 63650 La Monnerie Le Montel, France

has been assessed and certified as meeting the requirements of / a été audité et certifié selon les exigences de

**Directive 93/42/EEC**

on medical devices, Annex II (excluding Section 4)  
on medical devices, Annexe II (section 4 exclue)

For the following products / Pour les produits suivants

The scope of registration appears on page 2 of this certificate.  
Le domaine de certification apparaît page 2 de ce certificat.

This certificate is valid from 16 December 2019 until 31 August 2023  
and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 01 September 2003  
and first certified by SGS Belgium NV since 16 December 2019

Ce certificat est valable du 16 décembre 2019 au 31 août 2023  
et reste valide sous condition d'audits de surveillance satisfaisants.

Version 1. Certifié depuis 01 septembre 2003  
et initialement certifié par SGS Belgium NV depuis 16 décembre 2019

Certification is based on reports numbered / La certification est basée sur les rapports référence FR/MD 209213

Authorised by / Autorisé par

**SGS Belgium NV, Notified Body 1639**

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

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EC Certificate Full Quality Assurance System: Certificate  
Certificat CE Système complet d'assurance de qualité :  
FR19/81843446 continued

**MDJ sas**

**Directive 93/42/EEC**  
on medical devices, Annex II (excluding Section 4)  
on medical devices, Annexe II (section 4 exclue)

Issue / Version 1

For the following products / Pour les produits suivants

**Single use sets with sterile injector, cartridge and pusher  
for injection of intraocular lenses (IOL).**

**Sets stériles à usage unique composés de: injecteur, cartouche,  
et poussoir, pour l'injection de lentilles intra-oculaires.**

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.

Lorsque le champ d'application ci-dessus comprend un ou plusieurs dispositifs médicaux de classe III, un certificat EC Design Examination Certificate valide, conformément à l'annexe II (section 4), est obligatoire pour chaque dispositif, en addition du présent certificat, pour la mise sur le marché du dispositif.