EC Certificate Full Quality Assurance System: Certificate FR19/81843460

The management system of

EUROFEEDBACK SAS

ZI de la Petite Montagne Sud, 3, rue de l'Aubrac, 91017 EVRY, 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

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

This certificate is valid from 24 April 2020 until 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 07 February 2011
and first certified by SGS Belgium NV since 24 April 2020

Certification is based on reports numbered FR/MD 216647

Authorised by

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EUROFEEDBACK SAS

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

Issue 1

Detailed scope

Intense Pulsed Light (IPL) Medical device for dermatologic use (Acne, Hirsutism, Hypertrichosis, Rosacea, Lentigo and Melasma): Anthelia NG med, Anthelia G+, Anthelia LCD, Anthelia LCD G+, Ariane, Ariane G+, Ariane LCD, Ariane LCD G+, Galaxy, Galaxy G+, Galaxy LCD, Galaxy LCD G+, ADENA, ADENA-LCD

Intense Pulsed Light (IPL) Medical device for dermatologic use (hirsutism and hypertrichosis): FLUENCE

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.