EC Certificate Full Quality Assurance System: Certificate FR11/00245

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

**Intense Pulsed Light (IPL) Medical device for dermatologic use (acne, hirsutism and hypertrichosis): ANTHELIA NG Med, 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.**

This certificate is valid from 26 June 2017 until 06 February 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 07 January 2020
Issue 10. Certified since 07 February 2011

Certification is based on reports numbered FR/MD 216647

Authorised by

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