



EC CERTIFICATE

Certificate No 1698/MDD

Full Quality Assurance System Approval Certificate

On the basis of our examination carried out according to Annex II, excluding section 4, of the Directive 93/42/EEC and its revised version, we hereby certify that:

GMV SRL

00173 ROMA (RM) - VIA ROBERTO PARIBENI 37 (ITA) - Italy

manages in the factory of:

00173 ROMA (RM) - VIA ROBERTO PARIBENI 37 (ITA) - Italy

a quality assurance system ensuring the conformity of the following products:

Dermatological and plastic microsurgery equipment

Type ref. Vibrance; Plexr; O.F.F; Needle Shaping, PLEXR PLUS; PLEXR PRO; PLEXR VECRON; VECRON.
Trade mark GMV

Carbon dioxide therapy equipment

Type ref. CARBOMAX.
Trade mark GMV

Electroporation transdermal equipment

Type ref. VEIKOS LED.
Trade mark GMV

Cavitation ultrasound therapy equipment

Type ref. SONIK.
Trade mark GMV

Radiofrequency therapy equipment

Type ref. RF PLUS MED; RF POWER MED; THUZZLE.
Trade mark GMV

with the relevant essential requirements of the aforementioned directive (from design to final inspection and testing) and it is subject to surveillance as specified in section 5 of Annex II. For class III devices, this certificate is valid only with the relevant EC Design-Examination Certificate of Annex II.4.

Date: 2014-07-11
Updated: 2020-03-25
Substitution Date: 2019-07-10
Expiry Date: 2024-05-26

IMQ



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Reference to IMQ files Nos:

10AO00104; DM16-0001908-01; DM16-0003279-01; DM16-0005880-01; DM17-0011992-01; DM18-0026549-01;
DM19-0035566-01; DM19-0044983-01.

This Approval Certificate is issued by IMQ S.p.A. as Notified Body for the Directive 93/42/EEC and its revised version. Notified Body notified to European Commission under number: 0051.

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