



EC CERTIFICATE

Certificate No 1349/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:

AB MEDICA GROUP, S.A.

08035 BARCELONA - C/JERICO, 10 BIS INTERIOR (ESP) - Spain

manages in the factory of:

08210 BARBERÀ DEL VALLÈS (BARCELONA) - AVDA SALVATELLA, 4, P.I. SALVATELLA (ESP) - Spain

SANTIGA 08210 BARBERÀ DEL VALLÈS (BARCELONA) - C/ LLOBATERAS 14-18, TALLERES 7 NAVE 4 y 6
P.IND. (ESP) - Spain

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

Recording charts for medical equipment

Trade mark LESSA

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.

Reference to IMQ files Nos:

10AK00185; 10EO00024; 10EO00041; DM15E0397325-01; DM17-0020237-01; DM18-0024386.

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.

Date: 2010-06-29
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IMQ