

## **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 BARBERÁ DEL VALLÈS (BARCELONA) - AVDA SALVATELLA, 4, P.I. SALVATELLA (ESP) - Spain

SANTIGA 08210 BARBERÁ 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: Updated:	2010-06-29 2018-07-02		
Substitution Date:	2015-06-17 —		
Expiry Date:	2023-07-01	IMQ	
	cate is subjected to the provis	n of IMQ S.p.A.   I-20138 Milano	

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This is a translation of the Italian text, which prevails in case of doubts