EC CERTIFICATE

Certificate No 251/MDD

Production Quality Assurance System Approval Certificate

On the basis of our assessment carried out according to Annex V, section 3 and considering the Annex VII, section 5 of the Directive 93/42/EEC and its revised version, we hereby certify that:

LP ITALIANA SPA

20157 MILANO (MI) - VIA CARLO REALE 15/4 (ITA) - Italy

manages in the factories of:

20157 MILANO (MI) - VIA CARLO REALE 15/4 (ITA) - Italy

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

Sterile swabs
Type ref. CULTIPLAST

with the relevant essential requirements of the aforementioned directive (concerning the manufacturing stage relevant to the reaching and the keeping of the sterile conditions) and it is subject to surveillance as specified in section 4 of Annex V.

Reference to IMQ files Nos:
10A9900089; 10A9900090; 10AD00165; DM15E0387619-01; DM18-0023718-01.

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.

Date:	1999-10-04
Updated:	2018-05-14
Substitution Date:	2015-04-09
Expiry Date:	2020-04-08

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IMQ ISTITUTO ITALIANO DEL MARCHIO DI QUALITA'

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This Approval Certificate is subjected to the provisions laid down in the "Rules for managing the EC Certification of Medical Devices on the basis of the Directive 93/42/EEC".

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