

EC CERTIFICATE

Certificate No 1170/MDD

Full Quality Assurance System Approval Certificate

On the basis of our examination carried out according to Annex II, section 3 of Legislative Decree of 1997-02-24, No 46, national transposition of the Directive 93/42/EEC, we hereby certify that:

TEKNO-LIT SPA

manages in the factories of:

CASTENEDOLO (BS) - VIA MARTORELLO 1 (ITA) - Italy

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

Medical supply units

Series: Sistema H Type ref. miniblock; monoblock; trackblock.
Trade mark Tekno-lit S.p.A.

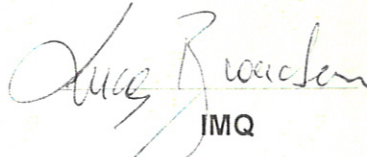
Series: Terapia Intensiva Type ref. singolo parete; singolo sospeso, doppio sospeso; singolo sospeso con interasse variabile.
Trade mark Tekno-lit S.p.A.

with the relevant essential requirements of the aforementioned national legislation transposing the Directive 93/42/EEC, from design to final inspection and testing.

Reference to IMQ files Nos: 10AI00205

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

Date: 2008-10-22


IMQ

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".
In any case, it does not remain valid after 2013-10-21 (article 11, clause 11 of the Directive).

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