



Ministero della Salute
**DIPARTIMENTO DELL'INNOVAZIONE- DIREZ. GENERALE DEI FARMACI E
 DISPOSITIVI MEDICI- UFFICIO III**

DGFDM/III/P 12478

HAVING REGARD to the 93/42/EEC Directive concerning medical devices;
HAVING REGARD to the Legislative Decree (D.L.vo) n. 46/97 (and its following amendments) reporting the accomplishment of 93/42 EEC Directive;
HAVING REGARD to request dated March 2nd, 2006 submitted by the Company RIMSA P. LONGONI S.r.l. located in 20038 Seregno (MI) – Via Monterosa, 18 - VAT. N. 00876160961;
WHEREAS this Company paid the fees required by Ministerial Decree (D.M.) May 24, 2004;
HAVING REGARD to the official deeds;

IT IS ATTESTED

that the Company RIMSA P. LONGONI S.r.l. located in 20038 Seregno (MI) – Via Monterosa, 18 - VAT. N. 00876160961, is manufacturer and has marked CE as medical devices, according to the Directive 93/42/EC, the following products:

OPERATING THEATRE LAMPS FOR MAJOR SURGERY:
 models D/900, D600, D/1500, D/1300, D/1200, D/1000

OT LAMPS FOR MINOR SURGERY
 models D/400, D/400x2, D/400x3, TRIS, SATURNO

EXAMINATION LAMPS
 models A/50, A/30, L/20, A/06, A/20, ALFA, 188 with magnifying glass, Wood-M, Wood-B, Shark-M, Shark-B

The above mentioned products, according to the art. 4 of the above mentioned Directive, can freely circulate and can be placed on the market in Italy and in all the European Union. This certificate is issued on the interested party's request in according to the law and for export in extracomunitary countries.

30 MAR 2006

The Office Manager
 Dott. Giuseppe Ruocco