MJS GROUP

DESIGN AND MANUFACTURE OF MEDICAL EQUIPMENT

EC DECLARATION OF CONFORMITY

Declares that the medical devices described hereafter:-

Hydropress 100 Hydropress 300
Hydropress 600 Hydropress 1200

Pulsepress Mini 3 Pulsepress Multi 3 Pro Pulsepress Multi 6
Pulsepress Multi 12 Pulsepress Multi 12 Auto Pulsepress Multi 12 Pro

Pulsepress Phyio 3 Pulsepress Physio 3 Pro Pulsepress Physio 6
Pulsepress Physio 12 Pro Pulsepress Multi 12 Lymph Pro
Pulsepress Venous 3

Pulsepress SLK Multi 3 Pulsepress SLK Multi 3 Pro
Pulsepress SLK Multi 6 Pro Pulsepress SLK Multi 12
Pulsepress SLK Multi 12 Auto Pulsepress SLK Vari-Lymph 12 Pro

Have been classified as Class IIa (Annex IX Rule 9) and is in conformity with the essential requirements and provisions of Council Directive 93 / 42 / EEC.

Is subject to the procedure set out in Annex II excluding section 4 of Directive 93/42/EEC under the supervision of notified body number 0120, SGS United Kingdom Limited, 202B Worle Parkway, Weston-Super-Mare, BS22 6WA

Barton Le Clay **27/07/12**

Chris Saggers
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