

EC Certificate

Full Quality Assurance System Directive 93/42/EEC on Medical Devices, Annex II excluding (4)

Registration No.: HD 1502429-1

Manufacturer: JUKA Spółka z ograniczoną odpowiedzialnością, Spółka komandytowa
ul. Fabryczna 4
32-005 Niepołomice
Poland

Products: -Partial Body Cryotherapy Chambers
-Whole Body Cryotherapy Chambers

Replaces EC Certificate, Registration No.: HD 60140975 0001

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II section 4 is required.

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Maciej Sciera

Maciej Sciera
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.