TÜVRheinland®

EC Certificate

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

Registration No.: HD 60140975 0001

Report No.: 26300295 009

Manufacturer:
JUKA Spolka z ograniczona
odpowiedzialnoscia,
Spolka komandytowa
ul. Fabryczna 4
32-005 Niepolomice
Poland

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

Replaces Certificate, Registration No.: ED 60114748 0001

Expiry Date: 2020-10-05

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.

Effective Date: 2019-09-04

Date: 2019-09-04

Notified Body

Jaroslaw Pyclik

TÜV Rheinland LGA Products GmbH - Zertifizierstelle

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.