TÜVRheinland®

EC Certificate

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

Registration No.: HD 60103674 0001

Report No.: 26300300 001

Manufacturer:
ASCOR MED Sp. z o.o.
Al. KEN 18 lok. 3B
02-797 Warszawa
Poland

Products:
- Syringe infusion pumps
- Volumetric infusion pumps

(see attachment for sites included)

Replaces EC Certificate, Registration No.: HD 60092919 0001

Expiry Date: 2020-07-14

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: 2015-08-18

Date: 2015-08-18

Notified Body

Sebastian Mniszek

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

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.