EC Certificate

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

Registration No.: HD 60133236 0001

Report No.: 21202387 013

Manufacturer:
NAWA Heilmittel GmbH
Ostendstr. 100
90482 Nürnberg
Deutschland

Products:
Products for wound healing and wound dressing

(see attachment for products included)

Replaces EC Certificate, Registration No.: HD 60089627 0001

Expiry Date: 2023-10-23

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: 2018-10-24

Date: 2018-10-11

Notified Body

Dr. K. Kluge

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.