


**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

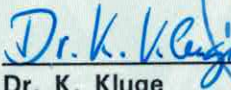
**Expiry Date:** 2023-10-23

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

Notified Body

**Date:** 2018-10-11

  
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.