

## **EG Declaration of Conformity**

### **According to Annex II of Directive 93/42/EEC for Medical Devices**

We herewith declare in sole responsibility that the following medical devices meet the essential requirements of Annex II of Directive 93/42/EEC and are appropriate for application according to this regulations:

Product specification: Blood lancets of sterile condition, Class II A  
Allergy lancets of sterile condition, Class II A  
Name of product (trade mark): **Medilance, product key 100**  
**Medilance safety, product key 1000**  
other products see HAB 18/1.1

- The manufacture takes place according to the implemented quality system fulfilling the essential requirements of DIN EN ISO 13485:2016+Ber. 1:2017-07 and Annex II of Directive 93/42/EEC.
- The Notified Body performed an audit of our quality system.
- The application of our quality system is subject to surveillance by the Notified Body.

#### **Notified Body:**

DQS Medizinprodukte GmbH  
August-Schanz-Straße 21  
D-60433 Frankfurt am Main



Period of validity acc. current certification Annex II - without section 4 of the directive 93/42/EEC suggestion of medical devices is:

26.05.2024

J. N. Eberle Federnfabrik GmbH  
Jürgen Brielmaier  
- director -

J. N. Eberle Federnfabrik GmbH  
Niels Weide  
-director-