EC- Declaration of conformity

Ref.
40366010 Blood lancets disposable sterile, central point, 200 pcs/box

J. N. Eberle Federnfabrik GmbH
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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 13485:2012-11 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 GmbH Deutsche Medizinprodukte GmbH
August-Schanz-Straße 21
D-60433 Frankfurt am Main

CE 0297

Period of validity acc. current certification Annex II - without section 4 of the directive 93/42/EEC suggestion of medical devices is:
23.10.2020

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

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

Schwabmünchen, 23.10.2015