

## Declaration of Conformity Cutimed Siltec (Re-Design)

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We.

#### **BSN** medical GmbH

# Quickbornstrasse 24 20253 Hamburg

hereby declare under our own responsibility, that the above mentioned product family

#### **CE-class IIb**

### containing the products:

```
73285-00000-00 CMED SLC STE 5X6CM 10 INT.
73285-00001-00 CMED SLC STE 10X10CM 10 INT.
73285-00002-00 CMED SLC STE 10X20CM 10 INT.
73285-00003-00 CMED SLC STE 15X15CM 10 INT.
73285-00004-00 CMED SLC STE 20X20CM 5 INT.
73285-00005-00 CMED SLC STE 5X6CM 12 DE
73285-00006-00 CMED SLC STE 10X10CM 12 DE
73285-00007-00 CMED SLC STE 10X20CM 12 DE
73285-00008-00 CMED SLC STE 15X15CM 12 DE
73285-00009-00 CMED SLC STE 20X20CM 6 DE
```

comply with the applicable regulations of the Medical Device Directive 93/42/EEC, including change directive 2007/47/EC, and that the products fulfil the essential requirements as defined in Annex I.

The Declaration of Conformity is performed in compliance with the Quality Management System according to EN ISO 13485 and in compliance with the Quality Assurance System according to MDD 93/42/EEC Annex II.

Date of Declaration: 04.11.2016

Valid until:

See expiry date of the attached EN ISO 13485 Quality

Management Certificate and the Quality Assurance Certificate

according to MDD 93/42/EEC Annex II.

This Declaration of Conformity is only valid in conjunction with the current certificate(s) issued by 0124 DEKRA Certification GmbH, Handwerkstrasse 15, D-70565 Stuttgart, Germany.

Compiled and released

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