

Konformitätserklärung Declaration of Conformity

Wir

We

**B. Braun Melsungen AG
Carl-Braun-Straße 1
34212 Melsungen
Deutschland/Germany**

erklären in eigener Verantwortung,
dass das/die Produkt/e

hereby declare in our own responsibility
that the product/s

Original Perfusor® Line

Original Perfusor® Line

Verlängerungsschlauch nur für enterale
Ernährung

Extension line for enteral nutrition only

(Artikelnummern siehe Anlage I)

(article numbers see attachment I)

mit den Anforderungen der folgenden Richtlinie
übereinstimmt/übereinstimmen

is/are in compliance with the following directive

Richtlinie 93/42/EWG des Rates vom 14. Juni
1993
über Medizinprodukte
geändert durch Richtlinie 2007/47/EG

Council Directive 93/42/EEC of 14th June
1993
concerning Medical Devices
amended by Directive 2007/47/EC

Konformitätsbewertungsverfahren
nach Anhang II (ausgenommen Abschnitt 4)

Conformity Assessment Procedure
according to annex II (excluding section 4)

Klassifizierung
gemäß Anhang IX der
oben genannten Richtlinie
Klasse IIa

Classification
according to annex IX of the
Council Directive named above
Class IIa

Benannte Stelle
TÜV SÜD Product Service GmbH
Ridlerstraße 65
80339 München
Deutschland
Kennnummer 0123

Notified Body
TÜV SÜD Product Service GmbH
Ridlerstraße 65
80339 München
Germany
Identification number 0123

Datum der ersten CE-Kennzeichnung
2013-07

Date of first CE-marking
2013-07

Gültig bis
2024-05-26

Valid until
2024-05-26

Anlage I / Attachment I

Art.-Nr. / Art. No.	Produktname / Product name	Klasse / Class
8722930	Original Perfusor® Line	Ila
87229910	Original Perfusor® Line	Ila

Amendment Information

Version	Description of the changes
04	Add art. no. 87229910 Change Address of the Notified Body
05	Update validity

Title: Declaration of Conformity - 121-002 - Original Perfusor Line (Enteral) Initiator: Caroline ? Herbst

This document is signed electronically in compliance with the B. Braun electronic signature policies and procedures by following persons:

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