



Full Quality Assurance

No. CE 74883

Issued to:

**Gambro Kathetertechnik Hechingen
Zweigniederlassung Gambro
Dialysatoren GmbH
Linsenaeker 1
72379 Hechingen
Germany**

In respect of:

The design, development and manufacture of Vascular Access Devices and accessories for use in renal replacement treatment

on the basis of our examination under the requirements of Council Directive 93/42/EEC, Annex II, Section 3.2.

For and on behalf of the British Standards Institution, a Notified Body for the above Directive (Notified Body Number 0086):

Alastair Trivett, Managing Director, BSI Product Services – Global

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Conditions of Approval

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

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