

Directive 93/42/EEC on Medical Devices (MDD), Annex II, excluding (4),
(Devices in Class IIa)

Corp. GmbH (Europe)

Eiffelstr. 80

Product

The Certification Body TÜV SÜD Product Service GmbH declares that the aforementioned

conforms in accordance with MDD Annex II

quality assurance

is mandatory. See also notes overleaf.

Report No. CU446420

Valid until: 2018-12-18



Date: 2012-1

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Product Service

CERTIFICATE

DO

EU Certificate

Directive 93/42/EEC on Medical Devices (MDD), Annex I (Devices in Class IIa, IIb or III)

No. G1 11 11 75606 004

IF

Facility(ies).

Purecan Medical (Shanghai) Co., Ltd.
24-6 Building

ΦΙΚΑΤ

CERTIFICATE

認證證書

CERTIFICATE

ZERTIF