



EC-CERTIFICATE

(Production quality assurance)

DQS Medizinprodukte GmbH

hereby certifies that the company

Remy & Geiser GmbH

Unternehmensgruppe für pharmazeutische Verpackungsmittel

Remy-Geiser-Straße 1
56584 Anhausen/Ww.
Germany

has implemented and maintains a quality assurance system which applies to the products with respect to the aspects of manufacture concerned with securing and maintaining sterile conditions.

An audit, documented in a report, performed by DQS, has verified that this quality assurance system fulfils the requirements of

Annex V of Council Directive 93/42/EEC concerning medical devices

with respect to the following medical devices:

steril medical devices according to annex

The manufacturer is subject to surveillance according to Annex V, Section 4. The CE marking with the Notified Body Number (0297) may be affixed on the devices listed in the certificate.

Certificate registration no.	001417 MR5s
Certification ID	170516194
Date of certification	2011-01-31
Valid until	2015-03-14

Frank Graichen
Managing Director

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Head of Certification Body

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DQS Medizinprodukte GmbH is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with the Notified Body Number 0297.



Annex to Certificate
Registration No. 001417 MR5s
Certificate ID 170516194
(Issued: 2011-01-31)

Remy & Geiser GmbH

Unternehmensgruppe für pharmazeutische Verpackungsmittel

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Products Groups

Pipettes and pipette assemblies sterile
Eye dropper systems sterile

Products

-Pipette assembly (classic) sterile
-Eye dropper closure aggregate
-Eye dropper closure Zentrop
-Eye dropper closure Retro
-Eye dropper closure