



EC-CERTIFICATE

(Full quality assurance system)



This is to certify that the company



BHT Hygienetechnik GmbH

Division

Messerschmittstr. 11
86368 Gersthofen
Germany

has implemented and maintains a full quality assurance system which applies to the products at every stage from design to final controls.

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of

Annex II – excluding Section 4 of Council Directive 93/42/EEC concerning medical devices

with respect to the following medical devices:

Device family	Device	Class
Disinfection equipment	INNOVA Serie M2 / E2 / E2 CMS	IIb
	INNOVA Serie M2s / E2s / E2s CMS	
	INNOVA Serie M3 / E3 / E3CMS	
	INNOVA Serie M4 / E4 / E4CMS	
	INNOVA Serie M5 / E5 / E5CMS / E5CMS EFF	
	INNOVA Serie 3s	
	INNOVA Serie 4s	

The manufacturer is subject to surveillance according to Annex II, Section 5. The CE marking with the Notified Body Identification Number (0297) may be affixed on the devices listed in the certificate. An EC Design Examination Certificate according to Annex II, Section 4 is required for class III devices covered by this certificate. The certificate is in the case of class I(s) devices (I(s) = class I products placed on the market in sterile conditions) limited to the aspects of manufacture concerned with securing and maintaining sterile conditions. The certificate is in the case of class I(m) devices (I(m) = class I devices with a measuring function) limited to the aspects of manufacture concerned with the conformity of the products with the metrological requirements.

Certificate registration no. 019906 MR2
Certificate unique ID 170722750
Effective date 2018-09-22
Expiry date 2022-07-15
Frankfurt am Main 2018-09-22

DQS Medizinprodukte GmbH

Sigrid Uhlemann
Managing Director

Dr. Thomas Feldmann
Head of Certification Body

August-Schanz-Straße 21, 60433 Frankfurt am Main,
Tel. +49 (0) 69 95427-300, medical.devices@dqs-med.de



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

