







(Full quality assurance system)

This is to certify that the company

Unintech GmbH

Monschauer Str. 12 40549 Düsseldorf 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:

Rigid Endoscopes according to annex

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. 544503 MR2
Certificate unique ID 170771088
Effective date 2020-12-13
Expiry date 2024-05-26
Frankfurt am Main 2020-12-13

DQS Medizinprodukte GmbH

Sigrid Uhlemann Managing Director Dr. Thomas Feldmann Head of Certification Body

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Annex to certificate

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Device family	Device	Rule	UMDNS	Class
Rigid Endoscopes (UMDSN 11-274)	Arthroscope Zystoscope Hysteroscope Laparoscope Laryngoscope Otoscope Resektoscope Rhinoscope / Sinoscope Thorakoscope Bronchoscope	6 6 6 6 6 6 6 6	10-198 11-112 12-081 12-291 12-293 12-849 13-335 13-396 14-047 15-074	lla lla lla lla lla lla lla lla
	Nephroscope Ureterenoscope	6 6	15-591 15-788	lla lla

