



EC-CERTIFICATE



(Full quality assurance system)

This is to certify that the company

RZ Medizintechnik GmbH

Unter Hasslen 20 78532 Tuttlingen 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:

Endoscopic Systems for minimal-invasive surgery, Equipment and accessories for minimalinvasive surgery, Monopolar/Bipolar instruments and Electrodes for HF-surgery according 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.	286629 MR2
Certificate unique ID	170772556
Effective date	2020-11-13
Expiry date	2023-08-02
Frankfurt am Main	2020-11-13

DQS Medizinprodukte GmbH

Mb lu

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

Sigrid Uhlemann Managing Director

Dr. Thomas Feldmann Head of Certification Body

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





Annex to certificate Certificate registration No.: 286629 MR2 Certificate unique ID: 170772556 Effective date: 2020-11-14

RZ Medizintechnik GmbH

Unter Hasslen 20 78532 Tuttlingen Germany

Device family	Device	Class
Endoscopy and accessories	Arthroscopy System Laparoscopy System Hysteroscopy System Ureterorenoscopy System Nephroscopy System Cystoscopy System Discectomy System	lla Ila Ila Ila Ila Ila
Laparoscopy	Laparoscopy Instruments sterile/non sterile	llb
HF-surgery	Bipolar Instruments Bipolar Electrodes Monopolar Instruments Monopolar Electrodes sterile/non-sterile High frequency generators	llb llb llb llb llb
Suction and irrigation system	Cannulas Suction and irrigation instruments (HF) Suction and irrigation cannulas	lla Ila Ila
Endoscopic sheath and accessories	Trocar systems	lla
Resectoscopy	Resectoscopy System Monopolar Electrodes sterile/non-sterile	llb + lla llb
Equipment	Suction and irrigation pump and accessories Insufflator and accessories Shaver System	lla Ilb + Ila Ilb
Retractors	Self-Retaining Retractors	lla

