



## **EC-CERTIFICATE**



(Full quality assurance system)

This is to certify that the company

## Max Hauser Süddeutsche Chirurgiemechanik GmbH

Föhrenstrasse 33 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:

Retractor, Spreader and Endoscopes 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. 449956 MR2
Certificate unique ID 170758585
Effective date 2019-11-11
Expiry date 2024-05-26
Frankfurt am Main 2019-11-11

**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	Class
Retractor	Rip Retractor Sterned Retractor	lla lla
Spreader	Wound Spreader Vertabone Spreader Laminectory Spreader	lla Ila Ila
Endoscopes	Endoscopes with working channel	lla

