



# EC-CERTIFICATE

(Full quality assurance system)



This is to certify that the company

## erbrich-instrumente GmbH

Eugenstraße 33  
78532 Tuttlingen - Nendingen  
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:

Suction and rinsing cannulas as listed in the 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.	516070 MR2
Certificate unique ID	170776143
Effective date	2021-05-20
Expiry date	2023-09-09
Frankfurt am Main	2021-05-20

### DQS Medizinprodukte GmbH

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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**  
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<b>Device</b>	<b>GMDN Code</b>	<b>Class</b>
Needle, Biopsy	12-734	Ila
Cannulae, Aspirating	10-566	Ila
Cannulae, Ear	10-571	Ila
Cannulae, Eye	10-573	Ila
Cannulae, Uterine	10-580	Ila
Cannulae, Nasal	16-422	Ila
Cannulae, Liposuction	17-640	Ila