



## **EC-CERTIFICATE**



(Production quality assurance)

This is to certify that the company

## Medizin & Service GmbH

Boettcherstrasse 10 09117 Chemnitz Germany

has implemented and maintains a quality assurance system which applies to the manufacture and final controls of the products.

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

## Annex V of Council Directive 93/42/EEC concerning medical devices

with respect to the following medical devices:

Keredusy® (KR2000/KR1000) Disir	nfection system Class IIa
Eye Diagnostic Equipment EP 1000	) Class IIa

The manufacturer is subject to surveillance according to Annex V, Section 4. The CE marking with the Notified Body Identification Number (0297) may be affixed on the devices listed in the 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.	530682 MR5
Certificate unique ID	170770110
Effective date	2020-06-28
Expiry date	2024-05-26
Frankfurt am Main	2020-06-28

## DQS Medizinprodukte GmbH

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Sigrid Uhlemann Managing Director

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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.