



# EU Quality Management Certificate



This is to certify that the company

## FUJIFILM medwork GmbH

Medworkring 1  
91315 Höchststadt a.d. Aisch  
Germany

SRN: DE-MF-0000084444

has established, implemented and maintains a Quality Management System in accordance with

### **Annex IX, Chapter I and III of the Regulation (EU) 2017/745 Conformity Assessment based on a Quality Management System and on Assessment of Technical Documentation**

for the device categories and products listed in the Annex of this certificate.

The conformity of the Quality Management System has been verified in an audit and is subject to regular surveillance in accordance with Annex IX, Chapter 1, Section 3. Limitations to this certificate are listed in the Annex.

Devices listed in the Annex may bear the CE marking with the identification number of the Notified Body (0297).

For placing of devices of class III and devices class IIb implantable according to Article 52(4) subparagraph 2 listed in the Annex on the market, an additional certificate according to Annex IX, Chapter II is required.

Certificate registration no.	230521 MDR2017Q
Certificate ID	1000204042
Effective date	2025-03-20
Expiry date	2028-10-01
Frankfurt am Main,	2025-03-20



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
www.zlg.de  
BS-MDR-094

## DQS Medizinprodukte GmbH

Heinrich von Mettenheim  
Managing Director



Accredited Body: DQS Medizinprodukte GmbH, August-Schanz-Str. 21, 60433 Frankfurt am Main  
**DQS Medizinprodukte GmbH is a Notified Body according to Regulation (EU) 2017/745  
of the Council concerning medical devices with the Identification Number 0297.**  
The validity of the certification can only be verified by the QR-code.



**Annex to EU Quality Management Certificate**  
**SRN of Manufacturer: DE-MF-0000084444**  
**Certificate ID: 1000204042**

**Device categories and variants covered by this certificate:**

Device category: **MDN 1203 - Non-active non-implantable guide catheters, balloon catheters, guidewires, introducers, filters, and related tools**  
Product name: Stone extraction balloon  
Risk classification: Is  
Basic-UDI-DI: 4044503056113U  
Intended purpose: **Stone extraction balloons** are used for endoscopic extraction of stones and sludge from the bile and pancreatic duct and for injection of contrast medium into the bile and pancreatic duct.

Device category: **MDA 0312 - Other active non-implantable surgical devices**  
Product name: Polypectomy snare  
Polypectomy snare with injection needle  
Risk classification: IIb  
Basic-UDI-DI: 40445030564349  
Intended purpose: **Polypectomy snares** are used for endoscopic tissue resection in the upper and lower gastrointestinal tract.

Device category: **MDN 1202 - Non-active non-implantable devices for administration, channelling and removal of substances, including devices for dialysis**  
Product name: Application catheter  
Risk classification: Is  
Basic-UDI-DI: 404450300100ZQ  
Intended purpose: **Application catheters** are used to apply hemostatic agents like synthetic peptide material or aqueous peptide solutions in endoscopic procedures in the upper and lower gastrointestinal tract..

Device category: **MDN 1202 - Non-active non-implantable devices for administration, channelling and removal of substances, including devices for dialysis**  
Product name: ERCP-Catheter  
Risk classification: Is  
Basic-UDI-DI: 40445030561748  
Intended purpose: **ERCP catheters** are used for endoscopic cannulation of the papilla and application of contrast medium into the biliary, pancreatic and hepatic ducts.



**Annex to EU Quality Management Certificate**  
**SRN of Manufacturer: DE-MF-0000084444**  
**Certificate ID: 1000204042**

Device category: **MDN 1202 - Non-active non-implantable devices for administration, channelling and removal of substances, including devices for dialysis**

Product name: EUS-FNA needle  
EUS-FNB needle

Risk classification: IIa

Basic-UDI-DI: 4044503017813S

Intended purpose: The **EUS-FNA/FNB needles** are used with a curvilinear echoendoscope for fine needle aspiration and biopsy of submucosal and extramural gastrointestinal lesions or for delivery of injectable materials.

**Examinations and tests performed:**

230521\_A210021MED\_01 dated 2022-10-28  
230521\_A213086MED\_01 BAL1 Series dated 2023-09-16  
230521\_A210021MED\_01 POL1 Series dated 2024-05-18  
230521\_A214650MED Application catheter dated 2024-07-30  
230521\_A216053MED\_02 EUS-FNA needle / EUS-FNB needle dated 2025-03-11

**Further conditions for or limitations to the validity of the certificate:**

In case of products that are placed on the market in sterile condition, the involvement of the Notified Body in the conformity assessment procedure is limited to the aspects of manufacture concerned with securing and maintaining sterile condition.

**Reference to previous certificates:**

Revision	Date of Issue	Certificate-ID	Description of change
01	2023-10-02	170780024	Addition of Polypectomy Snares
02	2024-05-23	1000171787	Addition of Application catheter, Certificate correction of typo files and change of address
03	2024-08-08	1000191195	Addition of ERCP-Catheter
04	2024-08-29	1000195468	Addition of EUS-FNA needle / EUS-FNB needle