



EU Declaration of Conformity

Richard Wolf
Pforzheimer Straße 32
75438 Knittlingen
Germany

SRN: DE-MF-000007048

We declare under our sole responsibility that the device/s meet(s) the applicable requirements / provisions of the European Medical Device Regulation 2017/745 and European Directive 2011/65/EU - Restriction of the use of certain hazardous substances in electrical and electronic equipment, and that we are solely responsible for issuing this Declaration of Conformity.

Conformity assessment procedure according to:	Article 52 (7), sentence 1 of Regulation EU 2017/745
Common specifications:	N/A
Notified body: (European Medical Device Regulation 2017/745)	N/A
Risk class:	I
EU quality management system certificate:	N/A
Basic UDI DI:	405520724072019-0017775NA
Intended purpose:	The products serve to convert an image of the inside of the patient's body generated by an endoscope into an electronic signal which is fed to the camera control unit.

All products covered by this declaration are listed on following page(s).

Place and date of issue: Knittlingen, 10.12.2025

Florian Happe
Vice-President Research and Development

12.12.2025 | 15:12 MEZ

Date

Signiert von:

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Signature

Ute Greiner
Director Global Regulatory Affairs

12.12.2025 | 17:28 MEZ

Date

Signed by:

2042D603C48B42C...
Signature

Wulf Brunow
Vice President Global Quality Assurance and
Regulatory Affairs

12.12.2025 | 17:32 MEZ

Date

DocuSigned by:

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Signature

Dokument: 10000188481 / R00 / 000 / 05



Product list | Produktliste

Product Produkt	Product name Produktbezeichnung
5525933	LOGIC HD PENDUAL CAMERA HEAD CABLE 3M LOGIC HD PENDUAL KAMERAKOPF KABEL 3M
85525902	LOGIC HD CAMERA HEAD CABLE 3M LOGIC HD KAMERAKOPF KABEL 3M
85525922	LOGIC HD CAMERA HEAD CABLE 3M LOGIC HD KAMERAKOPF KABEL 3M
85525942	LOGIC 4K CAMERA HEAD LOGIC 4K KAMERAKOPF