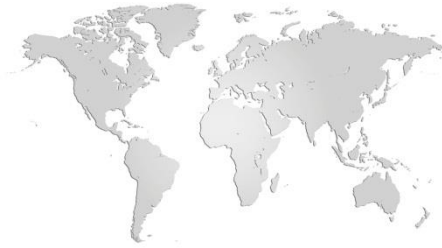


# CERTIFICATE



## EN ISO 13485:2016 + AC:2018 + A11:2021

DEKRA Certification GmbH hereby certifies that the organization

**Richard Wolf GmbH**

### Scope of certification:

Design and development, production, distribution, installation and service of systems, active medical devices (sterile, non-sterile), non-active medical devices (sterile, non-sterile) for human medicine, in particular for endoscopy and extracorporeal shockwave application.  
Design and development, production, and distribution of non-active implants in urology and surgery as well as accessories for processing (cleaning, disinfection, sterilization)

### Certified location:

Pforzheimer Str. 32, 75438 Knittlingen, Germany

(further locations see annex)

has established and maintains a quality management system according to the above mentioned standard. The conformity was adduced with audit report no. 50593-R2-00.

Certificate registration no.: 50593-21-00\_EN  
Validity of previous certificate: 2024-01-29

Certificate valid from: 2024-01-30  
Certificate valid to: 2024-11-28

Karin Leicht  
DEKRA Certification GmbH, Stuttgart, 2024-01-30



Deutsche  
Akkreditierungsstelle  
D-ZM-16029-08-00

# Annex to the Certificate No. 50593-21-00

valid from 2024-01-30 to 2024-11-28

The following locations/companies belong to the certificate above:

	Headquarters	Certified location	Scope of certification
	Richard Wolf GmbH	Pforzheimer Str. 32 75438 Knittlingen Germany	see page 1
	<b>at the following locations/at the companies at the following locations</b>		<b>Scopes of certification</b>
1.	Richard Wolf GmbH	Reuchlinstr. 10-11 10553 Berlin Germany	Manufacture of flexible and rigid endoscopes



*K. Leicht*

Karin Leicht  
DEKRA Certification GmbH, Stuttgart, 2024-01-30