EU Certificate

for the assessment of the quality management system

according to Medical Device Regulation (EU) 2017/745, Annex IX Chapter I

As a Notified Body of the European Union, DEKRA Certification GmbH certifies, that the manufacturer

Richard Wolf GmbH
Single Registration Number (SRN): ---
Pforzheimer Straße 32, 75438 Knittlingen, Germany

applies a quality management system according to Annex IX Chapter I of the Medical Device Regulation (EU) 2017/745 for the medical devices listed in the annex. The approval is based on the result of the (re-)certification audit, report no. 50593-R1-00, the decision dated 2020-12-14 and is only valid in connection with the successful performance of the annual surveillance audits.

EU Certificate no.: 50593-60-00

Certificate valid from: 2020-12-14
Certificate valid to: 2025-12-13

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
BS-MDR-092

Natascha Jezyschek
DEKRA Certification GmbH Stuttgart; 2020-12-14
Notified Body ID number: 0124

Annex to the EU Certificate no. 50593-60-00

Valid from 2020-12-14 to 2025-12-13

Revision status of the annex: 0 dated 2020-12-14

Following devices/device categories are included in this certificate:

Class IIa

MDA0204

- disposable flexible endoscope

Natascha Jezyschek
DEKRA Certification GmbH Stuttgart; 2020-12-14
Notified Body ID-number: 0124