



## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

### Manufacturer:

**AESCULAP AG**

Am Aesculap-Platz  
78532 Tuttlingen  
GERMANY

SRN Manufacturer - DE-MF-000005504

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has established, documented and implemented a quality management system as described in Article 10 (9) of the Regulation (EU) 2017/745 on medical devices. Details on device categories covered by the quality management system are described on the following page(s). The Report referenced below summarises the result of the assessment and includes reference to relevant CS, harmonized standards and test reports. The conformity assessment has been carried out according to Annex IX Chapter I and III of this regulation with a positive result.

The quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis.

The certified quality management system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. The surveillance assessment shall also include an assessment of the technical documentation for the device or devices concerned on the basis of further representative samples. All applicable requirements of the Testing, Certification, Validation and Verification Regulations TÜV SÜD Group have to be complied with.

For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G10 010066 0438 Rev. 05](http://www.tuvsud.com/ps-cert?q=cert:G10_010066_0438_Rev_05)

**Report No.:** 713347039

**Preceding Certificate No.:** G10 010066 0438 Rev. 04

**Valid from:** 2025-07-10

**Valid until:** 2030-07-09

**Date of Initial Issuance:** 2020-07-10

Christoph Dicks  
Head of Certification/Notified Body

**Issue date:** 2025-05-16



## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

**Classification:** Class IIa  
**Device Group:** L030101 - SUCTION AND IRRIGATION SURGICAL CANNULAS AND HANDPIECES, REUSABLE

**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** L031309 - SUTURE NEEDLE PASSERS, REUSABLE  
**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** L031401 - GENERAL SURGERY SPREADERS AND RETRACTORS, REUSABLE  
**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** L040901 - ABDOMINAL SPREADERS, REUSABLE  
**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** L060502 - NON-ENDOSCOPIC UROLOGY SPREADERS, REUSABLE  
**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** L070702 - CARDIAC DILATORS AND RETRACTORS, REUSABLE  
**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** L080602 - THORACIC SURGERY SPREADERS, REUSABLE  
**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** L110501 - VERTEBRAL SURGERY SPREADERS AND RETRACTORS, REUSABLE  
**Intended Purpose:** -



## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

<b>Classification:</b>	Class IIa
<b>Device Group:</b>	L110503 - CRANIAL SURGERY SPREADERS AND RETRACTORS, REUSABLE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	L149003 - ENT RETRACTORS, REUSABLE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	L031201 - THORACIC TROCAR, REUSABLE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	L031202 - ABDOMINAL TROCAR, REUSABLE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	L031280 - SURGICAL TROCAR, REUSABLE - ACCESSORIES
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A019001 - BLUNT NEEDLES
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	A070199 - ADAPTERS AND CONNECTORS - OTHER
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	C019019 - VESSEL STRIPPER SYSTEMS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	G020401 - HAEMORRHOID LIGATURE SETS
<b>Intended Purpose:</b>	-



## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

<b>Classification:</b>	Class IIa
<b>Device Group:</b>	H030102 - SINGULAR CLIPS FOR OPEN SURGERY
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	H030201 - MULTIPLE CLIP APPLIERS FOR VIDEOSURGERY
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	H030201 - MULTIPLE CLIP APPLIERS FOR VIDEOSURGERY
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	K010101 - TROCAR, SINGLE-USE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	K010201 - MINIMALLY INVASIVE SURGERY SURGICAL INSTRUMENTS, SINGLE-USE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	K010201 - MINIMALLY INVASIVE SURGERY SURGICAL INSTRUMENTS, SINGLE-USE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	K0104 - VERESS NEEDLES
<b>Intended Purpose:</b>	-



## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

<b>Classification:</b>	Class IIb
<b>Device Group:</b>	K020101 - MONO- AND BIPOLAR SURGICAL INSTRUMENTS, SINGLE-USE
<b>Intended Purpose:</b>	<p>Depending on the design of the working end, the instruments are used for cutting, dissecting, grasping and suturing tissues, as well as for biopsies and/ or for thermal tissue treatment during minimal-invasive procedures.</p> <p>Bipolar forceps are used for hemostatic coagulation as well as grasping and dissecting of tissue in surgical procedures.</p> <p>The modular monopolar electrodes are used to prepare, cut, dissect, and coagulate tissue are used in minimal invasive procedures by means of monopolar HF current.</p> <p>The single-use monopolar instrument is connected to an appropriate HF generator to cut or coagulate tissue in open surgical procedures.</p> <p>The suction/irrigation devices are used for the suction of body fluids, blood, tissue or secretions, for irrigation with rinsing solutions and for coagulation, cutting and dissecting tissue.</p>
<b>Classification:</b>	Class IIb
<b>Device Group:</b>	K020102 - ELECTROSURGERY PADS (NEUTRAL ELECTRODES) AND CABLES, SINGLE-USE
<b>Intended Purpose:</b>	The neutral electrodes are used in the monopolar HF technique, where they serve to pick up the HF current from the wider area of operation on the patient's body and conduct it back to the HF device.
<b>Classification:</b>	Class IIb
<b>Device Group:</b>	K020301 - RADIOFREQUENCY SURGERY INSTRUMENTS, SINGLE-USE
<b>Intended Purpose:</b>	Caiman instruments are bipolar HF seal and cut instruments. They are used for grasping, dissecting, sealing and cutting of soft tissue and vessels during open and / or laparoscopic surgical procedures.
<b>Classification:</b>	Class IIb
<b>Device Group:</b>	L180201 - OPEN ELECTROSURGERY SCISSORS, REUSABLE
<b>Intended Purpose:</b>	Bipolar scissors are used for cutting, dissecting and coagulating tissues in surgical operations.



## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

**Classification:** Class IIb  
**Device Group:** L180202 - ENDOSCOPIC ELECTROSURGERY SCISSORS, REUSABLE  
**Intended Purpose:** Depending on the design of the working end, the instruments are used for cutting, dissecting, grasping and suturing tissues, as well as for biopsies and/ or for thermal tissue treatment during minimal-invasive procedures.

**Classification:** Class IIb  
**Device Group:** L180301 - OPEN ELECTROSURGERY HANDPIECES, REUSABLE  
**Intended Purpose:** The reusable monopolar electrode handles are used in open surgical procedures to conduct the current from the HF generator to the operating site, to hold the required working electrode, and if applicable to activate the HF generator.

**Classification:** Class IIb  
**Device Group:** L180302 - ENDOSCOPIC ELECTROSURGERY HANDPIECES, REUSABLE  
**Intended Purpose:** Depending on the design of the working end, the instruments are used for cutting, dissecting, grasping and suturing tissues, as well as for biopsies and/ or for thermal tissue treatment during minimal-invasive procedures.  
  
The modular monopolar electrodes are used to prepare, cut, dissect, and coagulate tissue are used in minimal invasive procedures by means of monopolar HF current.

**Classification:** Class IIb  
**Device Group:** L180401 - OPEN ELECTROSURGERY FORCEPS, REUSABLE  
**Intended Purpose:** Bipolar forceps are used for hemostatic coagulation as well as grasping and dissecting of tissue in surgical procedures.  
  
The HF monopolar forceps are connected via a monopolar active electrode and monopolar handle with a HF generator to be used to dissect, grasp, hold, and coagulate tissue in general surgical procedures.

**Classification:** Class IIb  
**Device Group:** L180402 - ENDOSCOPIC ELECTROSURGERY FORCEPS, REUSABLE  
**Intended Purpose:** Depending on the design of the working end, the instruments are used for cutting, dissecting, grasping and suturing tissues, as well as for biopsies and/ or for thermal tissue treatment during minimal-invasive procedures.





## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

<b>Classification:</b>	Class IIb
<b>Device Group:</b>	L180601 - OPEN ELECTROSURGERY ELECTRODES, REUSABLE
<b>Intended Purpose:</b>	The monopolar HF electrodes are used in combination with appropriate handles and generators to prepare, cut, dissect, and coagulate tissue by means of monopolar HF current in open surgery.
<b>Classification:</b>	Class IIb
<b>Device Group:</b>	L180602 - ENDOSCOPIC ELECTROSURGERY ELECTRODES, REUSABLE
<b>Intended Purpose:</b>	The modular monopolar electrodes are used to prepare, cut, dissect, and coagulate tissue are used in minimal invasive procedures by means of monopolar HF current.
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	Q019001 - SALIVA ASPIRATORS AND SALIVA ABSORBENTS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	T030199 - COVERS, INSTRUMENTS AND EQUIPMENT - OTHER
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	V010101 - SCALPELS WITH SAFETY SYSTEMS, SINGLE-USE
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	V010302 - BLADES WITHOUT SAFETY SYSTEMS, SINGLE-USE - NOT INCLUDED IN OTHER CLASSES
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	V0199 - CUTTING DEVICES, SINGLE-USE - OTHER
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	Z120103 - DERMOTOMY EQUIPMENT
<b>Intended Purpose:</b>	-



## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

<b>Classification:</b>	Class IIb
<b>Device Group:</b>	Z120109 - ELECTROSURGICAL INSTRUMENTS
<b>Intended Purpose:</b>	The foot switch is used for activating compatible devices for HF surgery.  The bipolar HF generator is used for coagulation with bipolar instruments.  The HF generator is used for sealing and cutting of vessels with compatible seal and cut instruments.
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	Z120190 - VARIOUS INSTRUMENTS FOR GENERAL AND MULTIDISCIPLINARY SURGERY
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	Z120190 - VARIOUS INSTRUMENTS FOR GENERAL AND MULTIDISCIPLINARY SURGERY
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	Z120190 - VARIOUS INSTRUMENTS FOR GENERAL AND MULTIDISCIPLINARY SURGERY
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	Z120114 - SURGICAL NAVIGATION INSTRUMENTS
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	Z12011482 - SURGICAL NAVIGATION INSTRUMENTS - SOFTWARE ACCESSORIES
<b>Intended Purpose:</b>	-
<b>Classification:</b>	Class IIa
<b>Device Group:</b>	Z120204 - INSTRUMENTS FOR THE ACQUISITION AND MANAGEMENT OF ENDOSCOPIC AND MINIMALLY INVASIVE SURGERY IMAGES
<b>Intended Purpose:</b>	-





## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

**Classification:** Class IIa  
**Device Group:** Z120290 - VARIOUS INSTRUMENTS FOR ENDOSCOPY AND MINI-INVASIVE SURGERY

**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** Z120290 - VARIOUS INSTRUMENTS FOR ENDOSCOPY AND MINI-INVASIVE SURGERY

**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** Z120590 - VARIOUS INSTRUMENTS FOR CARDIOLOGY AND CARDIAC SURGERY

**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** Z121305 - MOTORISED ORTHOPAEDIC SURGERY SYSTEM INSTRUMENTS

**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** Z121305 - MOTORISED ORTHOPAEDIC SURGERY SYSTEM INSTRUMENTS

**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** Z121009 - INSTRUMENTS FOR MOTORISED NEUROSURGERY SYSTEMS

**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** Z121009 - INSTRUMENTS FOR MOTORISED NEUROSURGERY SYSTEMS

**Intended Purpose:** -

**Classification:** Class IIa  
**Device Group:** Z121009 - INSTRUMENTS FOR MOTORISED NEUROSURGERY SYSTEMS

**Intended Purpose:** -

**The validity of this certificate** . /.

Page 9 of 10

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany



## EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III  
(Class IIa and Class IIb Devices)

**No. G10 010066 0438 Rev. 05**

**depends on conditions and/or  
is limited to the following:**

### Revision History:

Rev.	Dated	Report	Description
00	2020-07-10	713175266	-
01	2021-12-09	713203407 / 713203404 / 713203403 / 713203400 / 713203397 / 713203393 / 713203388 / 713205439 / 713229575	-
02	2022-11-08	713203406 / 713205438 / 713218837 / 713218822	-
03	2022-11-17	713203406 / 713205438 / 713218837 / 713218822	-
04	2023-11-07	713218567 / 713218653 / 713218808 / 713230390 / 713303316	Supplemented: Device(s)/group of device(s) added
05	2025-07-10	713347039	Renewal of certificate