

Certificate

Quality Assurance System acc. to Directive 2014/68/EU

Certificate no.: 01 202 644/Q-18 B008

Name and address of the
certificate holder: **MELAG Medizintechnik GmbH & Co. KG**
Geneststr. 6-10
10829 Berlin
Germany

Herewith we certify that the above -mentioned manufacturer operates a quality system according to the European Directive 2014/68/EU.
The manufacturer has the permission to affix the following CE marking to pressure equipment described and manufactured in accordance to the scope covered by this Quality-Assurance System:

CE 0035

Test basis: **Directive 2014/68/EU: QA-System (Module D)**
(the QS-Modules E1, E, D1 are covered by Module D)

Audit report no.: 01 202 644/Q-18 B008

Scope: **Production of sterilizers for medical purposes and safety valves, see annex (Rev.:6, 2022-05-12) to certificate**

Manufacturing plant: see certificate holder

Validity: **This certificate is valid until 2024-05-31..**

Cologne, 2022-12-12

Dipl.-Ing. (FH) Vera Ruff



TÜV Rheinland Industrie Service GmbH
Notified Body for Pressure Equipment, ID-No. 0035
Am Grauen Stein, D-51105 Cologne

MS-0037317 E-008-Rev01