

# 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

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