

Certificate

Quality-Assurance System

acc. to Directive 2014/68/EU

Certificate no.:

01 202 644/Q-18 B008

Name and address of the
manufacturer:

MELAG Medizintechnik GmbH & Co. KG
Geneststr. 6-10
D - 10829 Berlin

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

Tested acc. to
Directive 2014/68/EU:

QS-System (Module D)
(the QS-Modules E1, E and D1 are covered by Module D)

Audit report no.:

644/Q-18 B008

Area of validity:

Production of sterilizers for medical purposes
and safety valves, see annex to certificate

Manufacturing plant:

MELAG Medizintechnik GmbH & Co. KG
Geneststr. 6-10
D - 10829 Berlin

Valid until:

May 31, 2024

Cologne, July 6, 2021

Dipl.-Ing. (FH) Vera Ruff

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



E-008-E-Rev22