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 svety valves, see annex to certificate

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

Valid until: May 31, 2021

Cologne, January 21, 2020

Dipl.-Ing. (FH) Roman Alexander Meyers

TÜVRheinland® 0035 Notified Body

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

E-008-E-Rev22

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