

EC Certificate of Conformity

In accordance with the requirements of the Pressure Equipment Directive 97/23/EC and the Pressure Equipment Regulations 1999, UK Statutory Instrument 1999 No. 2001 as amended by S.I. 2002 No. 1267 and S.I. 2015 No. 399

This is to certify that the Quality Management System of:

Steris Finn-Aqua Teollisuustie 2 FI-04300 Tuusula Finland

Has been assessed against the requirements of Annex III, Module H1 of the Pressure Equipment Directive 97/23/EC, and Schedule 4, Module H1 of the Pressure Equipment Regulations 1999 and conforms to the requirements for the products shown below:

Designing, manufacturing, testing and serving of equipment for decontamination, sterilization and water purification technologies for pharmaceutical, research, healthcare, food and dairy industries.

Approval is subject to the continued maintenance of the quality system in accordance with the requirements of the above Directive and Regulations, and continuing to comply with the EC Design Examination Certificate(s) as listed on the attached schedule.

Authorisation is hereby given to use the LRV Notified Body Identification Number in accordance with the requirements of the specified Directive and Regulations in relation to the products as identified above.

Certificate No:

0038/PED/HAM0662096/2

Original Approval: 01 February 2002

Current Certificate:

Certificate Expiry:

09 December 2015

31 October 2018

LRV Notified Body Number 0038

P. Mintzaridis on behalf of Lloyd's Register Verification

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LRV/ECD/PED/H1-System/April_2015/Rev.2



EC Certificate of Conformity Certificate 0038/PED/HAM0662096/2 Schedule

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Steris Finn-Aqua Teollisuustie 2 FI-04300 Tuusula Finland

Products

Certificate Number

FA-5000S DB Column 5000S, DB-Kolonni PED/H1/19004144, Issue 2 PED/H1/9006124, Issue 01

Issuing Notified Body

Lloyd's Register Verification 0038 Lloyd's Register Verification 0038

Schedule Issue:

04

Date of Schedule Issue:

09 December 2015

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