

Quality System Certificate

Certificate No.:
DGM – 893

Reference:
aur1i1901v60f841

Date of issue:
2019-03-06

Valid Until:
2022-03-06

Initial date of issue:
2016-09-27

This is to certify that the quality system of:

SSI Diagnostica A/S
Herredsvejen 2
3400 Hillerød
Danmark

fulfills the requirements in:

DS/EN ISO 13485:2016

The certificate covers the following activities:

Development, Manufacturing, Sales and Distribution of in vitro-diagnostics and blood products from animals

The certificate is valid provided that the quality system continues to conform to the above-mentioned scope and provided that the company does not introduce substantial changes to the quality system without the approval of Presafe Denmark A/S. Certifications to DS/EN ISO 13485:2012, EN ISO 13485:2012, ISO 13485:2003, DS/EN ISO 13485:2016, EN ISO 13485:2016, ISO 13485:2016, DS/EN ISO 9001:2015, EN ISO 9001:2015, and ISO 9001:2015 include the requirements of any applicable corrigendum. The quality system certificate is issued pursuant to Presafe Denmark A/S' terms and conditions for the certification of quality systems for medical devices.

Presafe Denmark A/S

Notified Body, Identification No. 0543
Tuborg Parkvej 8, 2900 Hellerup, Denmark



Heidi Jørgensen
Authorized person

For Presafe Denmark A/S



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Additional site(s) covered by the certificate:

Hvidesten
Frederiksborgvej 71
3450 Allerød
Danmark