Certificate of Registration



This is to certify that the quality management system of

Siemens Healthcare Oy

part of the cerificate for Siemens Healthcare AB

Main site: Johanneslundsvägen 12-14, SE-194 87 Upplands Väsby, Sweden Additional sites according to appendix

has been assessed and registered by Intertek as conforming to the requirements of

ISO 9001:2008

The quality management system is applicable to

Marketing, sales, installation, training and service of medical devices and systems within In-vitro and In-vivo diagnostics.

Certificate Number:

0033846-00/A

Initial Certification Date:

25 September 2015

Certificate Issue Date:

25 September 2015

Certificate Expiry Date:

22 September 2018

Duplicate / Translation from a Swedish original

1639 ISO/IEC 17021

Thomas Andersson, CEO Intertek Certification AB

P.O. Box 1103, SE-164 22 Kista, Sweden

In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at certificate.validation@intertek.com or by scanning the code to the right with a smartphone.



Appendix



This appendix identifies the locations by the management system of

Siemens Healthcare AB

Siemens Healthcare AB

Johanneslundsvägen 12-14, SE-194 87 Upplands Väsby, Sweden

Siemens Healthcare Oy

Tarvonsalmenkatu 19, Fl-02600 Espoo, Finland (HK) Vaeike-Paala 1, 11415 Tallinn, Estonia Biroju iela 10, Lidosta Riga, Marupes novads LV-2167, Latvia J. Jasinskio 16c, LT-03163 Vilnius, Lithuania.

Certificate Number: Initial Certification Date: Certificate Issue Date: Certificate Expiry Date:

0033846-00 25 September 2015 25 September 2015 22 September 2018

Thomas Andersson, CEO Intertek Certification AB P.O. Box 1103, SE-164 22 Kista, Sweden

In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at certificate.validation@intertek.com or by scanning the code to the right with a smartphone.

The certificate remains the property of Intertek, to whom it must be returned upon request.



1639 ISO/IEC 17021

