Certificate of Registration

This is to certify that the quality management system of

Hammarplast Medical AB
including company Hammarplast Medical AS

Main site: Kartåsgatan 8, Lidköping SE-531 40, Sweden
Additional sites according to appendix:

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

SS-EN ISO 13485:2012

The quality management system is applicable to

Development, manufacturing and marketing of medical and surgical products produced in clean room environment. Manufacture and market medicine handling systems to the health care sector.

Certificate Number:	0042619-00
Initial Certification Date:	14 December 2007
Certificate Issue Date:	11 February 2016
Certificate Expiry Date:	11 February 2019

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

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ISO/IEC 17021

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.

Appendix

Intertek

This appendix identifies the locations by the management system of

Hammarplast Medical AB
including company Hammarplast Medical AS

Västra vägen 10,
SE-330 10 Bredaryd,
Sweden

Manufacturing of medical products produced in clean room environment

Västberga allé 36B,
SE-126 30 Hägersten,
Sweden

Manufacturing of surgical products

Hammarplast Medical AS
Söstramäe 8, EST-114 15 Tallinn,
Estonia

Manufacturing of medical products produced in clean room environment

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