



CERTIFICATE OF APPROVAL

This is to certify that the Quality Management System of:

***Astra Tech AB
Möln dal
Sweden***

*has been approved by Lloyd's Register Quality Assurance
to the following Quality Management System Standards:*

ISO 13485 : 2003

The Quality Management System is applicable to:

***Design, manufacture, final inspection and distribution of
medical devices for surgery, urotherapy, blood transfusion,
respiratory training and dental implant systems.***

*Approval
Certificate No: LRQ 0945580/B*

Original Approval: 8 May 1996

Current Certificate: 10 March 2006

Certificate Expiry: 31 March 2008

Issued by: Lloyd's Register Quality Assurance Limited



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This document is subject to the provision on the reverse

71 Fenchurch Street, London EC3M 4BS, United Kingdom. Registration Number 1879370

This approval is carried out in accordance with the LRQA assessment and certification procedures and monitored by LRQA.

The use of the UKAS Accreditation Mark indicates Accreditation in respect of those activities covered by the Accreditation Certificate Number 001

Macro Revision 10.1



CERTIFICATE OF APPROVAL

This is to certify that the Quality Management System of:

**Astra Tech AB
Aminogatan 1, PO Box 14
SE 431 21, Mölndal
Sweden**

*has been approved by Lloyd's Register Quality Assurance
to the following Quality Management System Standards:*

ISO 13485:2003

The Quality Management System is applicable to:

***Design, Manufacture, Final Inspection and Distribution of
Medical Devices for Dental Implant Systems, Surgical Wound
Drainage Sets, Blood Autotransfusion Sets, Surgical Drainage Tubes/Bags/Catheter
Connectors, Surgical Suction Systems,
Surgical Vein Strippers, Haemorrhoid Ligators, Urological
Intermittent Catheters, Urological Stoma Drainage Catheters, Respiratory
Physiotherapy Training Systems.***

This certificate forms part of the approval identified by certificate number 0945580

Approval
Certificate No: 0945580/E

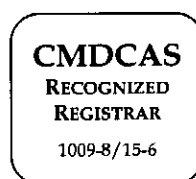
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LT Medcraft

Lorraine Medcraft, International Team Manager
Issued by: LRQA, Inc., Houston



*This document is subject to the provision on the reverse.
1401 Enclave Parkway, Suite 200, Houston, Texas 77077 USA*

This approval is carried out in accordance with the LRQA assessment and certification procedures and monitored by LRQA.

Macro Revision 12