



APPROVAL OF CONFORMITY CERTIFICATE

*In accordance with the requirements of the Directive 93/42/EEC and
Medical Devices Regulation 2002:618.*

This is to certify that the Quality Management System of:

**Astra Tech AB
Mölndal, Sweden**

*has been assessed against the requirements of Annex II of the Directive 93/42/EEC, and
the Medical Devices Regulation 2002:618 and conforms to the requirements for the
products shown below:*

Products as listed on Certificate Schedule.

*This certificate is valid only in association with the certificate schedule bearing the same
number on which the products applicable to this approval are listed.*

*Approval is subject to the continued maintenance of the quality system in accordance
with the requirements of LRQA approval 0945580/B and the requirements of the above
mentioned Directive and Regulation.*

*Authorisation is hereby given to use the LRQA Notified Body Registration Number in
accordance with the requirements of the specified Directives/Regulations in relation to
the products as identified above.*

Approval
Certificate No: LRQ 0945580/C

Original Approval: 8 May 1996

LRQA Notified Body
Registration No: 0088

Current Certificate: 1 April 2005

Certificate Expiry: 31 March 2008

RT Medcraft

Issued By: Issued by Lloyd's Register Quality Assurance Limited



APPROVAL OF CONFORMITY CERTIFICATE SCHEDULE

*In accordance with the requirements of the Directive 93/42/EEC and
Medical Devices Regulation 2002:618.*

**Astra Tech AB
Mölndal, Sweden**

Annex II Products

Dental Implant Surgical Components (Class IIa or Class IIb)
Dental Implant Prosthetics and Laboratory Components (Class IIa)
Dental Implant Instruments (Class IIa)
Surgical Suction Systems (Class IIa)
Varicose Vein Instruments (Class IIa)
Haemorrhoid Ligation Instruments (Class IIa)
Urotherapy Devices (Class IIa)
Surgical Wound Drainage and Blood Reinfusion Systems (Class IIa or Class IIb)
Bone Trap (Class IIa)

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This document is subject to the provision on the reverse
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LLOYD'S REGISTER QUALITY ASSURANCE