



APPROVAL OF CONFORMITY CERTIFICATE

In accordance with the requirements of the Medical Devices Directive 93/42/EEC and the Medical Devices Regulations 2002, UK Statutory Instrument 2002 No. 618

This is to certify that the Quality Management System of:

**AS Chemi-Pharm
Tallinn
Estonia**

has been assessed against the requirements of Annex V of the Medical Devices Directive 93/42/EEC, and the Medical Devices Regulations 2002 and conforms to the requirements for the products shown on the attached schedule.

Approval is subject to the maintenance of the quality system in accordance with the requirements of the above Directive and Regulations.

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.

Certificate No: LRQ 4001506/B
Original Approval: 25 May 2004
Current Certificate: 15 September 2009
Certificate Expiry: 7 April 2010
LRQA Notified Body Number 0088

Issued by: Lloyd's Register Quality Assurance Limited



**APPROVAL OF CONFORMITY CERTIFICATE
CERTIFICATE LRQ 4001506/B SCHEDULE**

**In accordance with the requirements of the Medical Devices
Directive 93/42/EEC and the Medical Devices Regulations 2002, UK
Statutory Instrument 2002 No. 618**

**AS Chemi-Pharm
Tallinn
Estonia**

Class II Products

BACTICID
CHEMIHYD DES
DES INSURANCE
STERISEPT
VIROBAC PLUS
ALCADENT
ACIDENT
CLEAN FOAM
WHITEWASH DES MD
CHEMISEPT WIPES MD
CHEMISEPT MD
CHLORINEX - 60 MD
HEALTHY SKIN GEL MD
SMELL NET MD
STERIDENT
CHEMIPHARM DES NEW MD
CHEMIDES PULVER
CHEMISEPT HF

Schedule Issue: 3

Date of Schedule Issue: 15 September 2009

LRQA Notified Body Number 0088

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Page 1 of 1

This document is subject to the provision on the reverse
71 Fenchurch Street, London EC3M 4BS United Kingdom. Registration number 1879370