

## 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:

**Vio Healthcare Ltd  
[incorporating Viomedex Ltd]  
Hailsham, East Sussex  
United Kingdom**

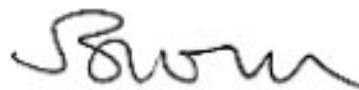
has been assessed against the requirements of Annex V of the Medical Devices Directive 93/42/EEC, and the Medical Devices Regulations 2002 and conforms to the requirements for the products shown on the attached schedule.

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 4000716/C  
Original Approval: 7 April 1997  
Current Certificate: 1 April 2011  
Certificate Expiry: 31 March 2014

LRQA Notified Body Number 0088



Issued by: Lloyd's Register Quality Assurance Limited

**APPROVAL OF CONFORMITY CERTIFICATE  
CERTIFICATE LRQ 4000716/C 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

**Vio Healthcare Ltd  
[incorporating Viomedex Ltd]  
Hailsham, East Sussex  
United Kingdom**

**Class II Products  
Non-sterile Breathing Circuits**

**Class I Sterile/Measuring Products  
Sterile Surgical Skin Markers  
Disposable Amniotic Membrane Perforator**

Schedule Issue: 01  
Date of Schedule Issue: 1 April 2011  
LRQA Notified Body Number 0088



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