Lloyd's Register
LRQA

EC CERTIFICATE – PRODUCTION QUALITY ASSURANCE

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:

Medical Technology Ltd
PO Box 223, Harbour Court, Les Amballes,
St Peter Port, Guernsey
Channel Islands

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 4005105/B
Original Approval:	5 November 2009
Current Certificate:	28 April 2015
Certificate Expiry:	27 April 2018

LRQA Notified Body Number 0088

Issued by: Lloyd's Register Quality Assurance Limited

1 Trinity Park, Bickenhill Lane, Birmingham B37 7ES, United Kingdom

Lloyd's Register LRQA

EC CERTIFICATE – PRODUCTION QUALITY ASSURANCE CERTIFICATE LRQ 4005105/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

Medical Technology Ltd
PO Box 223, Harbour Court, Les Amballes,
St Peter Port, Guernsey
Channel Islands

Class IIa Products
Tenease – Vibration Therapy Unit
Kneease – Vibration Therapy Unit

Schedule Issue: 01

Date of Schedule Issue: 28 April 2015

LRQA Notified Body Number 0088

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