EC Certificate Full Quality Assurance System: Certificate US98/13734

The management system of

Mui Scientific, Division of H & A Mui Enterprises Inc.

145 Traders Boulevard East,
Unit #34, Mississauga, Ontario, L4Z 3L3, Canada

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Manometric pumps; non sterile manometric catheters for gastrointestinal motility diagnosis and barostat bag pump and barostat non sterile catheters for the diagnosis of anorectal disorders.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 12 August 2016 until 17 May 2021
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 17 May 2019
Issue 11. Certified since 17 May 1998

Certification is based on reports numbered WW/MC 08933

Authorised by

SGS United Kingdom Ltd, Notified Body 0120
202B Worle Parkway, Weston-super-Mare, BS22 6WA UK
t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com

SGS CE 02 0215