EC Certificate Production Quality Assurance System: Certificate US07/1105

The management system of

PriMed Instruments Inc.

1080 Tristar Drive, Unit 14,
Mississauga, Ontario, L5T 1P1, Canada

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Sterile single use endoscopic biopsy forceps for gastroenterology use.

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

This certificate is valid from 26 September 2016 until 26 September 2021
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 27 August 2019
Issue 5. Certified since 26 September 2007

Certification is based on reports numbered WWMC 600540

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 14 0215