Certificate US11/82174

The management system of

The 3CPM Company

7402 York Road, Suite 100,
Towson, MD, 21204, United States

has been assessed and certified as meeting the requirements of

ISO 13485:2003
EN ISO 13485:2012

For the following activities

Design, manufacture, distribution, and service of Electrogastrograms.

This certificate is valid from 19 June 2015 until 7 October 2017 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 24 March 2017
Issue 4. Certified since 7 October 2011
Multiple certificates have been issued for this scope
The main certificate is numbered [# see GSP.02 5.5.8 (e.g. XX03/0000.0)]

Authorised by

SGS United Kingdom Ltd Systems & Services Certification
Rossmore Business Park Ellesmere Port Cheshire CH65 3EN UK
t +44 (0)151 350-6666 f +44 (0)151 350-6600 www.sgs.com

UKAS MANAGEMENT SYSTEMS
0005

SGS 13485-2 1114

Certificate US11/82173

The management system of

The 3CPM Company

7402 York Road, Suite 100,
Towson, MD, 21204, United States

has been assessed and certified as meeting the requirements of

ISO 13485:2003

For the following activities

Design, manufacture, distribution, and service of Electrogastrograms.

Effective Date 19 June 2015 Expiry Date 7 October 2017
Re certification audit due before 24 March 2017
Valid subject to satisfactory surveillance audits.
Issue 4. Certified since 7 October 2011

Authorised by
Jan Saunders – Business Manager

SCC Accredited
CB-MS
OCSM
Accrédité CCN

SGS United Kingdom Ltd Systems & Services Certification
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CMDCAS recognised registrar

SGS 13485 CMDCAS 0311

EC Certificate Full Quality Assurance System: Certificate US11/82172

The management system of

The 3CPM Company

7402 York Road, Suite 100,
Towson. MD. 21204. United States
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

Electrogastrograms used in the diagnosis of motility disorders in the gastrointestinal tract.

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 19 June 2015 until 7 October 2020 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 24 March 2017
Issue 2. Certified since 7 October 2011

Certification is based on reports numbered WW/ME 603058

Authorised by

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SGS CE 02 0215