TÜVRheinland®
Precisely Right.

Product Safety and Quality
Certification Department

TÜV Rheinland Product Safety GmbH D-51105 Cologne

MEDPLUS INC.
4th Floor, Building C-4,
Gaosha Industrial Zone
Mr. Zhong
Guangzhou
GUANGDONG PROVINCE 511495
CHINA

Tel. +49 180 3 112 112
Fax +49 180 3 000 169
Mail safety@de.tuv.com

Customer Service Center for
Product Safety and Quality
Tel. +49(0)221-806-1400
Mail service@de.tuv.com

Cologne, January 02, 2008

Application for : QMS Produktion, Anhang V MDD
Certificate No. : DD 60019660 SHEET 0001
Techn. appliance : Only for QM-System audit
Test requirements : Richtlinie 93/42/EWG

Dear Mr. Zhong,

Your Quality Management System has been tested and found to be in accordance with the above mentioned requirements.

Enclosed please find the certificate
No. DD 60019660 0001.

Kind regards

Certification body

X. Ren

Test sample: no, documentation available

TÜV Rheinland
Product Safety GmbH

TÜV Rheinland Group

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APPROVAL

EC Directive 93/42/EEC Annex V, Article 3
Quality Assurance System Production

Registration No.: DD 60019660 0001

Report No.: 15022512 001

Manufacturer:
MEDPLUS INC.
4th Floor, Building C-4,
Gaosha Industrial Zone
ZhongCun, Panyu District
Guangzhou

Guangdong Province 511495
China

Scope:
Manufacturing of Medical Devices

Products: see attachment

Date of Expiry: 29.12.2012

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex V, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex V, Article 4 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Notified Body

Cologne, 02.01.2008

X. Ren

TÜV Rheinland Product Safety GmbH - Am Grauen Stein - D-51105 Köln
Accredited by Zentralstelle der Länder für Sicherheitstechnik (ZLS) and
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten (ZLG).

Notified under No. 0197 to the EC Commission.

CE The CE marking may be used if all relevant and effective EC Directives are complied with. CE

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Doc. 1/1, Rev. 0

TÜV Rheinland
Product Safety GmbH

Am Grauen Stein, D-51105 Köln

Attachment to
Registration No.: DD 60019660 0001
Report No.: 15022512 001

Manufacturer:

MEDPLUS INC.
4th Floor, Building C-4,
Gaosha Industrial Zone
ZhongCun, Panyu District
Guangzhou

Guangdong Province 511495
China

Scope:

Products:

- Urethral catheters
- Tracheal tubes
- Breathing circuits
- Biopsy kits
- Loss of resistance syringes for single use

For the following medical devices the scope covers only the aspects of manufacture concerned with securing and maintaining sterile conditions:

- Tourniquets
- Skin markers

Köln, den 02.01.2008

X. Ren

TÜV Rheinland Product Safety GmbH
TÜVRheinland®
Zertifizierungsstelle