

**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
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CHINA

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Customer Service Center for
Product Safety and Quality
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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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Cologne HRB 25960
UST-ID Nr.: DE 811835490

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

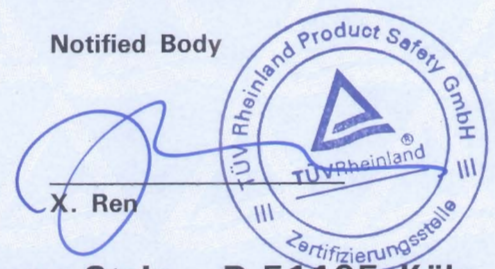
Products: see attachment

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



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

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

