

Circular Informativa

N.º 046/CD/550.20.001

Data: 05/02/2020

Assunto: **Certificado falso – Fabricante Changzhou Yihua Sanitation Material Co., Ltd**

Para: Divulgação geral

Contacto: Centro de Informação do Medicamento e dos Produtos de Saúde (CIMI); Tel. 21 798 7373;
Fax: 21 111 7552; E-mail: cimi@infarmed.pt; Linha do Medicamento: 800 222 444

O organismo notificado TÜV SÜD Product Service GmbH Zertifizierung Medizinprodukte (0123) identificou um certificado falso (em anexo) relativo aos dispositivos **luvas de exame estéreis, luvas de nitrilo estéreis e cateteres uretrais estéreis de uso único**, do fabricante **Changzhou Yihua Sanitation Material Co., Ltd**.

Em Portugal, não foram identificados registos da comercialização de dispositivos médicos deste fabricante, mas, atendendo a que existe livre circulação de produtos no Espaço Económico Europeu, o Infarmed recomenda que os dispositivos associados ao certificado supramencionado não sejam adquiridos nem utilizados, uma vez que apresentam aposta marcação CE 0123 falsa.

A deteção, em Portugal, destes dispositivos e deste certificado deve ser reportada à Direção de Produtos de Saúde do Infarmed através dos contactos: tel.: +351 21 798 72 35; e-mail: daps@infarmed.pt.

O Conselho Diretivo

Anexo – Certificado falso

ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ CERTIFICADO ◆ CERTIFICAT

EC Certificate
Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)
No. G2 14 09 36477 016

Manufacturer: **Changzhou Yihua Sanitation
Material Co.,Ltd**
No.160,Zhenlu Town.
213115Changzhou,JiangsuProvince
PEOPLE'S REPUBLIC OF CHINA

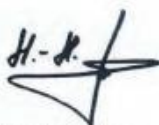
EC-Representative: **Shanghai International Holding
Corp. GmbH (Europe)**
Eiffestraße 80
20537 Hamburg
GERMANY

Product **Sterile Latex Examination Gloves,
Sterile Nitrile Examination Gloves,
Category(ies): Sterile Urethral Catheters for Single Use**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: SH14058EXT02

Valid from: 2014-11-24
Valid until: 2019-11-23

Date, 2014-11-24

Hans-Heiner Junker

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123
Page 1 of 2

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

TÜV SÜD
Product Service

TÜV SÜD
634172

TÜV

ZERTIFIKAT ♦ CERTIFICATE ♦ 認証証書 ♦ CERTIFICADO ♦ CERTIFICAT


Product Service

EC Certificate
Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)
No. G2 14 09 36477 016

Facility(ies): Changzhou Yihua Sanitation Material Co.,Ltd
No.160,Zhenlu Town,213115Changzhou,Jiangsu
Province,PEOPLE'S REPUBLIC OF CHINA

Page 2 of 2

A1 / 04.11

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

