



Product Service

# EC Certificate

## Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 16 11 61018 014

**Manufacturer:** **Medipro (Malaysia) Sdn. Bhd.**

Lot 15, LebuH Hishamuddin 3  
Kawasan 20, Bandar Sultan Suleiman  
42000 Port Klang, Selangor Darul Ehsan  
MALAYSIA



**Facility(ies):**

Medipro (Malaysia) Sdn. Bhd.  
Lot 15, LebuH Hishamuddin 3, Kawasan 20, Bandar Sultan  
Suleiman, 42000 Port Klang, Selangor Darul Ehsan, MALAYSIA

**Product  
Category(ies):**

**Sterile Surgical Drapes, Gowns, Mask, Caps,  
Surgical Supporting Products, Irrigation Pouch,  
Surgical Packs and Kits**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

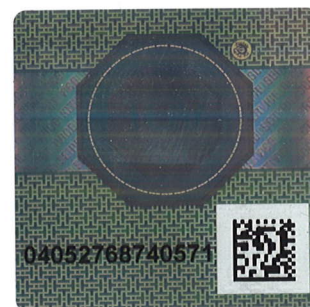
**Report No.:** MYQMH1116035-721416813

**Valid from:** 2017-01-04

**Valid until:** 2022-01-03

**Date,** 2017-01-02

Stefan Preiß



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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