

**EC Certificate**  
**Directive 93/42/EEC Annex II, excluding Section 4**  
**Full Quality Assurance System**  
**Medical Devices**

**Registration No.:** HD 60144003 0001

**Report No.:** 17055844 008

**Manufacturer:** Shenzhen Enmind Technology  
Co., Ltd.  
Room 201, Block A  
No. 1, Qianhai Road 1  
Qianhaishen Port Cooperative District  
Shenzhen  
518000 Guangdong  
China

**Products:** Infusion Pumps, Syringe Pumps

(see attachment for site included)

Replaces Approval, Registration No.: DD 60109366 0001

**Expiry Date:** 2024-05-27

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

**Effective Date:** 2019-12-02

**Date:** 2019-12-02

Notified Body



**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**  
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC  
concerning medical devices with the identification number 0197.

**TÜV Rheinland**  
**LGA Products GmbH**  
**Tillystraße 2, 90431 Nürnberg**

Doc. 1/1, Rev. 0

**Attachment to  
Certificate**

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Room 201, Block A  
No. 1, Qianhai Road 1  
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Shenzhen  
518000 Guangdong  
China

**Site included:**

5th Floor, Block A, Defengsheng Building,  
No.41 Dabao Road, Bao'an District 23,  
Shenzhen, 518101, P.R.China

**Date:** 2019-12-02

**Notified Body**



**Fuxiu Sheng**