

# EC Certificate



## Production Quality Assurance MDD Annex V

Registration No.: DD 2016352-1


Manufacturer: Jiangsu Caina Medical Co., Ltd.  
No. 23, Huanxi Road,  
Zhutang Town, Jiangyin,  
214415 Jiangsu  
P.R. China

Products: Disposable Syringes with Needle, Disposable Intravenous Needles, Disposable Insulin Syringes, Disposable Safelock Insulin Syringes, Disposable Huber Needles, Disposable Safety Huber Needles, Disposable Blood Collection Sets, Disposable Safelock Blood Collection Sets, Disposable Insulin Pen Needles, Disposable Safety Insulin Pen Needles, Disposable Neuraxial Fit Tip Syringes, Disposable Dental Needles, Blood Collection Needles, Safety Blood Collection Needles, Intravenous Catheters, Safety Intravenous Catheters, Retractable Safety Syringes, Disposable Syringes with Safety Needle, Disposable Safety Hypodermic Needles, Disinfection Caps.  
Aspects of manufacture concerned with securing and maintaining sterile conditions: Disposable Syringes without Needle, Enteral Feeding and Irrigation Syringes, Disposable LOR Syringes, Needleless Connectors, Disposable Medical Face Masks, Disposable Oral Specimen Collection Kits

The Notified Body hereby declares that the requirements of Annex V 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 V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Report No.: 15063456 008  
Effective date: 2021-01-27  
Expiry date: 2024-05-26  
Issue date: 2021-01-27

  
Herbert Zhong  
TÜV Rheinland LGA Products GmbH  
Tillystraße 2 · 90431 Nürnberg · Germany

The seal of TÜV Rheinland LGA Products GmbH, featuring a circular design with the company name and a stylized triangle logo in the center.

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