

## CERTIFICATE

### **Full Quality Assurance System**

Medical Devices Directive 93/42/EEC Annex II (Excluding Section 4)

M.2019.106.11727-1 Design Examination Certificate Was Prepared for Class III Products Defined in This Certificate.

: Shandong Haidike Medical Products Co., Ltd. Company Name

Company Address : Plant No.1, Science and Technology Enterprise Incubator Park,

Shan County, Heze City, Shandong Province, China.

: 93/42/EEC Medical Devices Directive - Annex II Related Directives and Annex

(Excluding Section 4)

Product : - Sterile Braided Coated Violet or Undyed Absorbable Polyglycolic Acid

(PGA) Suture with or without Needle - Class III

- Sterile Braided Coated Violet or Undyed Absorbable Polyglactin (PGLA)

Suture with or without Needle - Class III Sterile Single Use Non-Absorbable Silk Suture with or without needle - Class IIb

- Sterile Single Use Non-Absorbable Nylon Suture with or without needle - Class Ilb

- Sterile Single Use Non-Absorbable Polyester Suture with or without needle - Class Ilb

Sterile Single Use Non-Absorbable Polypropylene

Suture with or without needle - Class Ilb

**GMDN** : 13908, 17471, 13910, 13905, 13906, 13909

Product Types are attached.

Certificate Number : M.2019.106.11727

Report Number : MD.3626.IB Initial Assessment Date : 10.02.2018 Registration Date : 10.04.2019 Revision Date /No : 22.07.2019/01

Expiry Date : 09.04.2024

UDBM hereby declares that the requirements of Annex II, excluding section 4 of the 93/42/EEC Directive have been my for the listed products. The above named manufacturer has established and applied a quality assurance system, which is subject to periodic surveillance audits, defined by Annex II, section 5 of the forementioned directive. According to Annex Lisection 4 an EC design-examination certificate is required for placing the Class Lidevices on the market. UDBvfs responsibility for class I devices covered by the EC certificate is limited to manufacturing issues related to safeguarding and maintaining stelle conditions, if the device is sterile; and manufacturing issues related to product's conformity with metrological requirements. If it has measurement function. This certificate remains as the property of UDBM international Certification Auditing Training. Centre Industry and Trade Inc. Co. to whom it must be returned upon request. The above named company and UDBM must keep a copy of this certificate for 5 years from the registration of the certificate. Usage of the CE mark is under the responsibility of the manufacturer with the completion of EC Declaration of Conformity. The above mentioned company must notify all changes related with the approved product to UDBM. If UDBM will not renew the validity of this certificate in question, the mentioned company should stop placing the product on the market. The validity of the certificate can be checked through www.udem.com.tr.

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PARTNERS CERTIFICATE ASSURANCE



# ertificate

ISO 13485 : 2016

### Shandong Haidike Medical Products Co., Ltd.

Plant No.1, Science and Technology Enterprise Incubator Park, Shan County, Heze City, Shandong Province, China.

This certificate shows that the medical devices - quality management system (EN ISO 13485:2016) of the above company was approved by PCA Certification for the following scope, the validity of the certificate depends on the company's pass the annual surveillance audits and company's maintenance the related management system conditions according to international accreditation criteria

#### SCOPE

 $Design, development, manufacture\ and\ sales\ of\ Disposable\ suture\ needles\ with\ non-absorbable\ threads,\ sterile\ and\ sales\ of\ Disposable\ suture\ needles\ with\ non-absorbable\ threads,\ sterile\ needles\ with\ non-absorbable\ sterile\ needles\ need$ syringe for single use, sterile infusion sets for single use, disposable lancets for blood specimen collection, absorbable surgical sutures with or without needle. Medical face mask and Medical protective clothing

GROUP CODE

A

**Certificate No** 

: TC-75059

**Registration Date** 

: 27.04.2018

Reissue Date

: 19.04.2021/01

**Expiry Date** 

: 26.04.2024

**Certificate Period** Exclusion

: 3 Years (From the date of registration)

: 7.5.3

ACCREDITED

Management Systems Certification Body

**MSCB-103** 



PCA Certification Approval

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