MED CERT®

Certification. Medical Only.

EC Certificate of Conformity

The Notified Body

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany

herewith certifies that the company:

Anni And Anna Limited.
RM 16, 6F, Hong Leong Plaza 33 Lok.Yip Road
Fanling, NT
HongKong

has introduced and maintains a quality assurance system for the products / product categories listed in the appendix.

The compliance of this quality assurance system with the below mentioned requirements of the **Council Directive 93/42/EEC** was verified by an audit:

Annex V

This certification is subject to surveillance by MEDCERT.

This certificate is valid until 06 November 2023

Report No.: 7372IA01F
Process No.: QS – 7372
Certificate No.: 7372GB414190830

Hamburg, 30 August 2019

MEDCERT Certification Body
(Lorenz Runge)

The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15

MEDCERT Identification Number: 0482

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Appendix of EC Certificate of Conformity

Process No.: QS – 7372
Certificate No.: 7372GB414190830

List of products / product categories included in the scope of certificate

- **Disposable inflation device kits**
- **Haemostatic valves**
- **Pressure lines**

– End of list –

This appendix is integral part of the above-referenced certificate.
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The Notified Body

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany

herewith certifies that the company:

Anni And Anna Limited.
RM 16, 6F, Hong Leong Plaza 33 Lok.Yip Road
Fanling, NT
HongKong

has introduced and maintains a quality assurance system
for the aspects of manufacture concerned with securing and maintaining sterile conditions

for the products / product categories listed in the appendix.

The compliance of this quality assurance system with the below mentioned requirements of the **Council Directive 93/42/EEC** was verified by an audit:

Annex V

This certification is subject to surveillance by MEDCERT.

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Appendix of EC Certificate of Conformity

Process No.: QS – 7372
Certificate No.: 7372GB415190830

List of products / product categories included in the scope of certificate

- **Control syringes**
- **Disposable TR-Closure band/Radiseals**
- **Disposable balloon inflation devices**
- **Disposable angio-closure pads**
- **Disposable towel/sheet clips**

– End of list –

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