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Page 1 of 1

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Authorised by

Certification is based on reports numbered CN/XMN 6177MDD

This certificate is valid from 15 August 2018 until 14 August 2023 and remains valid subject to satisfactory surveillance audits. Re certification audit due before 18 April 2021 Issue 5. Certified since 18 March 2014

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

Sterile Circumcision Suture Device II
Sterile Circumcision Suture Device II

For the following products

on medical devices, Annex II (excluding Section 4)

Directive 93/42/EEC

has been assessed and certified as meeting the requirements of

North Side of the Biological Industrial Park, Enjiang Industrial Zone,
Yongfeng County, Ji'an City, Jiangxi Province, 331500, P.R. China

Jiangxi Lanhge Medical Instrument Co., Ltd.

The management system of

EC Certificate Full Quality Assurance System: Certificate CN14/30386

