EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60103517 0001

Report No.: 17049711 001

Manufacturer:

SHENZHEN SUPERLINE
TECHNOLOGY CO., LTD.
Room B-314, Century Holidays
Plaza, 9030 Shennan Ave., Nanshan
518053 Shenzhen
China

Products:

Orthodontic Wires, Dental Root-canal Instruments,
Gutta Percha Points, Sterile Absorbent Paper Points

(see attachment for additional site included)

Replaces Approval, Registration No.: DD 60082196 0001

Expiry Date: 2020-09-28

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.

Effective Date: 2015-09-29

Date: 2015-09-29

Notified Body

X. Ren

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.

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Doc. 1/1, Rev. 0

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: DD 60103517 0001
Report No.: 17049711 001

Manufacturer:
SHENZHEN SUPERLINE TECHNOLOGY CO., LTD.
Room B-314, Century Holidays Plaza, 9030 Shennan Ave., Nanshan
518053 Shenzhen
China

Site included:

5F-8F, Bldg. A, Zone C, Shiwei Datianyang Ind. Park,
Jiangshi Community, Guangming, Shenzhen, 518105 China

Notified Body

Date: 2015-09-29

X. Ren

TÜVRheinland LGA Products GmbH Zertifizierungsstelle