

**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

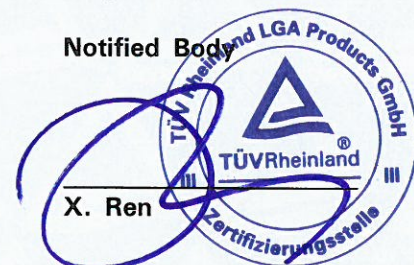
**Products:** Orthodontic Wires, Dental Root-canal Instruments,  
Gutta Percha Points, Sterile Absorbent Paper Points  
  
(see attachment for additional site included)  
  
Replaces Approval, Registration No.: DD 60082196 0001

**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



**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.

**TÜV Rheinland**  
**LGA Products GmbH**  
**Tillystraße 2, 90431 Nürnberg**

**Attachment to  
Certificate**

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518053 Shenzhen  
China

**Site included:**

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

**Date:** 2015-09-29

