

APPROVAL

EC Directive 93/42/EEC Annex V, Article 3
Quality Assurance System Production

Registration No.: DD 60034343 0001

Report No.: 16800186 001

Manufacturer: Specialty Medical Supplies
(Beijing) Co., Ltd.
Jin Yu Rui Science Zone Qiao
Zi Town Huai Rou District

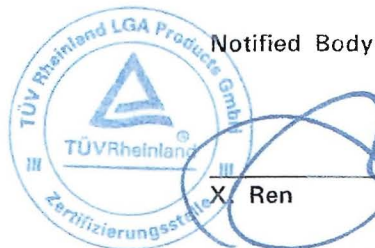
Beijing, 101402
China

Scope: Manufacture of Disposable Lancets
Replaces Approval, Registration No.: DD 60012938 0001

Date of Expiry: 05.12.2015

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex V, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex V, Article 4 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Date 06.12.2010



TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Accredited by Zentralstelle der Länder für Sicherheitstechnik (ZLS) and
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten (ZLG).

Notified under No. **0197** to the EC Commission.

CE The CE marking may be used if all relevant and effective EC Directives are complied with. CE