EC Declaration of Conformity

Manufacturer: Shandong Chengwu Medical Products Factory
274200 Southern End of Quancheng Road, Chengwu County,
Shandong Province, P.R.China

European Representative: SUNGO Europe B.V.
Olympisch Stadion 24, 1076DE Amsterdam

Product Name: **Disposable sterile venous blood specimen collection needle**

Model: soft-connection; hard-connection

UMDNS Code: 12736

Classification (MDD, Annex IX): **Class IIa, Rule 6**

We herewith declare that the above mentioned products meet the transposition into national law, the provisions of the following EC Council Directives and All applicable harmonised Standards. All supporting documentations are retained under the premises of the manufacturer.
Shandong Chengwu Medical Products Factory is exclusively responsible for the declaration of conformity

DIRECTIVES

Medical Device Directive:
COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices (MDD 93/42/EEC), Amended by DIRECTIVE 2007/47/EC of 5 September 2007.

Notified Body : TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339, MÜnchen, Germany

NB Identification number: 0123

(EC) Certificate(s): G2 103129 0002 Rev. 00

Expire date of the Certificate: 2024-02-22

Start of CE Marking: 2019-12-06

Place, Date of Issue: Shandong Chengwu 2019.12.06

Signature/date: Wang Jicun 2019.12.06

Name: Wang Jicun

Position: General Manager