EC Declaration of Conformity

Manufacturer: Chengdu Rich Science Industry Co., Ltd.
Enterprises Registration Address: No. 66 Tianqindong Street, West Hi-Tech Development Zone, Chengdu, China
Factory Address: No. 66 Tianqindong Street, West Hi-Tech Development Zone, Chengdu, China

We, the manufacturer, herewith declare that the products

Single Use Evacuated Blood Collection Needles

UMDNS-Code: 12736;

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class IIA according to Annex IX of the Directive 93/42/EEC. It bears the mark

CE 0197

The product concerned has been designed and manufactured under a quality management system according to Annex V of Directive 93/42/EEC.

Compliance of the designated product with the Directive 93/42/EEC has been assured via assessment of the quality management system by the Notified Body

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

following the procedure relating to the EC Declaration of Conformity set out in Annex V of Directive 93/42/EEC.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

Company: Chengdu Rich Science Industry Co., Ltd.
Address: No. 66 Tianqindong Street, West Hi-Tech Development Zone, Chengdu, China

Chengdu, January 14, 2016

Place, date

Legally binding signature, Function

EC Declaration of Conformity
RQ/CE-ZKCXZ-C-13(LOC no., Revision)
EC Declaration of Conformity

Manufacturer: Chengdu Rich Science Industry Co., Ltd.

Enterprises Registration Address:
No.66 Tianqindong Street, West Hi-Tech Development Zone, Chengdu, China

Factory Address:
No.66 Tianqindong Street, West Hi-Tech Development Zone, Chengdu, China

We, the manufacturer, herewith declare that the products
Evacuated Blood Collection Tubes

UMDNS-Code: 14183;

meet the provisions of Directive 98/79/EC which apply to them.

The medical device has been assigned to In Vitro Diagnostic Medical Devices Directive 98/79/EC. It bears the mark

The product concerned has been designed and manufactured under a quality management system according to In Vitro Diagnostic Medical Devices Directive 98/79/EC.

Compliance of the designated product with the Directive 98/79/EC has been assessed and certified by the Notified Body

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

following the procedure relating to the EC Declaration of Conformity set out in Annex III of Directive 98/79/EC.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

Company: Chengdu Rich Science Industry Co., Ltd.
Address: No.66 Tianqindong Street, West Hi-Tech Development Zone, Chengdu, China

Chengdu, January 14, 2016
Place, date

EC Declaration of Conformity
RG/CE-CXG-C-13(DOC no., Revision)