CE

EC Declaration of Conformity

Directive 93/42/EEC Medical Devices

(applicable to **Medical Device Class I**)

Manufacturer: **Shenzhen Zhonghe Headway Bio-Sci & Tech Co., Ltd.**

6/F, Bldg R1-B, South District, High-Tech Industrial Park, Shennan Road, Shenzhen, 518057, China

Product/s: Collection Card for Helicobacter Pylori Test

Category: **Medical Device Class I**

Conformity assessment route: **Annex VII**

Sterile: **No**

We, the Manufacturer, herewith declare with sole responsibility that our product/s mentioned above meet/s the provisions of the Directive 93/42/EEC of the European Parliament and of the Council on Medical Devices.

Signed on: Sept. 18, 2017. Place: Shenzhen, China.

Represented by

Signature (on behalf of the manufacturer) _____

Full Name of authorized signatory: Cui Haiping

Position held in the company: General Manager

Company Seal/Stamp: