CE

Declaration of conformity

CE

Manufacturer:
Wuxi NEST Biotechnology Co., Ltd.
No.230,Xida Road, New District, Wuxi, Jiangsu, 214112,China

whose single Authorized Representative:
TECHLABS MEDIKAL VE KIMYA TIC. LTD. STI
A BLOK NO: 536-537 K:6,PERPA TIC.MRK,HALIL RIFAT PASA , MAH.,SISLI,ISTANBUL.TURKEY

We, the manufacturer, herewith declare that the products
Cell Culture Plates, Cell Culture Dishes, Cell Culture Flasks, Multi-layer Cell Culture Flasks, Biofactory, Cell Scrapers, Glass Bottom Cell Cutlure Dishes/ Plates, Centrifuge Tube, Microcentrifuge Tubes, PCR Products, Pipette Tips, Cryogenic Vials, Deep Well Plate, Serological Pipette, Pasteur Pipette, Petri Dish, Elisa Plate

Type:See attached list

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

The In-vitro diagnostic medical device has been assigned to other IVD products according to Annex II of the Directive 98/79/EC. It bears the mark

CE

The product concerned has been manufactured under a quality management system according to EN ISO 13485, 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: Wuxi NEST Biotechnology Co., Ltd.

Address: No.230,Xida Road, New District, Wuxi, Jiangsu, 214112,China

无锡 4/26-2017
Wuxi April 26th

Place , date

Jack Fang

Legally binding signature, Function