CE

EC DECLARATION OF CONFORMITY

According to Directive 98/79/EC on *in vitro* diagnostic medical devices, Annex III

Manufacturer: Shanghai Kehua Bio-engineering Co., Ltd.
1189 North Qinzhou Road, 200233, Shanghai, PEOPLE'S REPUBLIC OF CHINA

EC- Representative: Qarad EC-REP BV
Pas 257
2440 Geel
Belgium

Product: Antibodies to Thyroid Peroxidase Quantitative Detection KIT
(Chemiluminescent Immunoassay)

Product code:
KH-G-L-335-01-CE
KH-G-L-335-02-CE
KH-G-L-335-03-CE
KH-G-L-335-04-CE

Classification: Other device (all devices except Annex II and self-testing devices)

We, manufacturer, herewith declare under our sole responsibility that the above-mentioned product meets the provisions of the relevant EC Council Directives and Standards. All supporting documentation is retained under the premises of the manufacturer.

Place, Date of Issue: Shanghai, P.R. China, 2021-11-26

Signature of RA Supervisor

Rachel Yan

KHB