CE Declaration of conformity CE

According to In Vitro Diagnostic Medical Device Directive 98/79/EC

Manufacturer: GUANGZHOU JET BIO-FILTRATION CO., LTD
No.1 DouTang Road, YongHe Development Zone,
511356 Guangzhou, PEOPLE'S REPUBLIC OF CHINA

Authorized Representative: Akralab, S.L.
Pol. Ind. Las Atalayas Av. de la Antigua Peseta, 77 Buzón 20212 · 03114 Alicante · Spain

Description/Category: Disposable Laboratory Plastic Products.
Plastic Pipets Products Series, Centrifuge Tubes Products Series, Cell and Tissue Culture Products, Disposable Filtration Products, ELISA Microplate, Inoculating Loops and Needles, Liquid Transfer Troughs, Cuvette, Pipette Micro Tips, Deep-Well Plates, Serum & Sample Tubes, Freezing Tubes and PCR Tubes for In Vitro Diagnostics.

Device Classification : Others, Self-Declaration

Conformity assessment Route: Annex III

THE UNDERSIGNED HEREWITH DECLARE THAT THE STATED MEDICAL DEVICES MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 98/79/EC CONCERNING MEDICAL DEVICES; ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER. WE, THE MANUFACTURER, ARE EXCLUSIVELY RESPONSIBLE FOR THE DOC.

Quality Manager

Guangzhou
February 21, 2019 (Place and date of issue)

York Zhang (Name and signature of authorized person)