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: Riomavix, S.L.
Calle de Almansa 55, 1D, Madrid 28039 Spain;
leis@riomavix.com

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 of ANNEX II of IVDD

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 IN VITRO DIAGNOSTIC MEDICAL DEVICES; ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER. WE, THE MANUFACTURER, ARE EXCLUSIVELY RESPONSIBLE FOR THE DOC.

Quality Director

Guangzhou
April 25, 2022 (Place and date of issue)

Xiaoxiao Ma Xiao Xiao Ma (Name and signature of authorized person)