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, Pipette Micro Tips, Tissue Culture Plate, Cell and Tissue Culture Dishes, Cell and Tissue Culture Flask, Erlenmeyer Flask, Roller Bottles, Deep-Well Plates, Serum&Sample Tubes, Cryovials 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 Manager

Guangzhou
Sep 30th, 2021
(Place and date of issue)

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