Certificate of Registration

This is to certify that the
Quality Management System for Medical Devices
of

CML BIOTECH LIMITED

at

MANUFACTURING SITE- 1 : A1, A2, B1& B2, TOWER 1, INKEL INDUSTRIAL COMPLEX, ANGAMALY SOUTH -PO, ERNAKULAM DISTRICT, KERALA - 683 573, INDIA

MANUFACTURING SITE 2 : PLOT NO: 2, INKID INDUSTRIAL AREA, INKEL BUSINESS PARK, ANGAMALY SOUTH - PO, ERNAKULAM DISTRICT, KERALA – 683573, INDIA

REG. OFFICE : 4/434B, KARUKUTTY P.O, ANGAMALY, ERNAKULAM DISTRICT, KERALA, INDIA, PIN-683576

has been independently assessed and
is compliant with the requirements of:

ISO 13485:2016

For the following scope of activities:

MANUFACTURE & MARKETING OF STERILE & NON STERILE BLOOD COLLECTION TUBES (VACUUM / NON VACUUM), LABDISPOSABLES, ALCOHOL PREP PAD, ADHESIVE PLASTER, BLOOD COLLECTION NEEDLES, PRP TUBES & KITS, MICROBIOLOGICAL TRANSPORT MEDIUM & SYSTEMS, SPECIMEN COLLECTING SWABS, MICROBIOLOGICAL CULTURE MEDIUM & STAINING REAGENTS

Certificate Number: UQ - 2022053149

Validity of this certificate can be verified at www.ukcertifications.org.uk/verify

Date of Certification	31st May 2022
1st Surveillance Audit Due	30th May 2023
2nd Surveillance Audit Due	30th May 2024
Certificate Expiry	30th May 2025

Authorised Signatory