Certification Services Inc.

VRC

Quality Excellence

Certificate of Compliance

This is to certify the products manufactured by :

LORD'S MARK INDUSTRIES PVT.LTD.

HEAD OFFICE: 404, 4TH FLOOR, COSMOS MARY PARK,
KOLBAD CROSS LANE, THANE (WEST) -400 601, MAHARASHTRA, INDIA.
ADDITIONAL SITE: FACTORY: A-71, ROYAL INDUSTRIAL HUB, BHIWANDI KAMAN ROAD,
POMAN VILLAGE, VASAI (EAST) PIN: 401 208. MAHARASHTRA, INDIA

The technical documentation and test reports maintained by the organization has been reviewed and found to comply with the requirements of EC Directive: Regulations 2017/746 on Invitro Diagnostic Medical Device Regulations (98/79/EC) .This certificate of compliance is based on the technical file of the below -mentioned product. Technical Report and Documentation is at the Organization's Disposal. This is to certify that the Product "As mentioned below" is in conformity with all revisions of EC Directive: Regulations 2017/746 on Invitro Diagnostic Medical Device Regulations (98/79/EC) and conforms to Harmonized Standards IP 2018, IP 2022, ISO 9001:2015, ISO 13485:2016, ISO 14971:2012, DIN EN ISO 15223-1:2013, NBN EN 13641:2002, EN 13640:2002, NBN EN 13612:2002. standards, this certificate does not imply assessment of the series-production of the product.

for the following product:

Invitro Diagnostic Kits, Reagents, Rapid Test Kits & Instruments/Analyzers

Details as mentioned in Appendix A

Certificate No.: VCELM63981938

Date of Expiry: 10.12.2028

1st Surveillance on or before: 10.12.2026

2nd Surveillance on or before: 10.12.2027

Recertification on or before : 10.12.2028

Date of Issue: 11.12.2025

Authorized Signatory

This compliance certificate is approved subject to the organisation maintaining its system, standard, test result as per prescribed standard(s), the test results are exclusive done by orginisation and has no connection to VRCS. This certificate is not transferable and remain valid only above mention address and scope. This certificate is related to the management system, not to the products or services of the certified organizations. This complaince certificate does not provide the certified organization with immunity from its legal obligations. Organisation must check the acceptability of the certificate in requirement with concerened institutions from time to time ,vrcs shall not be responsibile for the same.This certifificate is not for legal purpose and becomes invalid if yearly surveillance is not done.

Lord's Mark Industries Limited

CE

Certification Services Inc.

VRC

Quality Excellence

This is to certify the products manufactured by :

LORD'S MARK INDUSTRIES PVT.LTD.

HEAD OFFICE: 404, 4TH FLOOR, COSMOS MARY PARK, KOLBAD CROSS LANE,
THANE (WEST) -400 601, MAHARASHTRA, INDIA.
FACTORY: A-71, ROYAL INDUSTRIAL HUB, BHIWANDI KAMAN ROAD, POMAN VILLAGE,
VASAI (EAST) PIN: 401 208. MAHARASHTRA, INDIA

for the following product(s):

This certificate covers the following products:

Enzymes, NonEnzymes and Hematology (IVD Kits)	Alkaline Phosphatase Test Kit, Albumin Test Kit, Amylase Test Kit, Glucose Test Kit, Creatinine Test Kit, Total Protein Test Kit, Bilirubin (T&D) Test Kit Calcium (Ars III) Test Kit, Chloride Test Kit, Total Cholesterol Test Kit, Gama GT Test Kit, GOT/AST Test Kit. GPT/ALT Test Kit, HDL Cholesterol (PPT) Test Kit, HDL Cholesterol (Direct) Test Kit ,Hemoglobin Test Kit, Microprotein Test Kit, Phosphorous Test Kit, Sodium Test Kit, Triglycerides Test Kit, Urea UV Test Kit, Urea Test Kit (Mod Berthelot), Uric Acid Test, Potassium Test Kit, Diluent, Lyse, Rinse, LDL Cholesterol (Direct) Test Kit ,Probe Cleaner, EZ Cleaner, RF Turbilatex, CRP Turbilatex,G6PD,CKMB,LDH,Magnesium, Lipase, ADA, Microalbumin, CKNAC, Ammonia,Anti -CCP, Ferritin Kit, HBA1C Kit, IgE Test Kit, Widal Test Kit, Homocysteine Test Kit
Rapid Test Products	HIV 1& 2, HBsAg, HCV, SYPHILIS, MALARIA Pf/Pv Ag, MALARIA Pf/PAN, DENGUE NS1 ,DENGUE DUO, Dengue IgG/IgM, LEPTOSPIRA, S.TYPHI, CIKUNGUNYA IgG/IgM, TROPONIN I , HCG, HIV Ag/Ab 4th Generation, SICKLE CELL, H. Pylori ,Pregnancy HCG ,COVID -19,Urine Strips 10 Parameter, 3 Parameter (GPK), 2 Parameter (GP) & 2 Parameter (GK),Uine Strips 14 Parameter.
Analyzers	Semi Auto Biochemistry ,Fully Auto Biochemistry Analyzer,3 Part Hematology Analyzer, 5 Part Diff Hematology Analyzer , Multiple T/P Fully Auto Analyzer, CLIA Immunoassay, Electrolyte Analyzer, Coagulation Analyzer , ESR Analyzer, Hemodialysis Machine, Urine Analyzer, Vet Hematology Analyzer (3Part / 5 Part)

Certificate No.: VCELM63981938

Date of Expiry: 10.12.2028

1st Surveillance on or before: 10.12.2026

2nd Surveillance on or before: 10.12.2027

Recertification on or before : 10.12.2028

Date of Issue: 11.12.2025

Authorized Signatory

Lord's Mark Industries Limited

Certification Services Inc.

VRC

Quality Excellence

This is to certify the products manufactured by :

LORD'S MARK INDUSTRIES PVT.LTD.

HEAD OFFICE: 404, 4TH FLOOR, COSMOS MARY PARK, KOLBAD CROSS LANE, THANE (WEST) -400 601, MAHARASHTRA, INDIA.

Additional Site: FACTORY: A-71, ROYAL INDUSTRIAL HUB, BHIWANDI KAMAN ROAD, POMAN VILLAGE, VASAI (EAST) PIN: 401 208. MAHARASHTRA, INDIA

for the following product(s):

This certificate covers the following products:

Group	Product Name	Class
TurbiChem	Alpha-1-Microglobulin with Calibrator	Class B
TurbiChem	Anti CCP with Calibrator	Class B
TurbiChem	Apo A1- With Calibrator	Class B
TurbiChem	Apo B-With Calibrator	Class B
TurbiChem	Apo E-With Calibrator	Class B
TurbiChem	ASO-With Calibrator	Class B
TurbiChem	Beta-2-Microglobulin - With Calibrator	Class C
TurbiChem	C3-With Calibrator	Class B
TurbiChem	C4 - With Calibrator	Class B
TurbiChem	CRP-With Calibrator	Class C
TurbiChem	Cystanin C-With Calibrator 1	Class B
TurbiChem	D-Dimer - With Calibrator	Class C
TurbiChem	Ferittin Kit	Class B
TurbiChem	HbA1C(Direct) - With Calibrator	Class B
TurbiChem	Hs-CRP-With Calibrator	Class C
TurbiChem	IgA - With Calibrator	Class B

Group	Product Name	Class
TurbiChem	IgE-With Calibrator	Class B
TurbiChem	IgG-With Calibrator	Class B
TurbiChem	IgM-With Calibrator	Class B
TurbiChem	Lipoprotein (a)-With Calibrator	Class B
TurbiChem	Microalbumin Urea-With Calibrator	Class B
TurbiChem	Pre-Albumin-With Calibrator	Class B
TurbiChem	Procalcitotin-With Calibrator	Class B
TurbiChem	Retinol Binding Protein-With Calibrator	Class B
TurbiChem	RF-With Calibrator	Class B
SEROLOGY	ASO	Class B
SEROLOGY	CRP	Class C
SEROLOGY	RF	Class B
SEROLOGY	RPR	Class C
SEROLOGY	CombiWidal - S	Class B
SEROLOGY	CombiWidal - OH	Class B
TurbiChem	IgE-With Calibrator	Class B

Certificate No.: VCELM63981938

Date of Issue: 11.12.2025

Date of Expiry: 10.12.2028

1st Surveillance on or before: 10.12.2026

2nd Surveillance on or before: 10.12.2027

Recertification on or before : 10.12.2028

Authorized Signatory

Lord's Mark Industries Limited

Obelis GROUP

Registered Office Address
Bd Brand Whitlock 30
B-1200 Brussels Belgium

Registered Adresss
Bd Général Wahis 53
B-1030 Brussels Belgium

www.obelis.net

+32 2 732 59 54

CERTIFICATE OF
IVDR NOTIFICATION

Reference No.: KP 0683-2025 **Date:** 23/06/2025

Order No.: EU MD 0257-2025

This is to certify that, according to the Regulation (EU) 2017/746, we, here at Obelis s.a. performed all notification duties and responsibilities as the European Authorized Representative (EC REP) of:

Name:
Lords Mark Industries PVT LTD

Address: A-71, Royal Industrial Hub, Poman,
Kaman Bhiwandi Road, Vasai (East)
Palghar
Maharashtra 401208 India

AS STIPULATED AND DEMANDED BY THE AFOREMENTIONED REGULATION.

The manufacturer declares that the Class (A/B/C/D) device(s) comply(ies) with the Regulation including all general safety and performance requirements.

The Manufacturer has provided Obelis s.a. (EC REP) with all the appropriate declarations as per the Regulation (EU) 2017/746 article 48 requirements, including the EC Declaration of Conformity (according to annex IV) confirming that their Class (A/B/C/D) in vitro diagnostic medical device(s), as stipulated here below, is/are fulfilling the applicable requirements of the Regulation (EU) 2017/746.

The notification of the following in vitro diagnostic medical device(s) has been completed by Obelis s.a. (EC REP) in compliance with the Regulation (EU) 2017/746 on the 23/06/2025

CLASS OF IVD DEVICE(S): Please See Annex A - List of Devices (7 Pages, 11 Devices)

As of the 24/06/2025, and provided that the Manufacturer will continue complying with the hereabove mentioned requirements*, he therefore:

- Is required to affix the CE marking on this(ese) device(s);
- May place this(ese) device(s) in the European Union and EEA territory.

Obelis s.a. - O.E.A.R.C.
Registered Address :
Bld Général Wahis 53
1030 Bruxelles
Tél. +32 2 732 59 54 - Fax +32 2 732 60 03

Mr. G. Elkayam CEO
Obelis sa

Obelis European Authorized Representative Center is a member of the European Association of Authorized Representatives (E.A.A.R.).
Obelis s.a. is ISO 9001 : 2015 and ISO 13485 : 2016 certified.

*This certificate will become void automatically upon termination of the EAR agreement or removal of the products from EAR Mandate

ANNEX to IVDR Certificates

Manufacturer: LORDS MARK INDUSTRIES PVT LTD

Country: India

#	Name of device	Intended use	Class and Rule under IIVDR	EMDN Code	GMDN Code	Basic UDI-DI	Mandated (Y/N)	Mandate starting date	Mandated Vigilance (Y/N)
1	LordsMed 3 Part Hematology Analyser	3 part Hematology analyzer is used to conduct a complete blood count (CBC) which includes red blood cell count (RBC) & RBC distribution width.White blood cell count(WBC) & WBC differential count in percentage and absolute value.Mean of hemoglobin & Mean corpuscular hemoglobin concentrations. Platelet counts, Platelet mean volume, Platelet criteria, Platelet distribution width & Large platelet cell ratio, as well as Hematocrit levels and Mean corpuscular volume on Lords Mark 3 part hematology analysers for professional and laboratory environment use only.	Class A, Rule 5b	W02020102 02	N/A	890619447LM 001EX	Y	18/06/2025	Y
2	LordsMed 5-Part Auto Hematology Analyzer	5 part Hematology analyzer is used for a complete blood count	Class A, Rule 5b	W02020101 01	N/A	890619447LM 002EZ	Y	18/06/2025	Y

		(CBC) on the principle of flow cytometry to differentiatew hite blood cells count(WBC) into their five major sub-populations—neutrophils, lymphocytes, monocytes, eosinophils, and basophils.Red blood cell count (RBC) & RBC distribution width. Mean of hemoglobin & Mean corpuscular hemoglobin concentrations. Platelet counts, Platelet mean volume, Platelet criteria, Platelet distribution width & Large platelet cell ratio, as well as Hematocrit levels and Mean corpuscular volume on Lords Mark 5 part hematology analysers for professional and laboratory environment use only.							
3	LordsMed Semi Auto biochemistry Analyzer	Lords Mark semi Auto Biochemistry Analyzer is used for quantitative analysis of clinical chemistry of specific chemical components in bodily fluids like serum, plasma, urine and cerebrospinal fluid samples works on photometric colorimetry principle	Class A, Rule 5b	W0201010101	N/A	890619447LM003F3	Y	18/06/2025	

		which is for professional and laboratory environment use only.							
4	LordsMed Fully Auto Biochemistry Analyzer (400 TPH)	Lords Mark Fully Auto Biochemistry Analyzer is used for quantitative analysis of clinical chemistry of specific chemical components in bodily fluids like serum, plasma, urine and cerebrospinal fluidsamples works on photometric colorimetry principle with the minimal use of human assistancewhich is for professional and laboratory environment use only.	Class A, Rule 5b	W0201010102	N/A	890619447LM004F5	Y	18/06/2025	Y
5	LordsMed Fully Auto Biochemistry Analyzer (200 TPH)	Lords Mark Fully Auto Biochemistry Analyzer is used for quantitative analysis of clinical chemistry of specific chemical components in bodily fluids like serum, plasma, urine and cerebrospinal fluidsamples works on photometric colorimetry principle with the minimal use of human assistancewhich is for professional and laboratory environment use only.	Class A, Rule 5b	W0201010101	N/A	890619447LM005F7	Y	18/06/2025	Y

Attachments – DOC11.01.0129 V1 05/03/2025 ANNEX to MDR/IVDR Certificates

6	LordsMed Urine Analyser	The Lordsmed Urine Analyzer is used to analyse the components in urine samples and can be used for semi-quantitative and quantitative detection of biochemical components in urine samples by medical institutions by providing references for clinical references for institutions. Detectable items include white blood cells, urobilinogen, PH, occult blood, specific gravity, ketone bodies, bilirubin, glucose, ascorbic acid, urinary calcium, creatinine and microalbumin concentrations as well as qualitative detection of nitrite concentrations used for point of care, professional and laboratory environment use only.	Class A, Rule 5b	W020101020101	N/A	890619447LM006F9	Y	18/06/2025	Y
7	LordsMed CLIA Reader	LordsMed CLIA Reader a highly sensitive and specific method used in various	Class A, Rule 5b	W020102020101	N/A	890619447LM007FB	Y	18/06/2025	Y

		diagnostic tests for detecting biological molecules like							
8	LordsMed Diluent	LordsMed Diluent is In- Vitro diagnostic reagent, intended for automated dilution of human blood samples, quantitative and qualitative determination of erythrocytes (RBC), leucocytes (WBC) and the leucocyte subpopulations, thrombocytes (PLT) and measurement of hemoglobin (HGB) concentration on Lord Mark hematology analysers for professional and laboratory environment use only.	Class A, Rule 5a	W0103010105	N/A	890619447LM008FD	Y	18/06/2025	Y
9	LordsMed Rinse	LordsMed Rinse is In-Vitro diagnostic reagent, intended for cleaning and rinsing of fluidic path ofLord Mark hematology analysers for professional and laboratory environment use only.	Class A, Rule 5a	W0103010105	N/A	890619447LM009FF	Y	18/06/2025	Y
10	LordsMed Lyse	LordsMed Lyse is In-Vitro diagnostic reagent, intended for	Class A, Rule 5a	W0103010105	N/A	890619447LM010EY	Y	18/06/2025	Y

		automated dilution of human blood samples, quantitative and qualitative determination of erythrocytes (RBC),measurement of hemoglobin (HGB) concentration on Lord Mark hematology analysers for professional and laboratory environment use only.							
11	LordsMed Probe Cleaner	LordsMed Probe Cleaner is In-Vitro diagnostic reagent, intended for cleaning of sample probe of Lord Mark hematology analysers for professional and laboratory environment use only.	Class A, Rule 5a	W0103010105	N/A	890619447LM011F2	Y	18/06/2025	Y

Obelis s.a.
Date: 23/06/2025
Stamp

History log				
Version number	**Version Date**	**Order Form number**	**Reference Number**	**Action**
V1	23/06/2025	EU MD 0257-2025	KP 0683-2025	First issuance

Certificate of Registration

This is to certify that

LORD'S MARK INDUSTRIES PRIVATE LIMITED

HEAD OFFICE ADDRESS: 404, 4TH FLOOR, COSMOS MARY PARK, KOLBAD CROSS LANE, THANE, WEST, 400601, MAHARASHTRA, INDIA

FACTORY: A – 71, ROYAL INDUSTRIAL HUB, KAMAN BHIWANDI ROAD, POMAN VILLAGE, VASAI EAST, PALGHAR, 401208, MAHARASHTRA, INDIA

has been independently assessed by QRO

and is compliant with the requirement of:

ISO 9001:2015

Quality Management System

For the following scope of activities:

DESIGN, DEVELOPMENT, MANUFACTURE AND SUPPLY OF IN-VITRO DIAGNOSTIC BIOCHEMISTRY, HEMATOLOGY, IMMUNOLOGY (RAPID TEST) AND POCT - STRIPS, REAGENTS & KITS. RESALE, INSTALLATION AND SERVICING OF IN-VITRO DIAGNOSTIC ANALYZERS. TRADING & DISTRIBUTION OF IN-VITRO DIAGNOSTIC & MICROBIOLOGY PRODUCTS

Date of Certification: 21st November 2024

1st Surveillance Audit Due: 20th November 2025

2nd Surveillance Audit Due: 20th November 2026

Certificate Expiry: 20th November 2027

Certificate Number: 305024112126Q

CAB # 011905

Head of Certification

Validity of this certificate is subject to annual surveillance audits to be done successfully on or before 365 days from date of the audit.
(In case surveillance audit is not allowed to be conducted; this certificate shall be suspended / withdrawn).

The Validity of this certificate can be verified at www.qrocert.org

This certificate of registration remains the property of QRO Certification LLP, and shall be returned immediately upon request.

India Office : QRO Certification LLP
142, IInd Floor, Avtar Enclave, Near Paschim Vihar West Metro Station, Delhi-110063, (INDIA)
Website : www.qrocert.org, E-mail : info@qrocert.org

Certificate of Registration

This is to certify that

LORD'S MARK INDUSTRIES PRIVATE LIMITED

HEAD OFFICE ADDRESS: 404, 4TH FLOOR, COSMOS MARY PARK, KOLBAD CROSS LANE, THANE, WEST, 400601, MAHARASHTRA, INDIA
FACTORY: A – 71, ROYAL INDUSTRIAL HUB, KAMAN BHIWANDI ROAD, POMAN VILLAGE, VASAI EAST, PALGHAR, 401208, MAHARASHTRA, INDIA

has been independently assessed by QRO

and is compliant with the requirement of:

ISO 14001:2015

Environmental Management System

For the following scope of activities:

DESIGN, DEVELOPMENT, MANUFACTURE AND SUPPLY OF IN-VITRO DIAGNOSTIC BIOCHEMISTRY, HEMATOLOGY, IMMUNOLOGY (RAPID TEST) AND POCT - STRIPS, REAGENTS & KITS. RESALE, INSTALLATION AND SERVICING OF IN-VITRO DIAGNOSTIC ANALYZERS. TRADING & DISTRIBUTION OF IN-VITRO DIAGNOSTIC & MICROBIOLOGY PRODUCTS.

Date of Certification: 8th November 2025
1st Surveillance Audit Due: 7th November 2026
2nd Surveillance Audit Due: 7th November 2027
Certificate Expiry: 7th November 2028

Certificate Number: 305025110825E

CAB # 011905

Head of Certification

Validity of this certificate is subject to annual surveillance audits to be done successfully on or before 365 days from date of the audit.
(In case surveillance audit is not allowed to be conducted; this certificate shall be suspended / withdrawn).
The Validity of this certificate can be verified at www.qrocert.org
This certificate of registration remains the property of QRO Certification LLP, and shall be returned immediately upon request.

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142, IInd Floor, Avtar Enclave, Near Paschim Vihar West Metro Station, Delhi-110063, (INDIA)
Website : www.qrocert.org, E-mail : info@qrocert.org

CERTIFICATE

This is to Certify that

Occupational Health and Safety Management System

of

LORD'S MARK INDUSTRIES LTD.

HEAD OFFICE ADDRESS: 404, 4TH FLOOR, COSMOS MARY PARK, KOLBAD CROSS LANE, THANE (WEST) - 400 601, MAHARASHTRA, INDIA
FACTORY: A-71, ROYAL INDUSTRIAL HUB, KAMAN BHIWANDI ROAD, POMAN VILLAGE, VASAI (EAST), PALGHAR - 401 208, MAHARSHTRA, INDIA

has been independently assessed by DBS
and is compliant with the requirement of:

ISO 45001:2018

For the following scope of activities:

DESIGN, MANUFACTURE, FORMULATION, ASSEMBLING AND MARKETING OF INVITRO DIAGNOSTIC KITS, REAGENTS AND INSTRUMENTS / ANALYSERS AND MICROBIOLOGY PRODUCTS.

Certificate Number: HS-205224082410

Date of Certification:	**24th August 2024**
1st Surveillance Audit Due:	**23rd August 2025**
2nd Surveillance Audit Due:	**23rd August 2026**
Certificate Expiry:	**23rd August 2027**

This Certificate is property of DBS Certifications and remains valid subject to satisfactory surveillance audits

Validity of this certificate may be checked at www.iafcertsearch.org after 7 working days of issue.

Head of Certification

CB-MS-6529

DBS CERTIFICATIONS PVT. LTD.
142, IInd Floor, Avtar Enclave, Paschim Vihar, Delhi-110063, (INDIA) info@dbscertification.com, www.dbscertification.com

ACCREDITED BY :
United Accreditation Foundation Inc. 400 North Center DR STE 202, Norfolk, VA 23502, United States of America

BSCIC

Certificate

MEDICAL DEVICE-
QUALITY MANAGEMENT SYSTEM

This is to certify that:

LORD'S MARK INDUSTRIES PVT. LTD.

A-71, ROYAL INDUSTRIAL HUB, POMAN, KAMAN-BHIWANDI ROAD, VASAI E-401208, MAHARASHTRA, INDIA

Hereby granted the Certificate Number: **BN23622/22504** **Rev. 00**

Subsequent to the **Audit** of the organization, it has been found to be operating a Medical Device-Quality Management System which complies with the requirements of

ISO 13485:2016

For the following scope:

Design, Manufacturing, Distribution and Supply of In-Vitro Diagnostics Reagents, Reagent Products, Calibrators and Control Material for Clinical Chemistry, Immunology (Rapid Test, Elisa), Haematology, Serology & Poct.
Manufacturing, Sales, Installation & Service of Biochemistry, Haematology, Urine, Coagulation Analyzer/Instrument.
Technical Area: Reagents and Reagent Products, Calibrators and Control Materials for: Clinical Chemistry, A.1.4.1, Immunochemistry (Immunology), A.1.4.2, Haematology/ Haemostasis/ Immunohematology, A.1.4.3 In Vitro Diagnostic Instruments and Software, A.1.4.8

Originally Registered:	01-Feb-2025	1st Surveillance Due on:	31-Jan-2026
Issue Date:	01-Feb-2025	2nd Surveillance Due on:	31-Jan-2027
Expiry Date:	31-Jan-2028		

For
BSCIC CERTIFICATIONS PVT.LTD.

Sanjay Seth
Managing Director

(In case if Surveillance Audit is not allowed to be conducted; this Certificate shall be Suspended/Withdrawn).

Please Re-validate this certificate's status at www.bsc-icc.com at REGISTRATION STATUS.
This Certificate of Registration is granted subject to relevant provisions of the BSCIC Certifications PVT. LTD. Contract Terms & Scheme for Registration Form B018 (Latest Version). Please see B018 at our website www.bsc-icc.com.
The certificate of Registration remains the property of BSCIC Certifications Pvt. Ltd. and shall be returned immediately upon request.
BSCIC Headquarters: &work, 1st Floor, 5B/15A, Crown Plaza Mall, Faridabad - 121 007, Haryana, India.

Registered

BSCIC

NABCB
QM 030

MEMBER OF MULTILATERAL RECOGNITION ARRANGEMENT IAF

Certificate of Compliance

This is to certify that

LORD'S MARK INDUSTRIES PRIVATE LIMITED

HEAD OFFICE ADDRESS: 404, 4TH FLOOR, COSMOS MARY PARK, KOLBAD CROSS LANE, THANE, WEST, 400601, MAHARASHTRA, INDIA
FACTORY: A – 71, ROYAL INDUSTRIAL HUB, KAMAN BHIWANDI ROAD, POMAN VILLAGE, VASAI EAST, PALGHAR, 401208, MAHARASHTRA, INDIA

has been assessed and found working Satisfactorily as per the norms of
"Good Manufacturing Practice"
as laid down by World Health Organization

Good Manufacturing Practice (GMP) System

For the following scope of activities:

DESIGN, DEVELOPMENT, MANUFACTURE AND SUPPLY OF IN-VITRO DIAGNOSTIC BIOCHEMISTRY, HEMATOLOGY, IMMUNOLOGY (RAPID TEST) AND POCT - STRIPS, REAGENTS & KITS. RESALE, INSTALLATION AND SERVICING OF IN-VITRO DIAGNOSTIC ANALYZERS. TRADING & DISTRIBUTION OF IN-VITRO DIAGNOSTIC & MICROBIOLOGY PRODUCTS

Date of Certification: 21st November 2024
1st Surveillance Audit Due: 20th November 2025
2nd Surveillance Audit Due: 20th November 2026
Certificate Expiry: 20th November 2027

Certificate Number: 2024112119

Head of Certification

Validity of this certificate is subject to annual surveillance audits to be done successfully on or before 365 days from date of the audit.
(In case if surveillance audit is not allowed to be conducted; this certificate shall be suspended / withdrawn).
The Validity of this certificate can be verified at www.qrocert.com
This certificate of registration remains the property of QRO Certification Limited, and shall be returned immediately upon request.
QRO Certification Limited is accredited by UK Akkreditering Forum Limited, UK (www.ukaf.org.uk)
Winnington House 2 Woodberry Grove Finchley London N12 0DR

QRO Certification Limited
27 Old Gloucester Street, London, WC1N 3AX Company Number : 14230776
Website : www.qrocert.com, E-mail : info@qrocert.com

Certificate of Compliance

Certificate Number: UQ - 2024020710

This is to certify that

LORD'S MARK INDUSTRIES LIMITED

at

HEAD OFFICE: 404, 4TH FLOOR, COSMOS MARY PARK, KOLBAD CROSS LANE, THANE (WEST)-400 601, MAHARASHTRA, INDIA
FACTORY: A-71, ROYAL INDUSTRIAL HUB, KAMAN BHIWANDI ROAD, POMAN VILLAGE, VASAI (EAST) PIN: 401 208. MAHARASHTRA, INDIA

Has successfully implemented the Quality management System and been found working satisfactorily as per the norms of **"Good Manufacturing Practice"** as laid down by "**World Health Organisation"** which has been in conformance to the requirements of

WHO-GMP

DESIGN, MANUFACTURE, FORMULATION, ASSEMBLING AND MARKETING OF INVITRO DIAGNOSTIC KITS, REAGENTS, DNA KITS AND INSTRUMENTS / ANALYSERS

This certificate is issued under the following conditions:

1. It applies only to the quality system maintained in the manufacture of above referenced Models Products.
2. The certificate remains valid until the manufacturing conditions or the quality systems are changed and is subject to continuous surveillance according to the WHO-GMP Guidelines
3. The certificate validity is conditioned by positive results or surveillance audits.

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

Date of Certification	7th February 2024
1st Surveillance Audit Due	6th February 2025
2nd Surveillance Audit Due	6th February 2026
Certificate Expiry	6th February 2027

Authorised Signatory

Certification Services Inc.

VRC

Quality Excellence

Certificate of Compliance

This is to certify the products manufactured by :

LORD'S MARK INDUSTRIES LIMITED

HEAD OFFICE: B-101 RIDHI SIDHI COMPLEX, M.G. ROAD, BORIVALI (E), MUMBAI–400066

FACTORY: A-71, ROYAL INDUSTRIAL HUB, BHIWANDI KAMAN ROAD, POMAN VILLAGE, VASAI (EAST)–401208, MAHARASHTRA, INDIA

The VRC Certification Private Limited , Hereby Certifies that the characteristics of the sample of the products concerned have been found to be Conformity with applicable requirements of: Standard - IEC 61010-1:2010- Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements

for the following product(s):

Semi-Automatic Biochemistry Analyzer
Fully Automatic Biochemistry Analyzer
Urine Test Analyser
3 /5 Part Automated HematologyAnalyzer
Veterinary Hematology Analyser

Certificate No.: VLMO83490926

Date of Issue: 04.04.2025

Date of Expiry: 03.04.2028

1st Surveillance on or before: 03.04.2026

2nd Surveillance on or before: 03.04.2027

Recertification on or before : 03.04.2028

Authorized Signatory

This compliance certificate is approved subject to the organisation maintaining its system, standard, test result as per prescribed standard(s), the test results are exclusive done by orginisation and has no connection to VRCS. This certificate is not transferable and remain valid only above mention address and scope. This certificate is related to the management system, not to the products or services of the certified organizations. This complaince certificate does not provide the certified organization with immunity from its legal obligations. Organisation must check the acceptability of the certificate in requirement with concerened institutions from time to time ,vrcs shall not be responsibile for the same.This certifiificate is not for legal purpose and becomes invalid if yearly surveillance is not done.

TESTED

Certification Services Inc.

VRC

Quality Excellence

Certificate of Compliance

This is to certify the products manufactured by :

LORD'S MARK INDUSTRIES LIMITED

HEAD OFFICE: B-101 RIDHI SIDHI COMPLEX, M.G. ROAD,
BORIVALI (E), MUMBAI–400066
FACTORY: A-71, ROYAL INDUSTRIAL HUB, BHIWANDI KAMAN ROAD,
POMAN VILLAGE, VASAI (EAST)–401208, MAHARASHTRA, INDIA

The VRC Certification Private Limited , Hereby Certifies that the characteristics of the sample of the products concerned have been found to be Conformity with applicable requirements of: Standard -IEC 61010-2-081: 2019-Safety requirements for electrical equipment for measurement, control and laboratory use - Part 2-081: Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes

for the following product(s):

Semi-Automatic Biochemistry Analyzer
Fully Automatic Biochemistry Analyzer
Urine Test Analyser
3 /5 Part Automated HematologyAnalyzer
Veterinary Hematology Analyser

Certificate No.: VLMO427360933
Date of Expiry: 03.04.2028
1st Surveillance on or before: 03.04.2026
2nd Surveillance on or before: 03.04.2027
Recertification on or before : 03.04.2028

Date of Issue: 04.04.2025

Authorized Signatory

TESTED

Certification Services Inc.

VRC

Quality Excellence

Certificate of Compliance

This is to certify the products manufactured by :

LORD'S MARK INDUSTRIES LIMITED

HEAD OFFICE: B-101 RIDHI SIDHI COMPLEX, M.G. ROAD, BORIVALI (E), MUMBAI–400066

FACTORY: A-71, ROYAL INDUSTRIAL HUB, BHIWANDI KAMAN ROAD, POMAN VILLAGE, VASAI (EAST)–401208, MAHARASHTRA, INDIA

The VRC Certification Private Limited , Hereby Certifies that the characteristics of the sample of the products concerned have been found to be Conformity with applicable requirements of: Standard - IEC 61326-1:2020 -Electrical equipment for measurement, control and laboratory use - EMC requirements - Part 1: General requirements

for the following product(s):

Semi-Automatic Biochemistry Analyzer
Fully Automatic Biochemistry Analyzer
Urine Test Analyser
3 /5 Part Automated Hematology Analyzer
Veterinary Hematology Analyser

Certificate No.: VLMO509240943

Date of Issue: 04.04.2025

Date of Expiry: 03.04.2028

1st Surveillance on or before: 03.04.2026

2nd Surveillance on or before: 03.04.2027

Recertification on or before : 03.04.2028

Authorized Signatory

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VRC Certification Pvt.Ltd
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LORD'S MARK INDUSTRIES LIMITED

HEAD OFFICE: B-101 RIDHI SIDHI COMPLEX, M.G. ROAD, BORIVALI (E), MUMBAI–400066

FACTORY: A-71, ROYAL INDUSTRIAL HUB, BHIWANDI KAMAN ROAD, POMAN VILLAGE, VASAI (EAST)–401208, MAHARASHTRA, INDIA

The VRC Certification Private Limited , Hereby Certifies that the characteristics of the sample of the products concerned have been found to be Conformity with applicable requirements of: Standard - IEC 60601-1:2005+AMD1:2012+AMD2:2020 CSV, Medical electrical equipment - Part 1: General requirements for basic safety and essential performance

for the following product(s):

Semi-Automatic Biochemistry Analyzer, Fully Automatic Biochemistry Analyzer, Urine Test Analyzer, 3/5 Part Automated Hernatology Analyzer, Veterinary Hematology Analyzer, Immunoassay Analyzer, Electrolyte Analyzer, Hemodialysis Machine

Certificate No.: VLMCE76321828

Date of Expiry: 02.02.2029

1st Surveillance on or before: 02.02.2027

2nd Surveillance on or before: 02.02.2028

Recertification on or before : 02.02.2029

Date of Issue: 03.02.2026

Authorized Signatory

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#5A, 2nd Floor, Laxmanpuri, Lucknow-226016, U.P, India, www.vrccertification.in, vrcservices@gmail.com, info@vrccertification.in