CERTIFICATE

EUROGLOBAL
CERTIFICATIONS (UK) LTD.

Certificate of Compliance

We hereby declare that the technical file of

MAXSOL INSTRUMENTS

OFFICE ADDRESS- B-112, RAJESHREE INDUSTRIAL ESTATE, AGARWAL NAKA, SATIVALI ROAD, NEAR MSEB, VASAI EAST, PALGHAR-401208 (INDIA)
FACTORY ADDRESS- A-10, DARSHAK INDL. EST., VASAI PHATA, NEAR UMANG PHARMA, VASAI (EAST), DIST. PALGHAR-401208 (INDIA)

has been assessed & found to be in conformance with the provisions set forth by the requirement of Directive Low Voltage Directive 2014/35/EU.

PRODUCT DESCRIPTION: MANUFACTURER OF BLOOD BANK AND LABORATORIES MACHINERIES AND SPARES.

(More Details as per Appendix-I & II)

The Certification body has performed a sample audit of the above product quality system covering the design, manufacture & final inspection of the certified product(s). The quality system has been assessed, approved and is subject to continuous surveillance according to Directive Low Voltage Directive 2014/35/EU. No additional test report was carried out from submitted type sample of the product in Compliance with the Specification of the respective standards except those submitted by the Customer.

Certificate No.	**EU7502**		
Date of Initial Registration	04th July. 2023	*Issue Date*	04th July. 2023
1st Surveillance on or before	04th July. 2024	*Expiry Date*	03rd July. 2026
2nd Surveillance on or before	04th July. 2025		

Authorized Signatory

Euro Global Certifications UK Limited ★ EurO Global

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CERTIFIED FOR COMPLIANCE EUROGLOBAL CERTIFICATIONS

ISSUED BY THE Euroglobal CERTIFICATIONS (UK) LTD. CERTIFICATION BOARD

CERTIFICATE

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Certificate of Compliance Appendix-I to Certificate No.: EU7502

This Appendix shall be an integral part of the Certificate. All expressions and terms defined or used in the Certificate shall have the same meaning in this Addendum, unless the context clearly requires otherwise.

MANUFACTURER	:	MAXSOL INSTRUMENTS
PRODUCT GROUP	:	DEVICES
BRAND NAME	:	MAXOL

This certificate referred to above covers the following products:

- REFRIGERATED CENTRIFUGE (TABLE TOP)
- NON-REF CENTRIFUGE (TABLE TOP)
- BLOOD BANK CENTRIFUGE
- GEL CARD CENTRIFUGE
- BLOOD BANK REFRIGERATOR
- TUBE SEALER
- BLOOD COLLECTION MONITOR
- PORTABLE DONOR COACH
- DONOR COACH
- LAMINAR FLOW
- THAWING BATH
- PLASMA EXPRESSER
- SINGLE PAN BALANCE
- DOUBLE PAN BALANCE
- BLOOD BANK STRIPER
- PLATELET AGITATOR

The CE mark as shown can be used, under the responsibility of the manufacturer, after completion of an EC Declaration & Compliance with all relevant EC Directives. The statement is based on a single evaluation of one sample of above mentioned product(s). This certificate is issued under the conditions that the quality system maintained in the manufacturing of above referenced Models/Products & remains valid until the manufacturing conditions or the quality systems are changed subject to continuous surveillance according to the EC Guidelines. Further, Certificate is conditioned by positive results of surveillance audits.

Authorized Signatory

ISSUED BY THE Euroglobal CERTIFICATIONS (UK) LTD. CERTIFICATION BOARD

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Certificate of Compliance Appendix-II to Certificate No.: EU7502

This Appendix shall be an integral part of the Certificate. All expressions and terms defined or used in the Certificate shall have the same meaning in this Addendum, unless the context clearly requires otherwise.

MANUFACTURER	:	MAXSOL INSTRUMENTS
PRODUCT GROUP	:	DEVICES
BRAND NAME	:	MAXOL

This certificate referred to above covers the following products:

- PLATELET INCUBATOR
- WATER BATH
- WATER BATH SHAKER
- VDRL SHAKER
- ORBITAL SHAKER
- INCUBATOR SHAKER
- TUBE WARMER
- MICROSCOPE
- HOT AIR OVEN
- AUTOCLAVE AND STERILIZERS
- RH VIEW BOX
- CMS
- STABILITY TEST CHAMBERS
- PHOTO STABILITY TEST CHAMBERS
- BOD INCUBATOR
- PLASMA FREEZER (-40°C)
- ULTRA PLASMA FREEZER (-86°C)
- BIO FREEZER
- BIO SAFETY CABINET

The CE mark as shown can be used, under the responsibility of the manufacturer, after completion of an EC Declaration & Compliance with all relevant EC Directives. The statement is based on a single evaluation of one sample of above mentioned product(s). This certificate is issued under the conditions that the quality system maintained in the manufacturing of above referenced Models/Products & remains valid until the manufacturing conditions or the quality systems are changed subject to continuous surveillance according to the EC Guidelines. Further, Certificate is conditioned by positive results of surveillance audits.

Authorized Signatory

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