



		CALIB	RATIO	V CERT	IFICAT	E		
CERTIFICATE NO: SBS/CL/22/06703				MEDICAL DEVICES			Page No:1 of 1	
	ssue Date			25-07-2022			rage No.1 of 1	
SRF No & Date				SRF/22/0143-015 & 20-07-2022				
Receipt Date				20-07-2022				
Calibration Date				20-07-2022				
Calibration Due				19-07-2023				
	omer Name & Address			. 2020				
30V	ERNMENT PRIMARY HEA	LTH CENTRE,						
AND	HANALLLUR, TRICHY(DT	-639101.						
		Deta	ils of Device	Under Calibra	tion (DUC)			
Description : SEMI AUTO ANALYZER				Make & Model : ROBONIK & PRIETEST TOUCH				
Range				Sr. No : ATCD2680818RBK				
	ution :		Iden	tification No				
DUC Condition : SATISFACTORY				Location : LAB				
		Environment	al Conditions	& Calibration	Procedure De	etails		
Environmental Details Temperature:25.3 ° C			°C	Relative Humidity		49% RH		
Calibi	ration Procedure No	SBS/CP/MD/29		Calibration done at		ONSITE		
				Standards De	ails			
S.No	Description Make/ SI		ce/ SI No:	Certificate N		0	Validity	
	Electrical Safety Analyser Rigel Medica			3 44L-1059 M-210809-17-3			10-08-202	
RES	JLTS							
S.no	Specification	Measured values in Ω		Allowable limit in Ω		Uncertainty in Ω (±)		
1	Earth Resistance	Earth Resistance 0.099		<0.30		0.83		
	Measured values in		lues in µA	Allowable limit in µA		Uncertainty in % (±)		
2	Earth Leakage	101		<5000µAfor B,BF,CF		5.9		
	Measured values in		lues in µA	Allowable limit in µA		Uncertainty in µA (±)		
3	Patient Leakage	5		<100µAfor B,BF,<10µAfor CF		2.9		
		Measured values in µ		Allowable limit in µA		Uncertainty in % (±)		

Enclosure Leakage REMARKS

215 1. This Calibration certificate shall not be reproduced except in full, without written approval of the laboratory.
2. The user should determine the suitability of the instrument for its intended use.
3. The recalibration interval should be determined on the user requirement.

- 4. The results stated in this certificate relate only to the item calibrated.

 5. The indicated uncertainties are expanded uncertainty estimated for a confidence level of approximately 95% for a coverage factor k=2.00.

 6. Equipment used for Calibration were calibrated & traceable to National & International Standards.

Authorised Signatory Calibrated By

(Calibration Engineer)
C.SIVABALAN





<500µAfor B,BF,CF



5.9

SUNSHINE BIOMEDICAL SOLUTIONS

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