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REPORT ON ELECTRICAL SAFETY TESTING/ PERFORMANCE ANALYSIS/ CALIBRATION

Report No: TR/PUPHC/013/22-23

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1.1 CUSTOMER DETAILS

Reference and Date

Name and address of the organisation

UPHC PERIYASEMUR

Near panchayat office, Periyasemur

Calibration Date: 15-08-2022

Calibration Due: 14-08-2023

Tamil Nadu - 638004 Letter dated 14-08-2022

Date of receipt of item 14-08-2022

1.2 DESCRIPTION OF DEVICE UNDER TEST (DUT)

Nomenclature CENTRIFUGE

REMI A. Manufactured by R8C B. Model

PUPHC002 C. Serial No.

PUPHC/LAB/CENT/02 **Biomedical Product ID** D. 220-230 V AC, 50/60 Hz E. Supply Damped sinusoidal F. **Wave Form**

Type CF **Device Type** G.

Device Classification Class I equipment H. LABORATORY Location

1.3 CONDITION OF THE ITEM WHEN RECEIVED

No visible damage and in working order

1.4 ENVIRONMENTAL CONDITION OF MEASUREMENTS

32.4°C A. Temperature **B.** Relative Humidity 45-60% C. Ambient Barometric Pressure 753mmHg

IEC Specification IEC 60601-1,IEC 60601-2-4 1.5 Applicable Specification **Electrical Safety and Perfomance Testing** 1.6 Test Done

Tested BY:

Balamuralikrishnan K

Approved by:

Priya M

(Quality Managek)

FL BIOMEDICAL 189, Vasantham Paradise, Chithode, Erode-638102.

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1.7 STATUS

	factures fication	Users Specification	Within Specificatio	Out o		libration	Electrical Safety Test	Performance Analysis
1.8 TR/	ACEABILITY	Y DETAILS OF INS	TRUMENTS	USED FOR T	ESTING			
SI No		the Instrument	Make	Model	Serial	No	Cal Due	Traceability Reference
1.		l Tachometer I Safety Analyzer	WACO RGM	DT-2234C 288+	1804160 05H-06		JULY 2023 JULY 2023	Annexure 3 Annexure 1
1.9 PE	RFORMAN	ICE ANALYSIS OF	CENTRIFUGE					
Rpm / SI No	Speed O	utput In RPM			ons in DUT		wed Deviation ge (RPM)	Remarks

	-						~				
-	-	-	N. P.	-	10	A 1		C A	ECTY	<i>/</i> TI	25.1

SET VALUE IN

1000

2500

3000

DUT(J)

Z.U ELEL	TRICAL SAFETT TEST		Acceptable limits as per the Std.	Remarks
S.no	Parameter	Observed value	Acceptable liffits as per the star	
	Protective earth resistance	0.122 Ω	< 0.3 Ω	ok
1.		130.2 μΑ	< 5K μΑ NC	ok
2.	Earth leakage	7.8 µA	< 100 μA BF	ok
3.	Patient leakage current	0.3 A	As per manufacture spec	ok
4.	Equipment current		As per manufacture spec	ok
5.	Mains voltage	231.2 V	As per manajacture spec	

23

70

51

73

3.0 REMARKS

- This report is applicable to the sample tested only. 3.1
- The instruments used for testing are under valid calibration and are traceable to National Standards.
- Parameter of the DUT were verified and found to be within the specified limits. 3.3

MEASURED VALUE

1023

2570

3051

3573

IN DUT(J)

refer NABL Doc No. 121 Clause 7.0 Accommodation and environmental Conditions sub Clause see 3.4 7.2.11 below in line with ISO/IEC 17025:2005 Clause 5.3

Tested BY:

Balamuralikrishnan K



Approved by:

Priya M (Quality Manager) FL BIOMEDICAL

189, Vasantham Paradise, Chithode, Erode-638102.

PASS

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