

## Test Report issued under the responsibility of:

NCB Intertek Testing Services, NA, Inc.

## **TEST REPORT 102945703LAX-015**

IEC 60601-1-2 (ed.4) 2014-02-25

Date of issue...... 2017-03-10

Name of Testing Laboratory

preparing the Report ...... Intertek Testing Services NA, Inc.

Address...... 25800 Commercentre Drive

Lake Forest, CA 92630 USA

Applicant's name ...... Med-Logics, Inc.

Athens, TX 75751 USA

Test specification:

Standard ...... IEC 60601-1-2 (ed.4) 2014-02-25

Test procedure...... CB Scheme

Non-standard test method............ N/A
National Differences/Deviations....... N/A

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The test results presented in this report relate only to the object tested.

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Total Maria Depositation Contents	
Test Item Description:	Cataruise System
Trademark:	CataPulse®
Manufacturer:	Same as the applicant
Model/Type Reference:	CP1700 (CataPulse Console),
	CP1800 (CataPulse Footswitch),
	CP1900 (CataPulse Cart)
Ratings:	100 – 240 VAC, 5 A, 50/60 Hz

Responsible Testing Laboratory (as applicable), testing procedure and testing location(s):

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	CB Testing Laboratory	Intertek Testing Services NA, Inc.
Testing location/address		25800 Commercentre Drive
		Lake Forest, CA 92630 USA
Tested by (name, function, signature):		Julio Lopez, Project Handler,
Approved by(name, function, signature):		Julio Lopez, Project Handler,  Melvin Sanchez, Reviewer,
	Testing Procedure: CTF Stage 1	N/A
Testing location/address		
Tested by (name, function, signature):		
Appro	oved by(name, function, signature):	
	Testing Procedure: CTF Stage 2	N/A
Testing location/address:		
Tested by (name, function, signature):		
Witnessed by (name, function, signature):		
Approved by(name, function, signature):		
	Testing Procedure: CTF Stage 3	N/A
	Testing Procedure: CTF Stage 4	N/A
Testing location/address:		
Tested by (name, function, signature):		
Witnessed by (name, function, signature):		
Approved by(name, function, signature):		
Supe	rvised by:	
Manu	ufacturer's Declaration per sub-claus	e 4.2.5 of IECEE 02:
The application for obtaining a CB Test Certificate includes more than one factory location and a		
declaration from the Manufacturer stating that the sample(s) submitted for evaluation is (are)		
representative of the products from each factory has been provided ☐ Yes ☒ N/A		
When differences exist; they shall be identified in the full test report.		
Name and address of factory (ies): Same as the applicant		
General Product Information:		