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AUTHORIZATION TO MARK

This authorizes the application of the Certification Mark(s) shown below to the models described in the Product(s) Covered section when made in accordance with the conditions set forth in the Certification Agreement and Listing Report. This authorization also applies to multiple listee model(s) identified on the correlation page of the Listing Report.

This document is the property of Intertek Testing Services and is not transferable. The certification mark(s) may be applied only at the location of the Party Authorized To Apply Mark.

Applicant:	PEMF Systems, Inc.	**Manufacturer:**	Vesper Manufacturing Inc.
Address:	422 Kirkstone Way Las Vegas, NV 89123	**Address:**	14556 Weddington St Sherman Oaks, CA 91411
Country:	USA	**Country:**	USA
Contact:	Maria Carson, Tim Duggins	**Contact:**	Dane Davis, Tim Duggins
Phone:	818 267 0095, 602 200 4981	**Phone:**	818 267 0095, 602 200 4981
FAX:	NA	**FAX:**	NA
Email:	maria@pemfsystems.com timd@horizonphoenix.biz	**Email:**	dane@pemfsystems.com, timd@horizonphoenix.biz

Party Authorized To Apply Mark: Same as Manufacturer
Report Issuing Office: Lake Forest, CA

Control Number: 5010666 **Authorized by:** Carolyn Mattis
for Dean Davidson, Certification Manager

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This document supersedes all previous Authorizations to Mark for the noted Report Number.

Intertek Testing Services NA Inc.
545 East Algonquin Road, Arlington Heights, IL 60005
Telephone 800-345-3851 or 847-439-5667 Fax 312-283-1672

ATM Issued: 11/30/2018

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AUTHORIZATION TO MARK

Standard(s):	Medical Electrical Equipment - Part 1: General Requirements For Basic Safety And Essential Performance (R2012) [AAMI ES60601-1:2005 +C1;A2] Medical Electrical Equipment - Part 1: General Requirements For Basic Safety And Essential Performance (R2013) [CSA C22.2#60601-1:2008 Ed.2 +C2] Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability [IEC 60601-1-6:2010 Ed.3] Medical Devices - Application Of Usability Engineering To Medical Devices [IEC 62366:2007 Ed.1] Medical Elec. Equip.- Part 1-11: Gen. Req. For Basic Safety & Essential Perf.- Collateral Standard - Req. For Medical Elec. Equip. & Medical Elec. Systems Used In The Home Healthcare Environment; Corr. 1: 2011 [IEC 60601-1-11:2010 Ed.1 +C1]
Product:	Pulsed Electro Magnetic Fields (PEMF) Stimulation and Healing
Models:	High Power, Medium Power, Low Power

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