



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: WR Medical Electronics Co.
1700 Gervais Ave
Address: Maplewood, MN 55109
Country: USA
Contact: Mr. Kyle Maloney
Phone: (651) 604-8451
FAX: (651) 604-8499
Email: kam@wrmed.com

Manufacturer: WR Medical Electronics Co.
1700 Gervais Ave
Address: Maplewood, MN 55109
Country: USA
Contact: Mr. Kyle Maloney
Phone: (651) 604-8451
FAX: (651) 604-8499
Email: kam@wrmed.com

Party Authorized To Apply Mark: Same as Manufacturer
Report Issuing Office: Oakdale, MN

Control Number: 3025752

Authorized by: *for Michelle Lake*
William T. Starr, Certification Manager



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

This Authorization to Mark is for the exclusive use of Intertek's Client and is provided pursuant to the Certification agreement between Intertek and its Client. Intertek's responsibility and liability are limited to the terms and conditions of the agreement. Intertek assumes no liability to any party, other than to the Client in accordance with the agreement, for any loss, expense or damage occasioned by the use of this Authorization to Mark. Only the Client is authorized to permit copying or distribution of this Authorization to Mark and then only in its entirety. Use of Intertek's Certification mark is restricted to the conditions laid out in the agreement and in this Authorization to Mark. Any further use of the Intertek name for the sale or advertisement of the tested material, product or service must first be approved in writing by Intertek. Initial Factory Assessments and Follow up Services are for the purpose of assuring appropriate usage of the Certification mark in accordance with the agreement, they are not for the purposes of production quality control and do not relieve the Client of their obligations in this respect.

Intertek Testing Services NA Inc.
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Standard(s):	Standard for Medical Electrical Equipment, Part 1: General Requirements for Safety, UL 60601-1, 2003 First Edition, including Revision 2006 CAN/CSA C22.2 No. 601.1-M90 including update No. 2, November 2003, Reaffirmed 2005
Product:	Facial Nerve Stimulator
Brand Name:	Hilger
Models:	H3