



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.

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Control Number: 3131439 Authorized by: Mikael Goffné
Mikael Goffné for William T. Starr, Certification Manager

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

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Applicant:	<u>DisMark GmbH</u>	Manufacturer:	<u>Same as Applicant</u>
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Party Authorized To Apply Mark: Same as Manufacturer
Report Issuing Office: Intertek Kaufbeuren, Germany

Standard(s):	Medical Electrical Equipment, Part 1: General Requirements for Safety: UL60601-1, Issue: 2003-04-25 Ed.:1 Rev: 2006/04/26; CAN/CSA C22.2 NO 601.1-M90, Issue:1990/01/01 Rev: 1998/02; Supplement 1; 1994; Amendment 2 - February 1998; Update No. 2 (R2001) Part 1-2: General Requirements for Safety - Collateral Standard Electromagnetic Compatibility - Requirements and Tests: CAN/CSA-C22.2 NO. 60601-1-2-03; Amendment 1:2006; IEC 60601-1-2, Issue:2001/09/01 Ed: 2 Safety of Laser Products - Part 1: Equipment Classification, Requirements and User's Guide: CAN/CSA-E60825-1-03; IEC 60825-1, Ed. 1.2
Product :	Model MedicLaser is a Class 3R laser device for treatment of a variety of diseases, for example acne, allergies, arthritis, etc. It is part of a complete delivery package with the model designation 'EarLaser' (consisting of Model MedicLaser and Model EarTool, a head bracket for fixing the laser device to the ear), which is used for treatment of inner ear diseases (i.e. tinnitus).
Models:	<u>MedicLaser</u>

CRH