



# AUTHORIZATION TO MARK

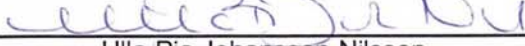
This authorizes the application of the Certification Mark(s) shown below to the models described in the Product(s) Covered section when made in accordance with the conditions set forth in the Certification Agreement and Listing Report. This authorization also applies to multiple listee model(s) identified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

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**Applicant:** Sa.Vir Spa  
**Address:** Via G.Galilei, 22  
IT-46040 Guidizzolo (MN)  
**Country:** ITALY  
**Contact:** Mr. Cristiano Michelazzi  
**Phone:** +39 0376 840404  
**FAX:** +39 0376 840419  
**Email:** cristiano.michelazzi@savir.it

**Manufacturer:** Eurohm srl  
**Address:** Via Volta, 3  
IT-20054 Nova Milanese (MI)  
**Country:** ITALY  
**Contact:** Mr. Claudio Volpi  
**Phone:** +39036240901  
**FAX:** +390362450818  
**Email:** volpi@eurohm.it

**Party Authorized To Apply Mark:** Same as Manufacturer  
**Report Issuing Office:** Intertek Semko AB, Stockholm, Sweden

**Control Number:** 4000568      **Authorized by:**   
Ulla-Pia Johansson-Nilsson  
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.  
165 Main Street, Cortland, NY 13045  
Telephone 800-345-3851 or 607-753-6711 Fax 607-756-6699

<b>Standard(s):</b>	UL 60601-1 Issued: 2003-04-25 Ed.:1 Rev: 2006/04/26 Standard for Safety Medical Electrical Equipment, Part 1: General Requirements for Safety; CSA C22.2#601.1 Issued: 1990/01/01 Medical Electrical Equipment - Part 1: General Requirements for Safety General Instruction No 1; Supplement 1; 1994; Amendment 2 - February 1998; Update No. 2 (R2001); IEC 60601-1-2 Issued: 2007: Medical electrical equipment Part 1-2: General requirements for safety - Collateral standard: Electromagnetic compatibility - Requirements and tests IEC 60601-1-4 2000/04/01 Ed:1 Medical Electrical Equipment - Part 1-4: General Requirements for Safety - Collateral Standard: Programmable Electrical Medical Systems Consolidated with Amd. 1:1999
<b>Product:</b>	Patient transfer system
<b>Models:</b>	MOBILIZER_3

*MMP*

*ES*