

ELECTRICAL LISTING PROGRAM

Customer: Rostrum Medical Innovation Inc.
 Class: Medical Electrical Equipment
 Location: Vancouver, B.C. Canada
 Website: www.rostrummedical.com
 Product Site: N/A
 Listing No. E10581
 Report No. E10581-2301
 Project No. E10581-2301
 Effective Date: September 12, 2022
 Last Revised Date: February 28, 2023
 Expires: N/A
 Standards: AAMI ES60601-1
 CAN/CSA-C22.2 No. 60601-1:14
 Product: The VQm Pulmonary Health Monitor (PHM), Medical Electrical Equipment, Class I, Cord-set connected, Mobile
 Markings: Products are marked in a permanent manner where it is readily visible after installation with the following:

- a) Manufacturer's name or trademark
- b) Model designation
- c) Month and year of manufacture
- d) Product ratings: voltage, frequency, amps
- e) QAI file Number: E10581
- f) The wording "Conforms with CAN/CSA 60601-1 and AAMI ES60601-1"
- g) Warning markings as required by the standards in English and French
- h) QAI logo shown here:



Models / Ratings:

Model	Rating	Description	Environmental Locations
VQm-S06	100-240V~, 50/60Hz, 2A	Pulmonary Health Monitor (PHM)	Dry Location only

- Notes:
1. Requirements for biocompatibility (Subclause 11.7) and electromagnetic compatibility (Clause 17) are not evaluated (N/E) in this evaluation. Consideration shall be made by the Manufacturer in the end use.
 2. Hazards resulting from intended physiological effects or from the abnormal use of the Product are not considered in this evaluation.
 3. The Manufacturer's risk management file for the Product is reviewed in accordance with the requirements of ISO 14971 as part of this evaluation. The Manufacturer's risk management decisions inform the applicable test and evaluation requirements. The Manufacturer shall notify QAI of any changes or revisions in the risk management file that affect the compliance of the Product with the applicable Standards.
 4. If this Product is subsequently combined with other equipment through a functional connection or by use of a multiple socket-outlet, consideration of the requirements of Clause 16 for Medical Electrical Systems shall be made by the Manufacturer or the Responsible Organization in the end use.

The materials, products or systems listed herein have been qualified to bear the QAI Listing Mark under the conditions stated with each Listing. Only those products bearing the QAI Listing Mark are considered to be listed by QAI. No warrantee is expressed or implied, and no guarantee is provided that any jurisdictional authority will accept the Listing found herein. The appropriate authorities should be contacted regarding the acceptability of any given Listing. Visit the QAI Online Listing Directory located at www.qai.org for the most up to date version of this Listing and to validate that this QAI Listing is active. Questions regarding this listing may be directed to info@qai.org. Please include the listing number in the request.