Reprocessed by

Sustainability Solutions

Instructions for Use
Reprocessed ECG 5-Lead Grabber

For Use with Philips IntelliVue MX40

Reprocessed Device for Single Use

Caution: Federal (U.S.A.) law restricts this device to sale by or on the order of a physician.

- **NON-STERILE (Exposed to EO gas)**
- **Not made with natural rubber latex**

Explanation of Symbols


Date of Reprocessing

Batch Code

Catalog Number


Do Not Reuse


Consult Instructions for use

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Reprocessed Philips ECG 5-Lead Grabber

Product Overview

Indications for Use
The Reprocessed ECG 5-Lead Grabber are intended to provide ambulatory and bedside monitoring of ECG/Impedance Respiration, for adults who weigh greater than 40kg and pediatrics who weigh greater than 20kg. The Reprocessed ECG 5-Lead Grabber are for use with IntelliVue MX40 patient wearable monitors, and are intended to be used by trained healthcare personnel in a professional healthcare facility.

Cable Configurations
5-Lead* ECG Only

REF	**ECG Color Coding**
989803172031	AAMI color coded grabber connectors

**All ECG leads are 0.85m (34in) long.*

Warnings
- For Single-patient use only. Do not use on multiple patients. Multi-patient use may result in patient infection or cross-infection.
- Do not use device in magnetic resonance imaging (MRI) environments or during MRI procedures.
- Ensure that lead sets are carefully positioned to avoid entanglement, choking, strangulation, or inhibit circulation in extremities.
- Protect the patient cable trunk connector and ECG lead connectors from becoming soaked with any liquid.

Precautions
- Inspect the device prior to use. Do not use if visual inspection reveals damage from liquid, lint or other contaminants.
- Ensure that lead set is fully and properly inserted into trunk cable or instrument.
- Do NOT use lead set in excessively wet environments or under massive influence of fluids (e.g. rain).

Directions for Use
Visually inspect the reprocessed ECG 5-Lead Grabber before applying the patient cable to a patient.
- If any signs of damage or deterioration are seen, immediately discard the device following approved medical waste disposal methods specified by local regulations or your patient care facility.
- If not visual damage is seen, apply the ECG leads to the patient following standard AAMI practices (refer to the *Philips IntelliVue MX40 Instructions for Use).*

Additional Information

For more detailed information describing patient cable use with the MX40, including warnings, patient safety, and ECG patient application instructions, refer to the *Philips IntelliVue MX40 Instructions for Use.*

Reprocessed Philips ECG 5-Lead Grabber

Warranty

Reprocessed Products

Stryker warrants all reprocessed products, subject to the exceptions provided herein, to be free from defects in reprocessing and to substantially conform to the product specifications contained in the documentation provided by Stryker with the products for one use in accordance with the instructions for use of such product.

STRYKER SHALL NOT BE LIABLE FOR ANY DAMAGES TO THE EXTENT CAUSED BY ANY DEFECT IN MATERIAL, WORKMANSHIP OR DESIGN BY THE ORIGINAL MANUFACTURER OF THE PRODUCT OR ANY ACT OR OMISSION OF THE ORIGINAL MANUFACTURER OF THE PRODUCT.

Products for which Stryker is the Original Manufacturer

Stryker warrants all products for which it is the original manufacturer, subject to the exceptions provided herein, to be free from defects in design, materials and workmanship and to substantially conform to the product specifications contained in the documentation provided by Stryker with the products for a period of one year from the date of purchase.

General Warranty Terms Applicable to All Products

TO THE FULLEST EXTENT PERMITTED BY LAW, THE EXPRESS WARRANTY SET FORTH HEREIN IS THE ONLY WARRANTY APPLICABLE TO THE PRODUCTS AND IS EXPRESSLY IN LIEU OF ANY OTHER WARRANTY BY STRYKER, EXPRESSED OR IMPLIED, INCLUDING, BUT NOT LIMITED TO, ANY IMPLIED WARRANTY OR MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. IN NO EVENT WILL STRYKER'S LIABILITY ARISING IN CONNECTION WITH THE SALE OF THE PRODUCT (WHETHER UNDER THE THEORIES OF BREACH OF CONTRACT, TORT, MISREPRESENTATION, FRAUD, WARRANTY, NEGLIGENCE, STRICT LIABILITY OR ANY OTHER THEORY OF LAW) EXCEED THE PURCHASE PRICE, CURRENT MARKET VALUE OR RESIDUAL VALUE OF THE PRODUCTS, WHICHEVER IS LESS. STRYKER SHALL NOT BE LIABLE FOR INDIRECT, SPECIAL, INCIDENTAL, PUNITIVE OR CONSEQUENTIAL DAMAGES RESULTING FROM ANY BREACH OF WARRANTY OR UNDER ANY OTHER LEGAL THEORY.

This warranty shall apply only to the original end-user purchaser of products directly from Stryker or a Stryker authorized distributor. This warranty may not be transferred or assigned without the express written consent of Stryker.

This warranty does not apply to: (1) products that have been misused, neglected, modified, altered, adjusted, tampered with, improperly installed or refurbished; (2) products that have been repaired by any person other than Stryker personnel without the prior written consent of Stryker; (3) products that have been subjected to unusual stress or have not been maintained in accordance with the instructions in the user manual or as demonstrated by a Stryker representative; (4) products on which any original serial numbers or other identification marks have been removed or destroyed; or (5) products that have been repaired with any unauthorized or non-Stryker components.

If a valid warranty claim is received within thirty (30) days of the expiration of the applicable warranty period, Stryker will, in its sole discretion: (1) replace the product at no charge with a product that is at least functionally equivalent to the original product or (2) refund the purchase price of the product. If a refund is provided by Stryker, the product for which the refund is provided must be returned to Stryker and will become Stryker's property. In any event, Stryker's liability for breach of warranty shall be limited to the replacement value of the defective or non-conforming part or component.

If Stryker determines in its reasonable discretion that the claimed defect or non-conformance in the product is excluded from warranty coverage as described hereunder, it will notify the customer of such determination and will provide an estimate of the cost of repair of the product. In such an event, any repair would be performed at Stryker's standard rates.

Products and product components repaired or replaced under this warranty continue to be warranted as described herein during the initial applicable warranty period or, if the initial warranty period has expired by the time the product is repaired or replaced, for thirty (30) days after delivery of the repaired or replaced product. When a product or component is replaced, the item provided in replacement will be the customer's property and the replaced item will be Stryker's property. If a refund is provided by Stryker, the product for which the refund is provided must be returned to Stryker and will become Stryker's property.

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The OEM information listed on the label is provided as device ID prior to reprocessing and may contain the trademarks of unrelated third parties that do not sponsor this device.

This product and its packaging have been exposed to ethylene oxide gas (EtO). Even though the product then is processed in compliance with all applicable laws and regulations relating to EtO exposure, Proposition 65, a State of California voter initiative, requires the following notice:

Warning: This product and its packaging have been exposed to ethylene oxide. The packaging may expose you to ethylene oxide, a chemical known to the State of California to cause cancer or birth defects or other reproductive harm.

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