Reprocessed by
ReNu Medical, Inc.

Instructions for Use
Reprocessed Blood Pressure Cuff

Reprocessed Device for Single Use

Symbol Legend:

	Date of Reprocessing		Reprocessor/Manufacturer
	Do Not Reuse		Not made with natural rubber Latex
	Consult instructions for use		Fragile, handle with care
	Do not use if package is damaged		Keep dry
	Non-Sterile – High Level Disinfection	Rx Only	Caution: Federal (USA) law restricts this device to sale by or on the order of a physician.
	Caution See Instructions for use	OEM	Original Equipment Manufacturer Catalog#
REF	ReNu Medical Catalog#	LOT	ReNu Medical Sales Order
CID	Customer ID#, if none specified; ReNu Medical Catalog#	QTY	Qty of Devices included in Pkg/Cs

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Reprocessed Blood Pressure Cuff

Indications for Use

Clinical applications for the Blood Pressure Cuffs are as follows: Used to monitor the vital signs of the patient

Contraindications

The Cuff should NOT be used in the following conditions:
Any local condition in which the garment would interfere:

- open wounds
- untreated infected wounds
- gangrene
- recent skin graft
- dermatitis

Warnings

- Do not apply cuff to areas on patient where skin is delicate or damaged.
- Check cuff site frequently for irritation.
- Allow space for 1 to 2 fingers between patient and cuff.
- Do not apply cuff to limbs used for IV infusion.
- For best results - minimize cuff movement and limb motion during readings.
- Ensure an airtight seal at all connection points prior to use.

Cautions

- Proper cuff sizing and application is essential.
- Cuff should be removed immediately if the patient experiences tingling, numbness, or pain. The physician should be notified.
- Cuff should not be used over an open wound, gangrene, untreated infected wound, recent skin grafts, or dermatitis.
- Cuff should not be inflated too tightly so as not to occlude circulation.
- If inflated too loosely, the cuff will not deliver maximum benefits.

Directions for Use

1. Select cuff size appropriate for the patient's arm circumference.
2. Place cuff under upper arm of patient.
3. Fold Velcro side of cuff over the first side. Secure Velcro attachment.
4. Ensure that the tubes are directed down away from the cuff.
5. Ensure that the cuff is securely attached to the patient.
6. Inflate cuff.

The names of the actual Original Equipment Manufacturer and products mentioned within this document and any information listed on the label is provided as identification prior to High-Level Disinfection reprocessing and may contain trademarks of unrelated third parties who may not represent the device after reprocessing.

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