Symbols description			
CE	CE Marking	ART.	Artery mark
LOT	LOT number	LATEX FREE	Not made with natural rubber latex
	Temperature limitation	MAX RANGE MIN / MIN RANGE MAX	Range indicator of arm circumferences to help selection of the correct cuff size
	Humidity limitation	REF	Reference catalogue number
	Atmospheric pressure limitation		Need for the user to consult the instruction manual
	Identifier of cuffs compatible for the device		Arm circumference
	Cuff positioning indicator for the left arm	MD	Medical device
INDEX	Range pointer and brachial artery alignment position		
Product production date is integrated in the Lot number, which is placed on the product and/or sales package: the first 4 digits mean year of production, the next 2 digits mean month of production and the last 2 digits mean day of production.			

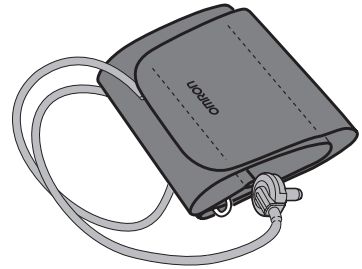
Depending on the product, some of the above symbols may not be applicable on the product.

Manufacturer	**OMRON HEALTHCARE Co., Ltd.** 53, Kunotsubo, Terado-cho, Muko, KYOTO, 617-0002 JAPAN
EU-representative EC REP	**OMRON HEALTHCARE EUROPE B.V.** Scorpius 33, 2132 LR Hoofddorp, THE NETHERLANDS www.omron-healthcare.com
Importer in EU	
Asia Pacific HQ	**OMRON HEALTHCARE SINGAPORE PTE LTD.** 438A Alexandra Road, #05-05/08 Alexandra Technopark, Singapore 119967 www.omronhealthcare-ap.com
Production facility	**OMRON HEALTHCARE MANUFACTURING VIETNAM CO., LTD.** Binh Duong Province, VIETNAM

Made in Vietnam

This OMRON product is produced under the strict quality system of OMRON HEALTHCARE Co., Ltd., JAPAN.

OMRON



HEM-RML31

Wide Range Soft Cuff

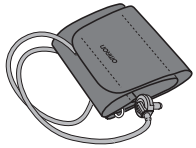
Instructions for use

All for Healthcare

Issue Date:2019-10-03
5342305-6C

OMRON

HEM-RML31



Thank you for purchasing the OMRON HEM-RML31. Arm circumference for this cuff is 22 to 42 cm.

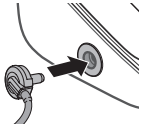
Follow this instruction manual and applicable OMRON blood pressure monitor manual thoroughly before use.
Keep for future reference. For specific information about your own blood pressure, CONSULT YOUR PHYSICIAN.

⚠ ***Caution:*** **Indicates a potentially hazardous situation which, if not avoided, may result in minor or moderate injury to the user or patient or damage to the equipment or other property.**

- Use only the approved Blood Pressure Monitor for this cuff.
- Use the same arm cuff written by the side of the air jack. Do not use the other types of the arm cuff.
- Do not wash the cuff or make it wet.

Instructions for use

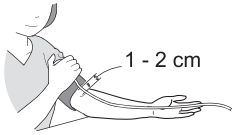
Remove tight-fitting clothing or tight rolled up sleeve from your left upper arm.
Do not place the arm cuff over thick clothes.



1. Insert the air plug into the air jack securely.

2. Apply the arm cuff to your left upper arm snugly.

The bottom edge of the arm cuff should be 1 to 2 cm above the elbow. Air tube is centred on the middle of your inner arm.



3. Close the fabric fastener FIRMLY.

Taking measurements on the right arm
The air tube will be at the side of your elbow.
Be careful not to rest your arm on the air tube.

Note: The blood pressure can differ between the right arm and the left arm, and therefore also the measured blood pressure values can be different. OMRON recommends to always use the same arm for measurement. If the values between both arms differ substantially, please check with your physician which arm to use for your measurement.

Maintenance

- Use a soft and moistened cloth and neutral soap to clean the arm cuff.
- Do not use petrol, thinners of similar solvents to clean the arm cuff.
- Do not store the arm cuff to extreme temperatures, humidity, direct sunlight, dust or corrosive vapours.
- Do not forcibly bend the arm cuff or the air tube excessively.

Specifications:

Product description	Blood pressure monitor cuff
Model	HEM-RML31
Arm circumference	22 to 42 cm
Cuff pressure range	Pressure: 0 to 299 mmHg
Operating conditions	+10 to +40°C / 15 to 90% RH (non-condensing) 700 to 1060 hPa
Storage/ Transport conditions	-20 to +60°C / 10 to 95% RH (non-condensing) 700 to 1060 hPa
Accuracy	Pressure: ±3 mmHg Pulse: ±5 % of display reading
Durable period (Service life)	1 year

CE

Please report to the manufacturer and the competent authority of the Member State in which you are established about any serious incident that has occurred in relation to this device.