

- Fully automatic
- 2 User function
- Traffic light indicator
- PC connection via USB

BP A3L Comfort

- · Cuff check indicator
- Arm movement indicator

- recommended by doctors
- 🖢 Large display



IMT - Inflation Mode Technology

With the inflation mode technology the measurement is performed during the inflation, thus ensuring a quick and comfortable blood pressure measurement with proven Microlife accuracy.



MAM - Microlife Average Mode 3 measurements in 1 click

US Patent 6,447,457 B1

- Automatically takes three measurements with just one click
- Following clinical recommendations to perform at least two automated readings in a row and provide an average of the results
- Patented technology: accurate blood pressure measurement in patients with heart arrhythmia and atrial fibrillation
- Clinically tested by leading experts



Conical cuff fits normal and large arms

- Proper cuff size is crucial for accurate measurements
- Conical shape provides a better fit for improved accuracy
- Suitable for all common arm sizes
- Clinically tested



PAD - Pulse Arrhythmia Detection

US Patent 6,485,429 B2

- Early alerts to possible heart arrhythmias
- Clinically tested pulse arrhythmia detection

Technical specifications

10 - 40 °C / 50 - 104 °F Operating conditions:

15 - 95 % relative maximum humidity

Storage conditions: -20 - +55 °C / -4 - +131 °F

15 - 95 % relative maximum humidity

385 g (including batteries) Weight: Dimensions: 143 x 85 x 58 mm

Measuring procedure: oscillometric, corresponding to Korotkoff

method: Phase I systolic, Phase V diastolic

Measurement range: 20 - 280 mmHg - blood pressure 40 - 200 beats per minute - pulse

Cuff pressure display

range: 0 - 299 mmHg Resolution: 1 mmHg

pressure within ± 3 mmHg Static accuracy: Pulse accuracy: ± 5 % of the readout value

Voltage source: 4 x 1.5 V alkaline batteries; size AA Mains adapter DC 6V, 600 mA (optional)

Reference to standards: EN 1060-1 /-3 /-4; IEC 60601-1;

IEC 60601-1-2 (EMC); IEC 60601-1-11

This device complies with the requirements of the Medical Device Directive 93/42/EEC. Technical alterations reserved.

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