

DECLARATION OF BLOOD PRESSURE MEASURING DEVICE EQUIVALENCE 2013

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SECTION A - Please complete all items.

I **Tomohiro Kukita,** a Director of **Omron Healthcare Europe B.V.,**
Name of a Company Director Company name

hereby state that there are no differences that will affect blood pressure measuring accuracy between the

Maker^a Omron Healthcare Co., Ltd. **Address** 53, Kunotsubo, Terado-cho, Muko, Kyoto 617-0002 Japan
Manufacturer^b Omron Healthcare Co., Ltd. **Address** 53, Kunotsubo, Terado-cho, Muko, Kyoto 617-0002 Japan
Brand^c Omron **Model^d** M3 IT (HEM-7131U-E)

Blood pressure measuring device for which validation is claimed. If alternative model names are used, include all.

blood pressure measuring device and the validated blood pressure measuring device

Maker^a Omron Healthcare Co., Ltd. **Address** 53, Kunotsubo, Terado-cho, Muko, Kyoto 617-0002 Japan
Manufacturer^b Omron Healthcare Co., Ltd. **Address** 53, Kunotsubo, Terado-cho, Muko, Kyoto 617-0002 Japan
Brand^c Omron **Model^d** M6 AC (HEM-7322-E)

Existing validated blood pressure measuring device.

which has previously passed the ESH2010 protocol, the results of which were published as follows:

dablEducational Trust; 2014 Jan 22. 4 p. Available from: [http://www.dableducational.org/Publications/2014/ESH-IP-2010-Validation-of-Omron-M6-AC-\(HEM-7322-E\).pdf](http://www.dableducational.org/Publications/2014/ESH-IP-2010-Validation-of-Omron-M6-AC-(HEM-7322-E).pdf)

Full reference

The only differences between the devices involve the following components:

Tick one box for each item 1-18.

Part I	1	Algorithm for Oscillometric Measurements	Yes <input type="checkbox"/>	No <input checked="" type="checkbox"/>	N/A ^e <input type="checkbox"/>
	2	Algorithm for Auscultatory Measurements	Yes <input type="checkbox"/>	No <input type="checkbox"/>	N/A ^f <input checked="" type="checkbox"/>
	3	Artefact/Error Detection	Yes <input type="checkbox"/>	No <input checked="" type="checkbox"/>	
	4	Microphone(s)	Yes <input type="checkbox"/>	No <input type="checkbox"/>	N/A ^f <input checked="" type="checkbox"/>
	5	Pressure Transducer	Yes <input type="checkbox"/>	No <input checked="" type="checkbox"/>	
	6	Cuffs or Bladders	Yes <input type="checkbox"/>	No <input checked="" type="checkbox"/>	
	7	Inflation Mechanism	Yes <input type="checkbox"/>	No <input checked="" type="checkbox"/>	
	8	Deflation Mechanism	Yes <input type="checkbox"/>	No <input checked="" type="checkbox"/>	
Part II	9	Model Name or Number	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/>	
	10	Casing	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/>	
	11	Display	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/>	
	12	Carrying/Mounting Facilities	Yes <input type="checkbox"/>	No <input checked="" type="checkbox"/>	
	13	Software other than Algorithm	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/>	
	14	Memory Capacity/Number of stored measurements	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/>	
	15	Printing Facilities	Yes <input type="checkbox"/>	No <input type="checkbox"/>	N/A ^g <input checked="" type="checkbox"/>
	16	Communication Facilities	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/>	N/A ^g <input type="checkbox"/>
	17	Power Supply	Yes <input type="checkbox"/>	No <input checked="" type="checkbox"/>	
	18	Other Facilities	Yes <input checked="" type="checkbox"/>	No <input type="checkbox"/>	N/A ^g <input type="checkbox"/>

An explanation of each item ticked "Yes" must be included in Section B or on a separate sheet.

- Notes:
- a Provide the name and address of the actual maker of the device.
 - b Provide the name and address of the legal manufacturer of the device, even if it is the same as that of the maker.
 - c Provide the name of the brand under which it is sold, even if it is the same as that of the manufacturer or maker.
 - d Provide the model name. If alternative or internal model names are used, include all. Each device must be uniquely identifiable.
 - e Only tick N/A (Not Applicable) if neither device measures blood pressure using the oscillometric method.
 - f Only tick N/A (Not Applicable) if neither device measures blood pressure using the auscultatory method.
 - g Only tick N/A (Not Applicable) if neither device provides printing, communication or other facilities, as appropriate.

SECTION B An explanation for each item, 1 to 18, ticked "Yes" in Section A must be provided here or in an attached document. All differences between the devices must be described.

- 9) The model number is changed to M3 IT (HEM-7131U-E) from M6 AC (HEM-7322-E).
- 10) The USB port is added to the M3 IT. The weekly average button is removed from the M3 IT.
- 11) The morning average symbol, the evening average symbol and the morning hypertension symbol are removed from the M3 IT. The OK symbol, the transfer indicator and the DATA/FULL symbol are added to the M3 IT.
- 13) The average at morning and night function and the morning hypertension detection function are removed from the M3 IT. The M3 IT can manage to blood pressure value by software "BiLink".
- 14) The memory capacity has 2 users each 60 sets.
- 16) The USB interface to connect with personal computer are added.
- 17) The M3 IT has AC adapter as optional parts.
- 18) The M3 IT can manage to blood pressure value by software "BiLink".

SECTION C Please check that the following are included with the application

- A manual for the validated device
- A manual for the device for which equivalence is being sought
- An image of the validated device
- An image of the device for which equivalence is being sought
- An image of the screen layout of validated device*
- An image of the screen layout of the device for which equivalence is being sought*

* Screen layouts shown complete, and without obscuring labels or lines, in manuals need not be included separately.

SECTION D Complete all items, bar signatures and seal, online and print. Sign and seal it then send the original to our address below. Please email a signed copy of this form, together with the manuals and images for both devices, to info@dablededucational.org.

Signature of Director Tomohiro Kukita

Company Stamp/Seal

Name Tomohiro Kukita

Date 17 Mar, 2014

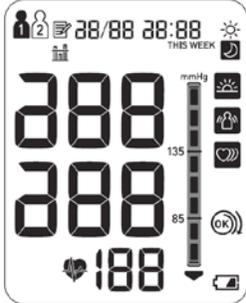
Signature of Witness Anita Kecskes

Name Anita Kecskes

Address 17 Mar, 2014

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 NL-2132 LR Hoofddorp
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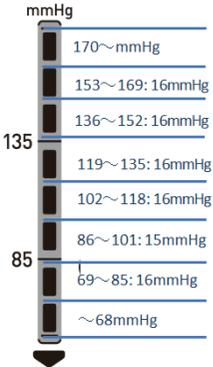
Comparison of the Omron M3 IT (HEM-7131U-E) with the Omron M6 AC (HEM-7322-E)

Devices	Omron M3 IT (HEM-7131U-E)	Omron M6 AC (HEM-7322-E)
Pictures		
Display		
Validation		ESH 2010
Device 1 Criteria	<p>Display/Symbols/Indicators</p> <p><i>Communication</i></p> <p>USB Connection ^{Query 3} 11, 16</p> <p>Data transmission completed  ^{Query 3} 11, 16</p> <p>Data awaiting transmission (Data/Full ) ^{Query 3} 11, 14, 16</p> <p>Algorithms</p> <p><i>Communication</i></p> <p>Data transfer to online database 13, 16</p> <p>Casing</p> <p><i>Ports</i></p> <p>USB port, cable and downloadable PC software 16, 18</p>	
Same Criteria	<p>Measurement</p> <p><i>Accuracy</i></p> <p>BP accuracy ± 3 mmHg 1, 5</p> <p>Pulse accuracy ± 5% 1, 5</p>	<p>Measurement</p> <p><i>Accuracy</i></p> <p>BP accuracy ± 3 mmHg 1, 5</p> <p>Pulse accuracy ± 5% 1, 5</p>

Devices	Omron M3 IT (HEM-7131U-E)	Omron M6 AC (HEM-7322-E)
Same Criteria (continued)	<p>Measurement (continued)</p> <p><i>Method</i></p> <p>Oscillometric measurement method 1, 5</p> <p>BP 0 mmHg to 299 mmHg^{Query 1} 1, 5, 7, 8</p> <p>Pulse 40 bpm to 180 bpm 1, 5, 8</p> <p>Manually initiated measurements 13</p> <p>Measurements are from single inflations 13</p> <p>Prevent storing of result (Guest mode) 13, 14</p> <p><i>Inflation</i></p> <p>Inflation 0 mmHg to 299 mmHg 1, 5, 7</p> <p>Automatic Inflation 7</p> <p>Fuzzy Logic 7</p> <p>Press button if BP > 210 mmHg 7</p> <p><i>Deflation</i></p> <p>Automatic Deflation 8</p> <p><i>Cuffs</i></p> <p>Large (Arm circ. 22 cm to 42 cm) No. HEM-RML31 6</p> <p>Buttons/Switches</p> <p><i>Power</i></p> <p>On/Off with Start/Stop (Start/Stop Label) 10</p> <p><i>Measurement Records</i></p> <p>Memory 10</p> <p>User ID slider 10</p> <p><i>Function</i></p> <p>Date/Time set 10</p> <p>Up and down 10</p> <p>Display/Symbols/Indicators</p> <p><i>Measurement Procedure</i></p> <p>Deflation symbol 11</p> <p>During Measurement: BP Level & Heartbeat 11</p> <p><i>Post Measurement</i></p> <p>SBP, DBP and Pulse 11</p> <p>Measurement error E1 E2 E3 E4 E5 Er 11</p> <p>Hypertension (Indicator strip)^{Query 2} 11, 13</p> <p>Hypertension (Indicator LEDs) 11, 13</p> <p>Memory zone average 11, 13, 14</p>	<p>Measurement (continued)</p> <p><i>Method</i></p> <p>Oscillometric measurement method 1, 5</p> <p>BP 0 mmHg to 299 mmHg^{Query 1} 1, 5, 7, 8</p> <p>Pulse 40 bpm to 180 bpm 1, 5, 8</p> <p>Manually initiated measurements 13</p> <p>Measurements are from single inflations 13</p> <p>Prevent storing of result (Guest mode) 13, 14</p> <p><i>Inflation</i></p> <p>Inflation 0 mmHg to 299 mmHg 1, 5, 7</p> <p>Automatic Inflation 7</p> <p>Fuzzy Logic 7</p> <p>Press button if BP > 210 mmHg 7</p> <p><i>Deflation</i></p> <p>Automatic Deflation 8</p> <p><i>Cuffs</i></p> <p>Large (Arm circ. 22 cm to 42 cm) No. HEM-RML31 6</p> <p>Buttons/Switches</p> <p><i>Power</i></p> <p>On/Off with Start/Stop (Start/Stop Label) 10</p> <p><i>Measurement Records</i></p> <p>Memory 10</p> <p>User ID slider 10</p> <p><i>Function</i></p> <p>Date/Time set 10</p> <p>Up and down 10</p> <p>Display/Symbols/Indicators</p> <p><i>Measurement Procedure</i></p> <p>Deflation symbol 11</p> <p>During Measurement: BP Level & Heartbeat 11</p> <p><i>Post Measurement</i></p> <p>SBP, DBP and Pulse 11</p> <p>Measurement error E1 E2 E3 E4 E5 Er 11</p> <p>Hypertension (Indicator strip)^{Query 2} 11, 13</p> <p>Hypertension (Indicator LEDs) 11, 13</p> <p>Memory zone average 11, 13, 14</p>

Devices	Omron M3 IT (HEM-7131U-E)	Omron M6 AC (HEM-7322-E)
Same Criteria (continued)	<p>Display/Symbols/Indicators (continued)</p> <p><i>Post Measurement (continued)</i></p> <p>Body movement error 3, 11, 13, 18</p> <p>Irregular heartbeat 11, 13, 18</p> <p><i>Measurement Records</i></p> <p>Memory icon 11</p> <p>Memory recall number (Replaces pulse rate momentarily) 11</p> <p>User (1, 2 and Guest) 11</p> <p><i>Date and Time</i></p> <p>Date and Time (During memory recall) 11</p> <p><i>Power</i></p> <p>Low & Exhausted battery 11, 17</p> <p>Algorithms</p> <p><i>Averages and Differences</i></p> <p>Last 3 measurements (within 10 min) memory zone mean 13</p> <p><i>Diagnostic</i></p> <p>BP classification ^{Query 2} 13</p> <p>135 / 85 mmHg thresholds 13</p> <p>Irregular heartbeat detection 13</p> <p>Body movement error detection 3, 13</p> <p><i>Functions</i></p> <p>Correct cuff wrapping detection 13</p> <p>Casing</p> <p><i>Display</i></p> <p>Single screen display 10</p> <p>Segment LCD 10</p> <p><i>Power</i></p> <p>4 “AA” batteries ~ 1000 measurements 17</p> <p>AC adapter (S-9515336-9 or UK-9983666-5) (Optional) 17</p> <p>Automatic switch-off when not used for 2 min 17</p> <p>Rechargeable batteries not permitted 17</p>	<p>Display/Symbols/Indicators (continued)</p> <p><i>Post Measurement (continued)</i></p> <p>Body movement error 3, 11, 13, 18</p> <p>Irregular heartbeat 11, 13, 18</p> <p><i>Measurement Records</i></p> <p>Memory icon 11</p> <p>Memory recall number (Replaces pulse rate momentarily) 11</p> <p>User (1, 2 and Guest) 11</p> <p><i>Date and Time</i></p> <p>Date and Time (During memory recall) 11</p> <p><i>Power</i></p> <p>Low & Exhausted battery 11, 17</p> <p>Algorithms</p> <p><i>Averages and Differences</i></p> <p>Last 3 measurements (within 10 min) memory zone mean 13</p> <p><i>Diagnostic</i></p> <p>BP classification ^{Query 2} 13</p> <p>135 / 85 mmHg thresholds 13</p> <p>Irregular heartbeat detection 13</p> <p>Body movement error detection 3, 13</p> <p><i>Functions</i></p> <p>Correct cuff wrapping detection 13</p> <p>Casing</p> <p><i>Display</i></p> <p>Single screen display 10</p> <p>Segment LCD 10</p> <p><i>Power</i></p> <p>4 “AA” batteries ~ 1000 measurements 17</p> <p>AC adapter (S-9515336-9 or UK-9983666-5) (Optional) 17</p> <p>Automatic switch-off when not used for 2 min 17</p> <p>Rechargeable batteries not permitted 17</p>
Comparable Criteria	<p>Measurement</p> <p><i>Measurement Records</i></p> <p>Memory: 60 measurements × 2 users (Guest not recorded) 14</p> <p>Display/Symbols/Indicators</p> <p><i>Post Measurement</i></p> <p>Correct cuff wrapping indicator (icon + LED) 11, 13, 18</p>	<p>Measurement</p> <p><i>Measurement Records</i></p> <p>Memory: 100 measurements × 2 users (Guest not recorded) 14</p> <p>Display/Symbols/Indicators</p> <p><i>Post Measurement</i></p> <p>Correct cuff wrapping indicator (icon + 2 LEDs) 11, 13, 18</p>

Devices	Omron M3 IT (HEM-7131U-E)	Omron M6 AC (HEM-7322-E)
Comparable Criteria (continued)	<i>Display/Symbols/Indicators (continued)</i> <i>Date and Time</i> Date and Time (alternating) 11	<i>Display/Symbols/Indicators (continued)</i> <i>Date and Time</i> Date and Time 11
Device 2 Criteria		<i>Buttons/Switches</i> <i>Analysis</i> Average 10 <i>Display/Symbols/Indicators</i> <i>Post Measurement</i> Morning hypertension 11, 13 7-day morning memory zone average 11, 13, 14 7-day evening memory zone average 11, 13, 14 <i>Algorithms</i> <i>Averages and Differences</i> 7-day morning and evening memory zone means (8 weeks) 13 <i>Diagnostic</i> Morning hypertension 13

<p>Queries</p>	<p>1</p>	<p>Query Each of the manuals states that the blood pressure measurement range is 0 mmHg to 299 mmHg. They also state that the monitor will not inflate above 299 mmHg. This means that the measurement ranges must be below this.</p> <p>According to ISO 80601-2-30 (2009), the device must be capable (in non-neonatal mode) of indicating at least 60 mmHg to 230 mmHg for SBP and 40 mmHg to 130 mmHg for DBP (201.12.1.103), so specifying these separately is necessary. It also requires that the pressure ranges provided are the rated pressures (201.7.9.2.9 h and 201.12.1.101) and that those measurements that are outside of these ranges trigger a technical alarm (201.12.1.106).</p> <p>a) What are the SBP and DBP rated ranges for each of the devices?</p> <p>b) Are there technical alarm ranges for each of the devices?</p> <p>Response a) There is no SBP and DBP rated ranges because we have not defined the rated range of cuff pressure which is actually limited by measurement range of the pressure (not blood pressure) 0 to 299 mmHg. The capability to measure the required SBP and DBP range (201.12.1.103) are confirmed by technical validation test.</p> <p>b) There is no technical alarm because of the reason above.</p> <p>Query The rated ranges for SBP and DBP are simply the ranges, within the inflation range, for which SBP and DBP values are displayed. Where a pulse is detected close to the maximum inflation pressure or the wave envelope suggests SBP as being close to the maximum inflation pressure, it may be rejected an unreliable estimate of SBP. DBP estimates close to zero can be similarly rejected. These are the technical alarm conditions.</p> <p>The reply suggests that there are no upper or lower limits to either SBP or DBP i.e. $299 \text{ mmHg} \geq \text{SBP} > \text{DBP} \geq 0 \text{ mmHg}$. Is this correct?</p> <p>Response Yes, this is correct.</p> <p>Comment The reply answers the query fully.</p>
	<p>2</p>	<p>Query Both devices have vertical indicator strips. Each strip contains 8 sections with a 135 mmHg mark between the 3rd and 4th sections and an 85 mmHg mark between the 6th and 7th sections. From the diagrams in the respective manuals, their function appears to be to display the pulse pressure, with a series of sections lit from one indicating a range of SBP values to one indicating a range of DBP values. However, no explanation is provided in either manual. What range of pressure does each section represent and are they the same for both devices?</p> <p>Response Please find the range of the pressure for each section as shown. These are same for both devices.</p> <p>Comment The reply answers the query fully.</p> 

	3	<p>Query The M3 IT (HEM-7131U-E) has Transfer, OK and Data/Full symbols that are not described in the manual. Please outline the functions of these symbols.</p> <p>Response The Transfer indicator ⇐ is displayed when the USB cable is connected.</p> <p>The OK symbol is displayed when the data transfer has finished.</p> <p>The DATA/Full symbol ■ is not displayed, for the selected user, if there are fewer than 48 measurements not transferred. It will blink if there are between 48 and 59 measurements not transferred. It will be shown as a steady symbol if the memory is full (60 measurements) and none of them are transferred.</p> <p>Comment The reply answers the query fully.</p>
Note	1	<p>The main differences between these devices are that the M3 IT (HEM-7131U-E) has additional features to allow measurements to be transferred to an online database whereas the M6 AC (HEM-7322-E) provides morning and evening averages. The M3 IT (HEM-7131U-E) stores 60 measurements per user, as distinct from 100 per user for the M6 AC (HEM-7322-E) and also uses the same segments to display an alternating date and time whereas the M6 AC (HEM-7322-E) can display both together.</p> <p>Both devices have a green LED symbol to indicate if the cuff was wrapped correctly The M6 AC (HEM-7322-E) has an extra separate orange LED symbols to indicate if the cuff was wrapped too loosely; no light is shown on the M3 IT (HEM-7131U-E) in this instance. The feature is duplicated on the screen of both devices where the cuff wrap symbol is shown with either the “OK” or “tighten it” arrow sections to indicate correct or loose cuff wrapping.</p>
Recommendation	Equivalence is Recommended	
Date	20 March 2014	