Medical Device Assessment



Medaval Accreditation Assessment

Volume 2016 Report 1629 05 August 2016

Accreditation assessment of the blood pressure measurement technology used in the Omron HEM-7252G-HP upper arm monitor, as validated according to the European Society of Hypertension International Protocol revision 2010

Approved by the Medaval Advisory Board

Eoin O'Brien (Chair)
George S. Stergiou (Deputy Chair)
Roland Asmar G
Alejandro de la Sierra T
Peter W. de Leeuw
Eamon Dolan G
Geoffrey A. Head

Yutaka Imai Andrew Shennan
Martin Myers Jan Staessen
Gbenga Ogedegbe Martin J. Turner
Takayoshi Ohkubo Paolo Verdecchia
Paolo Palatini Bernard Waeber
Gianfranco Parati J-Guang Wang

Reference

Medaval Ltd. Accreditation assessment of the blood pressure measurement technology used in the Omron HEM-7252G-HP upper arm monitor, as validated according to the European Society of Hypertension International Protocol revision 2010. *Medical Device Assessment*. 2016 Aug 5;**2016**(1629). 5 p. Epub: 2019 Jan 31. Available from: https://www.medaval.ie/MDA/2016/MDA1629.pdf.

Medical Device Assessment is published by

Medaval Ltd., Unit 107, SBC, Serpentine Ave., Ballsbridge, Dublin D04 H522, IRELAND.

© 2016-2019 Medaval Ltd. All rights reserved.

Permissions: Requests for permissions to reproduce figures, tables, or portions of reports or articles originally published in *Medical Device Assessment* can be obtained by email request to info@medaval.ie.

Accreditation assessment of the blood pressure measurement technology used in the Omron HEM-7252G-HP upper arm monitor, as validated according to the European Society of Hypertension International Protocol revision 2010

Medaval Accreditation-Assessment Report – 5th August 2016

Test Device Details

Assessment **Full Name** Omron HEM-7252G-HP Requirement satisfactory Model HEM-7252G-HP Requirement satisfactory **Measurement Site** Upper Arm Requirement satisfactory Client Use Suitable for self-measurement. Requirement satisfactory **Operation Method** Oscillometry, automatic during Requirement satisfactory deflation **Measurement Occurrence** Single Measurements Only Requirement satisfactory **Device Photograph** Photograph not in paper. Standard image shown.



Manufacturer(s) Sole: Omron Healthcare, Kyoto

Head Office, Shiokoji Horikawa, Shimogyo ku, Kyoto 600 8530,

JAPAN.

Cuffs Omron HEM-CS24-L: Small 17

cm to 22 cm

Omron HEM-CR24-L: Medium

22 cm to 32 cm

Omron HEM-RML31-L: Large

32 cm to-42 cm

Requirement satisfactory

Cuffs Listed: Requirement satisfactory

Arm Circumferences: Requirement satisfactory

Assessment

Study Details

Original Publication Takahashi H, Yoshika M, Yokoi T. Validation of two automatic devices: Omron HEM-7252G-HP and

Omron HEM-7251G for self-measurement of blood pressure according to the European Society of Hypertension International Protocol revision 2010. *Blood Press Monit*. 2015 Oct;**20**(5):286-90.

Epub: 2015 Apr 29. doi: 0.1097/MBP.000000000000127. PMID: 25932887.

Protocol The European Society of Hypertension International Protocol revision 2010 for the validation of

blood pressure measuring devices in adults¹

AdherenceNot stated but clear from textRequirement satisfactoryAdjustmentsNoneRequirement satisfactoryStudy Meas. MethodOscillometricRequirement satisfactoryStudy Measurement SiteUpper ArmRequirement satisfactory

Observers

Supervisor + 2 ObserversYesRequirement satisfactoryObserver TrainingBHS online trainingRequirement satisfactoryObserver FamiliarisationGeneral testingRequirement satisfactory

Observer Blinding From each other Modification: Missing "from device" accepted by review

Sample

PopulationA general populationRequirement satisfactoryCircumstancesNoneRequirement satisfactoryHBP Subjects SelectionOutpatientsRequirement satisfactoryNBP Subjects SelectionHospital staff & volunteersRequirement satisfactory

Test Device Details and Study Details Assessment
Checks
Permitted Modifications
2
Violations
0

Procedure

Table 1: Screening and Recruitment Details

	3	creening and Recruit	ment			Assessment	:	
Total Screened 44						Value within requirements		
Total Excluded 11					11	Value within requirements		
Ranges Complete 1						Value within requirements		
Range Adjustment 0						Value within requirements		
Arrhythmias 4						Value within requirements		
Device Failure 0 Poor Quality Sounds 1						Value within requirements Value within requirements		
	Cuff Size U	Jnavailable	0			Value within requirements		
	Observer	Disagreement	0			Value within requirements		
	Distribution	on	0			Value within requirements		
	Other Rea	sons*	5			Value within requirements		
Total F	Recruited				33	Value within requirements		
*Expla	nation Sum							
	The device	e displayed a body m	oveme	nt erro	r in five	Details satisfactory		
	individuals	who were excluded	for this	reaso	٦.			
		December 15						
SBP	Total	Recruitment Range	es		33	Value within requirements		
SDP	Low			12	33	Value within requirements		
	LOW	< 90 mmHq	0	12				
		90 – 129 <i>mmHq</i>	0 12			Value within requirements Value within requirements		
	Medium	130 – 160 mmHg	12	11		Value within requirements		
		150 – 160 minny		10				
	High	161 – 180 mmHg	6	10		Value within requirements Value within requirements		
		> 180 mmHq	4			Value within requirements		
		> 100 mming	-			value within requirements		
DBP	Total				33	Value within requirements		
	Low			11		Value within requirements		
		< 40 mmHg	0			Value within requirements		
		40 –79 mmHg	11			Value within requirements		
	Medium	80 – 100 <i>mmHg</i>		11		Value within requirements		
	High			11		Value within requirements		
		101 – 130 mmHg	11			Value within requirements		
		> 130 mmHg	0			Value within requirements		
				_				
I otal E	xtremes			4		Value within requirements		
		On Treatment Rang	zes					
SBP	Low	< 130 mmHg	,	0		Value within requirements		
JUF	Medium	130 – 160 mmHg		2		Value within requirements		
	High	> 160 mmHq		2		Value within requirements		
	6.,	, 100 mming		_		value within requirements		
DBP	Low	< 80 mmHg		0		Value within requirements		
	Medium	80 – 100 <i>mmHg</i>		2		Value within requirements		
	High	> 100 mmHg		2		Value within requirements		
Table	1 Assessme	nt				Checks	36	
						Permitted Modifications	0	
						Violations	0	

Study Results

Table 2: Subject Details

			Assessment		
Sex	Male:Female	19:14	Value within requirements	Value within requirements	
Ago (vogra)	Range (Low:High)	28:78	Value within requirements	Value within requirements	
Age (years)	Mean (SD)	50 (12)	Value within requirements	Value within requirements	
Arm Circumference	Range (Low:High)	20.1:37.2	Value within requirements	Value within requirements	
(cm)	Mean (SD)	28.1 (4.6)	Value within requirements	Value within requirements	
Cuff for Test Device	Small <i>(17 –22)</i>	4			
(cm)	Medium (22 – 32)	23			
	Large <i>(32 – 42)</i>	6			
	Total	33	Value within requirements		
Recruitment SBP	Range (Low:High)	90:207	Value within requirements	Value within requirements	
(mmHg)	Mean (SD)	144 (32.7)	Value within requirements	Value within requirements	
Recruitment DBP	Range (Low:High)	48:129	Value within requirements	Value within requirements	
(mmHg)	Mean (SD)	90 (20.2)	Value within requirements	Value within requirements	
Table 2 Assessment			Checks	19	
			Permitted Modifications	0	
			Violations	0	

Table 3: Observer Measurements in each Recruitment Range

			Assessment		
SBP	Overall Range mmHg (Low:High)	86:218	Value within requirements	Value within requirements	
	Low (< 130 mmHg)	41	Value within	requirements	
	Medium (130 – 160 mmHg)	28	Value within	requirements	
	High (> 160 mmHg)	30	Value within	requirements	
	Maximum Difference	13	Value within	requirements	
DBP	Overall Range mmHg (Low:High)	48:134	Value within requirements	Value within requirements	
	Low (< 80 <i>mmHg</i>)	26	Value within	requirements	
	Medium (80 – 100 <i>mmHg</i>)	42	Value within	requirements	
	High (> 100 <i>mmHq</i>) 31		Value within requirements		
	Maximum Difference	16	Value within	requirements	
Table 3 Assessment			Checks	12	
			Permitted Modifications	0	
			Violations	0	

Table 4: Observer Differences

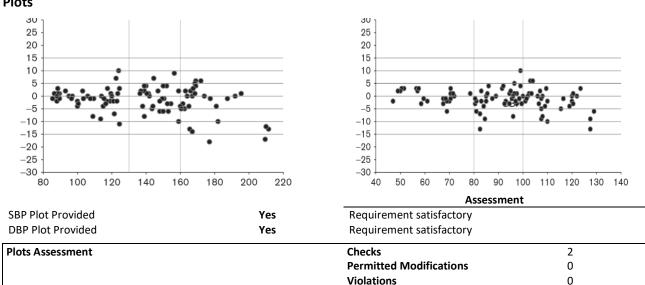
			Assessment		
Observer 2 – Obser	ver 1				
SBP (mmHg)	Range (Low:High)	-4:+4	Value within requirements	Value within requirements	
	Mean (SD)	0 (1.3)	Value within requirements	Value within requirements	
DBP (mmHg)	Range (Low:High)	-4:+4	Value within requirements	Value within requirements	
, 5,	Mean (SD)	0 (1.3)	Value within requirements	Value within requirements	
Repeated Measurements		0	Value within	requirements	
Table 4 Assessment			Checks	9	
			Permitted Modifications	0	
			Violations	0	

Table 5: Validation Results

Part 1	Pass Req.		Achieved		Assessment	
	Two of	All of	SBP	DBP		
<u><</u> 5 mmHg	73	65	76	83	Value within passing criteria	Value within passing criteria
<u><</u> 10 mmHg	87	81	92	97	Value within passing criteria	Value within passing criteria
<u><</u> 15 mmHg	96	93	97	99	Value within passing criteria	Value within passing criteria
Grade 1			Pass	Pass	Value within passing criteria	Value within passing criteria
Mean <i>mmHg</i>			-1.5	-1.2	Value within requirements	Value within requirements
SD mmHg			5.1	3.9	Value within requirements	Value within requirements
Part 2		Pass	Achieved			
		Req.	SBP	DBP		
2/3 <u><</u> 5 mmHg	'	<u>> 24</u>	28	29	Value within passing criteria	Value within passing criteria
0/3 <u><</u> 5 mmHg	1	<u><</u> 3	1	2	Value within passing criteria	Value within passing criteria
Grade 2			Pass	Pass	Value within passing criteria	Value within passing criteria
Grade 3			Pass	Pass	Value within passing criteria	Value within passing criteria
Part 3						
Result			Pa	ass	Value within passing criteria	

Table 5 Assessment	Checks	21
	Permitted Modifications	0
	Violations	0

Plots



Recommendations

Overall Summary

Number of checks	121
Number of permitted modifications	2
Number of violations	0

Assessment Summary

The validation has been checked and is verified as having been conducted in accordance with the protocol requirements. Therefore, the results are considered to be valid, the null hypothesis, that the device is inaccurate in measuring blood pressure, is rejected and the conclusion, that the device is accurate for self-measurement in adults, is correct.

Certification Decision

The Omron HEM-7252G-HP, with the HEM-CR24-L: medium 22 cm to 32 cm or HEM-RML31-L: large 32 cm to-42 cm is certified by Medaval Ltd., for blood pressure measurement in adults, as it fulfilled the conditions required for a pass in a validation study carried out in accordance with the requirements of the International Protocol of the European Society of Hypertension 2010 Revision.

Date of Advisory Board Approval: 29th July 2016.

Reference

O'Brien E, Atkins N, Stergiou G, Karpettas N, Parati G, Asmar R, Imai Y, Wang J, Mengden T, Shennan A; Working Group on Blood Pressure Monitoring of the European Society of Hypertension. European Society of Hypertension International Protocol revision 2010 for the validation of blood pressure measuring devices in adults. Blood Press Monit. 2010;15:23-38. doi: 10.1097/MBP.0b013e3283360e98. PMID: 20110786. Erratum in Blood Press Monit. 2010;15(3):171-2.