Medical Device Assessment



Medaval Accreditation Assessment

Volume 2016 Report 1620 05 August 2016

Accreditation assessment of the blood pressure measurement technology used in the Omron HEM-7500F upper arm monitor, as validated according to the European Society of Hypertension International Protocol revision 20100

Approved by the Medaval Advisory Board

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Reference

Medaval Ltd. Accreditation assessment of the blood pressure measurement technology used in the Omron HEM-7500F upper arm monitor, as validated according to the European Society of Hypertension International Protocol revision 2010. *Medical Device Assessment*. 2016 Aug 5;**2016**(1620). 5 p. Epub: 2019 Jan 31. Available from: https://www.medaval.ie/MDA/2016/MDA1620.pdf.

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Accreditation assessment of the blood pressure measurement technology used in the Omron HEM-7500F 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-7500F Requirement satisfactory Model HEM-7500F Requirement satisfactory **Measurement Site** Requirement satisfactory Upper Arm Suitable for self-measurement. Requirement satisfactory Client Use **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.

Omron HEM-FSM50: Wide Cuffs Cuffs Listed: Requirement satisfactory

Range 17 cm to-36 cm Arm Circumferences: Requirement satisfactory

Study Details

Original Publication Takahashi H, Yoshika M, Yokoi T. Validation of three automatic devices for the self-measurement

> of blood pressure according to the European Society of Hypertension International Protocol revision 2010: the Omron HEM-7130, HEM-7320F, and HEM-7500F. Blood Press Monit. 2015

Requirement satisfactory

Apr;20(2):92-7. doi: 10.1097/MBP.000000000000096. PMID: 25462531.

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

blood pressure measuring devices in adults1

Assessment Adherence Not stated but, apart from Requirement accepted as satisfactory paper, inferred from text.

Adjustments None Requirement satisfactory Study Meas. Method Oscillometric Requirement satisfactory **Study Measurement Site** Upper Arm Requirement satisfactory

Observers

Supervisor + 2 Observers Yes Requirement satisfactory **Observer Training** BHS online training Requirement satisfactory

Observer Familiarisation Requirement accepted as satisfactory Not described but assumed

completed

Observers Blinded From each other and assumed Requirement satisfactory

from device

Sample

Population A general population Requirement satisfactory Circumstances None Requirement satisfactory Outpatients **HBP Subjects Selection** Requirement satisfactory

NBP Subjects Selection Hospital staff & volunteers Requirement satisfactory

Test Device Details and Study Details Assessment Checks 22 **Permitted Modifications** 1 **Violations** 0

Procedure

Table 1: Screening and Recruitment Details

	S	creening and Recruit	ment			Assessment	
Total S	Screened				48	Value within requirements	
Total Excluded 15						Value within requirements	
Ranges Complete			0			Value within requirements	
	Range Adj	ustment	2			Value within requirements	
	Arrhythmi	ias	6			Value within requirements	
	Device Fai	lure	0			Value within requirements	
	Poor Qual	ity Sounds	2			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	mary					
	The only r	eason for exclusion,	stated i	in the te	ext and	Broad explanation accepted	
	not includ	led above, is body r	noveme	ent erro	or. It is		
	assumed 5	subjects were exclu	ded for	this rea	ason.		
	Tatal	Recruitment Range	es			Value within as with a section	
SBP	Total				33	Value within requirements	
	Low	.00	_	11		Value within requirements	
		< 90 mmHg	1			Value within requirements	
		90 – 129 <i>mmHg</i>	10			Value within requirements	
	Medium	130 – 160 <i>mmHg</i>		11		Value within requirements	
	High		_	11		Value within requirements	
		161 – 180 <i>mmHg</i>	9			Value within requirements	
		> 180 mmHg	2			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 mmHg		10		Value within requirements	
	High	J		12		Value within requirements	
	•	101 – 130 mmHg	12			Value within requirements	
		> 130 mmHg	0			Value within requirements	
Total E	xtremes			3		Value within requirements	
		On Treatment Rang	265				
SBP	Low	< 130 mmHg	,	0		Value within requirements	
55.	Medium	130 – 160 <i>mmHg</i>		6		Value within requirements	
	High	> 160 mmHg		1		Value within requirements	
	J	3				4	
DBP	Low	< 80 mmHg		0		Value within requirements	
	Medium	80 – 100 <i>mmHg</i>		4		Value within requirements	
	High	> 100 mmHg		3		Value within requirements	
Table	1 Assessme	nt				Checks	36
		-				Permitted Modifications	0
						Violations	0
							-

Study Results

Table 2: Subject Details

			Assessment		
Sex	Male:Female	21:12	Value within requirements	Value within requirements	
	Range (Low:High)	33:73	Value within requirements	Value within requirements	
Age (years)	Mean (SD)	50 (11)	Value within requirements	Value within requirements	
	iviean (SD)	30 (11)	value within requirements	value within requirements	
Arm Circumference	Range (Low:High)	18.9:35.8	Value within requirements	Value within requirements	
(cm)	Mean (SD)	28.5 (4.4)	Value within requirements	Value within requirements	
Cuff for Test Device	Standard <i>(17 – 36)</i>	33			
(cm)	Total	33	Value within requirements		
Recruitment SBP	Range (Low:High)	85:188	Value within requirements	Value within requirements	
(mmHg)	Mean (SD)	143 (30.4)	Value within requirements	Value within requirements	
Recruitment DBP	Range (Low:High)	49:126	Value within requirements	Value within requirements	
(mmHg)	Mean (SD)	89 (21.7)	Value within requirements	Value within requirements	
Table 2 Assessment			Checks	19	
			Permitted Modifications	0	
			Violations	0	

Table 3: Observer Measurements in each Recruitment Range

			Asses	sment
SBP	Overall Range mmHg (Low:High)	88:190	Value within requirements	Value within requirements
	Low (< 130 mmHg)	41	Value within	requirements
	Medium (130 – 160 mmHg)	33	Value within	requirements
	High (> 160 mmHg)	25	Value within	requirements
	Maximum Difference	16	Value within	requirements
DBP	Overall Range mmHg (Low:High)	49:129	Value within requirements	Value within requirements
	Low (< 80 <i>mmHg</i>)	39	Value within	requirements
	Medium (80 – 100 mmHg)	24	Value within	requirements
	High (> 100 <i>mmHg</i>)	36	Value within	requirements
	Maximum Difference	15	Value within	requirements
Table 3	3 Assessment		Checks	12
			Permitted Modifications	0
			Violations	0

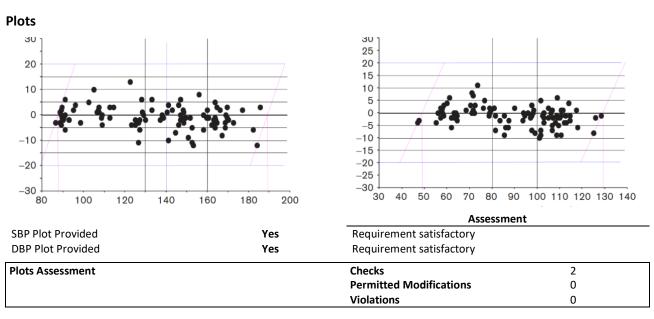
Table 4: Observer Differences

			Assessment		
Observer 2 – Observ	ver 1				
SBP (mmHg)	Range (Low:High)	-4:+4	Value within requirements	Value within requirements	
	Mean (SD)	+0.4 (1.4)	Value within requirements	Value within requirements	
DBP (mmHg)	Range (Low:High)	-4:+2	Value within requirements	Value within requirements	
	Mean (SD)	-0.3 (1.5)	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	79	78	Value within passing criteria	Value within passing criteria
<u>−</u> 10 <i>mmHg</i>	87	81	93	98	Value within passing criteria	Value within passing criteria
_ ≤ 15 mmHg	96	93	99	99	Value within passing criteria	Value within passing criteria
Grade 1			Pass	Pass	Value within passing criteria	Value within passing criteria
Mean mmHg			-1.0	-1.1	Value within requirements	Value within requirements
SD mmHg			4.5	4.0	Value within requirements	Value within requirements
Part 2		Pass	Achi	eved		
		Req.	SBP	DBP		
2/3 < 5 mmHg	•	<u>></u> 24	28	28	Value within passing criteria	Value within passing criteria
0/3 <u><</u> 5 mmHg		<u><</u> 3	0	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			Pass		Value within passing criteria	
Table F Assessmen	•				Chasks	21

Table 5 Assessment	Checks	21
	Permitted Modifications	0
	Violations	0



Recommendations

Overall Summary

Number of checks	121
Number of permitted modifications	1
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-7500F, with the HEM-FSM50: wide-range 17 cm to-36 cm cuff, 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.