

Medaval Accreditation Assessment

Volume 2016

Report 1608

05 August 2016

Accreditation assessment of the blood pressure measurement technology used in the Andon KD-5965 (KP-5965) 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)	Yutaka Imai	Andrew Shennan
George S. Stergiou (Deputy Chair)	Martin Myers	Jan Staessen
Roland Asmar	Gbenga Ogedegbe	Martin J. Turner
Alejandro de la Sierra	Takayoshi Ohkubo	Paolo Verdecchia
Peter W. de Leeuw	Paolo Palatini	Bernard Waeber
Eamon Dolan	Gianfranco Parati	J-Guang Wang
Geoffrey A. Head		

Reference

Medaval Ltd. Accreditation assessment of the blood pressure measurement technology used in the Andon KD-5965 (KP-5965) upper arm monitor, as validated according to the European Society of Hypertension International Protocol revision 2010. *Medical Device Assessment*. 2016 Aug 5;2016(1608). 5 p. Epub: 2019 Jan 31. Available from: <https://www.medaval.ie/MDA/2016/MDA1608.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 Andon KD-5965 (KP-5965) 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	Andon KD-5965	Requirement satisfactory
Model	KP-5965	Requirement satisfactory
Measurement Site	Upper Arm	Requirement satisfactory
Client Use	Suitable for self-measurement.	Requirement satisfactory
Operation Method	Oscillometry, automatic during deflation	Requirement satisfactory
Measurement Occurrence	Single Measurements Only	Requirement satisfactory
Device Photograph		Modification: No photograph in paper. Standard image shown in report.
Manufacturer(s)	Andon Health Co. Ltd., 3 Jinping Road, Ya'an Street, Nankai District, Tianjin 300190, CHINA	Requirement satisfactory
Cuffs	Small (15 – 24) Standard (20 – 34) Large (30 – 44) X Large (40 – 48)	Cuffs Listed: Requirement satisfactory Arm Circumferences: Requirement satisfactory

Study Details

Original Publication	Huang J, Li Z, Li G, Liu Z. Validation of the Andon KD-5965 upper-arm blood pressure monitor for home blood pressure monitoring according to the European Society of Hypertension International Protocol revision 2010. <i>Blood Press Monit.</i> 2015 Oct; 20 (5):283-5. Epub: 2015 May 5. doi: 10.1097/MBP.0000000000000129. PMID: 25968093.
Protocol	The European Society of Hypertension International Protocol revision 2010 for the validation of blood pressure measuring devices in adults ¹

		Assessment
Adherence	Followed Precisely	Requirement satisfactory
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 tutorial	Requirement satisfactory
Observer Familiarisation	12 measurements	Requirement satisfactory
Observers Blinded	From device and each other	Requirement satisfactory
Sample		
Population	A general population	Requirement satisfactory
Circumstances	None	Requirement satisfactory
HBP Subjects Selection	Inpatients and outpatients	Requirement satisfactory
NBP Subjects Selection	Hospital staff	Requirement satisfactory

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

Procedure

Table 1: Screening and Recruitment Details

Screening and Recruitment				Assessment
Total Screened			42	Value within requirements
Total Excluded			9	Value within requirements
	Ranges Complete		7	Value within requirements
	Range Adjustment		0	Value within requirements
	Arrhythmias		2	Value within requirements
	Device Failure		0	Value within requirements
	Poor Quality Sounds		0	Value within requirements
	Cuff Size Unavailable		0	Value within requirements
	Observer Disagreement		0	Value within requirements
	Distribution		0	Value within requirements
	Other Reasons*		0	Value within requirements
Total Recruited			33	Value within requirements
*Explanation Summary				No details required
Recruitment Ranges				
SBP	Total		33	Value within requirements
	Low	< 90 mmHg	0	Value within requirements
		90 – 129 mmHg	10	Value within requirements
		130 – 160 mmHg	12	Value within requirements
	Medium	161 – 180 mmHg	10	Value within requirements
		> 180 mmHg	1	Value within requirements
	High		11	Value within requirements
DBP	Total		33	Value within requirements
	Low	< 40 mmHg	0	Value within requirements
		40 – 79 mmHg	11	Value within requirements
		80 – 100 mmHg	12	Value within requirements
	Medium	101 – 130 mmHg	10	Value within requirements
		> 130 mmHg	0	Value within requirements
High		10	Value within requirements	
Total Extremes			1	Value within requirements
On Treatment Ranges				
SBP	Low	< 130 mmHg	2	Value within requirements
	Medium	130 – 160 mmHg	10	Value within requirements
	High	> 160 mmHg	11	Value within requirements
DBP	Low	< 80 mmHg	4	Value within requirements
	Medium	80 – 100 mmHg	9	Value within requirements
	High	> 100 mmHg	10	Value within requirements
Table 1 Assessment				Checks 36
				Permitted Modifications 0
				Violations 0

Study Results

Table 2: Subject Details

			Assessment	
Sex	Male:Female	13:20	Value within requirements	Value within requirements
Age (years)	Range (Low:High)	26:82	Value within requirements	Value within requirements
	Mean (SD)	65.1 (12.5)	Value within requirements	Value within requirements
Arm Circumference (cm)	Range (Low:High)	25:35	Value within requirements	Value within requirements
	Mean (SD)	29.7 (2.5)	Value within requirements	Value within requirements
Cuff for Test Device (cm)	Small (15 – 24)	0		
	Standard (20 – 34)	26		
	Large (30 – 44)	7		
	X Large (40 – 48)	0		
	Total	33	Value within requirements	
Recruitment SBP (mmHg)	Range (Low:High)	96:182	Value within requirements	Value within requirements
	Mean (SD)	144.7 (22.4)	Value within requirements	Value within requirements
Recruitment DBP (mmHg)	Range (Low:High)	50:125	Value within requirements	Value within requirements
	Mean (SD)	88.1 (16.7)	Value within requirements	Value within requirements
Table 2 Assessment			Checks	19
			Permitted Modifications	0
			Violations	0

Table 3: Observer Measurements in each Recruitment Range

			Assessment	
SBP	Overall Range mmHg (Low:High)	83:180	Value within requirements	Value within requirements
	Low (< 130 mmHg)	26 to 41	Modification: Generality accepted by paper review	
	Medium (130 – 160 mmHg)	26 to 41	Modification: Generality accepted by paper review	
	High (> 160 mmHg)	26 to 41	Modification: Generality accepted by paper review	
	Maximum Difference	≤ 15	Modification: Generality accepted by paper review	
DBP	Overall Range mmHg (Low:High)	48:121	Value within requirements	Value within requirements
	Low (< 80 mmHg)	26 to 41	Modification: Generality accepted by paper review	
	Medium (80 – 100 mmHg)	26 to 41	Modification: Generality accepted by paper review	
	High (> 100 mmHg)	26 to 41	Modification: Generality accepted by paper review	
	Maximum Difference	≤ 15	Modification: Generality accepted by paper review	
Table 3 Assessment			Checks	12
			Permitted Modifications	8
			Violations	0

Table 4: Observer Differences

			Assessment	
Observer 2 – Observer 1				
SBP (mmHg)	Range (Low:High)	-4:+4	Value within requirements	Value within requirements
	Mean (SD)	+0.1 (1.4)	Value within requirements	Value within requirements
DBP (mmHg)	Range (Low:High)	-4:+4	Value within requirements	Value within requirements
	Mean (SD)	+0.2 (1.4)	Value within requirements	Value within requirements
Repeated Measurements		2	Value within requirements	
Table 4 Assessment			Checks	9
			Permitted Modifications	0
			Violations	0

Table 5: Validation Results

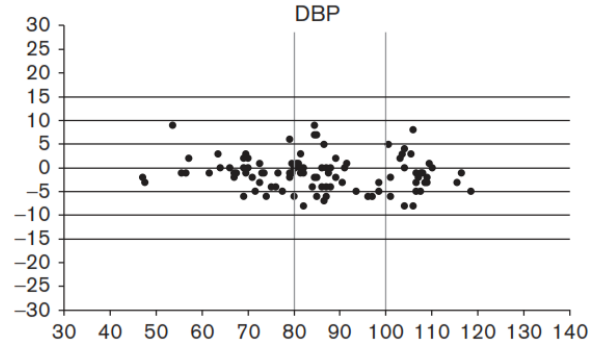
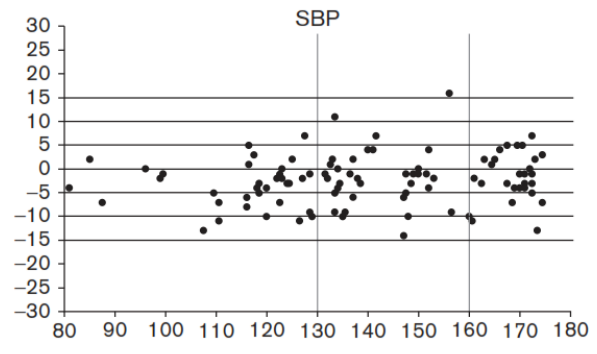
Part 1	Pass Req.		Achieved		Assessment	
	Two of	All of	SBP	DBP		
≤ 5 mmHg	73	65	70	81	Value within lower passing criteria	Value within passing criteria
≤ 10 mmHg	87	81	91	99	Value within passing criteria	Value within passing criteria
≤ 15 mmHg	96	93	98	99	Value within passing criteria	Value within passing criteria
Grade 1			Pass	Pass	Value within lower passing criteria	Value within passing criteria
Mean mmHg			-2.37	-1.10	Value within requirements	Value within requirements
SD mmHg			5.28	3.65	Value within requirements	Value within requirements

Part 2	Pass Req.	Achieved		Assessment	Assessment
		SBP	DBP		
2/3 ≤ 5 mmHg	≥ 24	25	29	Value within passing criteria	Value within passing criteria
0/3 ≤ 5 mmHg	≤ 3	2	1	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 lower passing criteria	Value within passing criteria

Part 3	Result	Pass	Value within lower passing criteria
--------	--------	------	-------------------------------------

Table 5 Assessment	Checks	21
	Permitted Modifications	0
	Violations	0

Plots



SBP Plot Provided	Yes	Requirement satisfactory
DBP Plot Provided	Yes	Requirement satisfactory

Plots Assessment	Checks	2
	Permitted Modifications	0
	Violations	0

Recommendations

Overall Summary

Number of checks	121
Number of permitted modifications	9
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 Andon KD-5965, with the 20 cm to 34 cm standard cuff or the 30 cm to 44 cm large 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.