dabl®Educational Trust

Declaration of Equivalence Form

DECLARATION OF BLOOD PRESSURE MEASURING DEVICE EQUIVALENCE

A SIGNED COPY WILL BE POSTED ON THE www.dableducational.org WEBSITE

SECTION A - Please complete all items.

l	Bill Huan Name of a C		ector		a Director of AViTA Corporation,		
hereby state that there are no differences that will affect blood pressure measuring accuracy between the							
Maker ^a BEURER GmbH			R GmbH	Address	Soeflinger Strasse 218, 89077 Ulm	, Germany	
Mar	oufacturer ⁶	AVITA (Corporation	Address	9F, NO.78, SEC.1, KWANG-FU RD Taipei City 24158 Taiwan R.O.C.	., SAN –Chung	District, New
Brar Bloo		Beurer leasuring de	evice for which validation is claimed.	Model^d If alternative	BM 51 model names are used, include all.		
blo	od pressi	ure mea	suring device and the valid	dated blo	ood pressure measuring device		
Mak	er ^a	AViTA (Corporation	Address	9F, NO.78, SEC.1, KWANG-FU RD Taipei City 24158 Taiwan R.O.C.	., SAN –Chung	District, New
Mar	oufacturer ^b	AViTA (Corporation	Address	9F, NO.78, SEC.1, KWANG-FU RD Taipei City 24158 Taiwan R.O.C.	., SAN –Chung	District, New
Brar Exist		AVITA I blood pres	sure measuring device.	Model ^d	BPM64		
wh	ich has p	revioush	y passed the ESH-2010 p	rotocol,	the results of which were published	l as follows:	
Kang Y.Y., Chen Q., Liu C.Y., Li Y. and Wang J.G. Validation of the AVITA BPM64 upper-arm blood pressure monitor for home blood pressure monitoring according to the European Society of Hypertension International Protocol revision 2010. 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 Oscillomet	ric Meas	urements Yes □	No ⊠	N/A ^e 🔲
		2	Algorithm for Auscultato	ry Meas	urements Yes □	No 🗌	N/A ^f ⊠
		3	Artefact/Error Detection		Yes 🗌	No 🛛	
		4	Microphone(s)		Yes □	No 🗌	N/A ^f ⊠
		5	Pressure Transducer		Yes □	No 🖂	
		6	Cuffs or Bladders		Yes 🗌	No 🖾	
		7	Inflation Mechanism		Yes □	No 🖂	
		8	Deflation Mechanism		Yes □	No ⊠	
_	Part II	9	Model Name or Number		Yes ⊠	No □	
		10	Casing		Yes 🔀	No 🗆	
		11	Display		Yes ⊠	No □	

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

Notes:

12

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a Provide the name and address of the actual maker of the device.

Printing Facilities

Power Supply

Other Facilities

Carrying/Mounting Facilities

Communication Facilities

Software other than Algorithm

b Provide the name and address of the legal manufacturer of the device, even if it is the same as that of the maker.

Memory Capacity/Number of stored measurements

- 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.

Yes 🖂

Yes 🔲

Yes 🛛

Yes 🗌

Yes 🔲

Yes 🛛

Yes 🗌

No 🗌

No 🗵

No 🗌

No 🔲

No 🗌

No 🗌

No 🛛

N/Ag 🖾

N/A^g

N/A^g

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SECTION B	differences between the devices must be described.	ached document. All
9) The model n	ame is different. for new device and validated device is BPM64	
10) The designs	s of the case are different.	
1 1) The size an	d displayed data are different.	
12) Carrying/M	lounting Facilities are different.	
14) has 2*100	memories	
17) New mode	l use AA*1.5V batteries or AC adaptor 100 – 240 V, 50 – 60 Hz, 0.5 A max	
SECTION O		
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	
	Completed DET9 Form	
	An image of the device for which equivalence is being sought	
	An image of the screen layout of validated device*	Warning Co.
	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 s	narately
	Screen layouts snown complete, and without obscuring labels of lines, in mandals need not be included s	eparately.
SECTION D	Complete all items, bar signatures and seal, online and print. Sign and seal it then send the original to our email a signed copy of this form, together with the manuals and images for both devices, to info@dabledu	
Signature of Di	rector Company Stamp/Seal	
Name <	Del Man 9 8/9. 2019	
Date		
Signature of W	litness	
	onathan Chen 2019.8.19.	

Address

9F, NO.78, SEC.1, KWANG-FU RD., SAN -Chung District, New Taipei City 24158 Taiwan R.O.C.

Device Equivalence Evaluation Form

Comparison of the Beurer BM51 with the AViTA BPM64

Devices – Item 9	Beurer BM51	AVITA BPM64
Pictures	(a) D3	
Display Image		18-38 0 18:88 M AVG M3 HISH HISH
Validation		ESH 2010
Category	Upper Arm Type Blood Pressure Monitor	Upper Arm Type Blood Pressure Monitor
Casing – Item 10	Dimensions approx. 138 mm x 103 mm x 44 mm (W * H *D) Ports Cuff Port DC Jack Port Features	Ports Cuff Port Features ABS Plactic part
	ABS plastic part Printing	ABS Plastic part Printing
Display – Item 11	LCD	LCD

Carrying/Mounting Facilities – Item 12	Storage Box	Storage Box
Software other than Algorithm – Item 13	N/A	N/A
Memory Capacity Item 14	2*100 times with date and time	1*90 times with date and time
Printing Facilities Item 15	Artwork logo, gift box and manual is different from AViTA BPM64 for different functions	Artwork logo, gift box and manual is different for different functions
Communication Facilities – Item 16	N/A	N/A
Power Supply Item 17	4 * AA Batteries AC adaptor Input: 100 – 240 V, 50 – 60 Hz, 0.5 A max Output: 6V DC, 600 mA	4 * AA Batteries
Other differences	Accuracy Blood Pressure Accuracy ± 3 mmHg Pulse Accuracy ± 5%	Accuracy Blood Pressure Accuracy ± 3 mmHg Pulse Accuracy ± 4%
Same Criteria	Measurement	Measurement
	Method Oscillometri	Method Oscillometri
	Ranges Cuff pressure 0–300 mmHg, systolic 60–255 mmHg, diastolic 40–200 mmHg,	Ranges Cuff pressure 0–300 mmHg, systolic 60–255 mmHg, diastolic 40–200 mmHg,
	Inflation Automatic inflation by internal pump	Inflation Automatic inflation by internal pump
	Deflation Automatic speed deflation system	Deflation Automatic speed deflation system
	Cuffs (Please state sizes and materials used) approx. 585 X 178 cm Bladder dimension: 230 X 125mm	Cuffs (Please state sizes and materials used) approx. 580 X 150 cm Bladder dimension: 230 X 125mm

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Page 2 of 5

Sensors

US-9111-006-S

Measurement Records

2*100 times with date and time

Measurements other than Blood Pressure

Pulse rate

Buttons/Switches

Power

START/POWER Button (on / off)

Measurement Records

Memory Recall Button - MEM

Function

Date and Time Set Button - SET

"-/+" function buttons - select the relevant value

Analysis

N/A

Event Marking

N/A

Communication

N/A

Display/Symbols/Indicators

Preparation

N/A

Measurement Procedure

Inflation symbol

Deflation symbol

Heartbeat symbol during deflation

Irregular Heartbeat symbol

Post Measurement

Sensors

US-9111-006-S

Measurement Records

1*90 times with date and time

Measurements other than Blood Pressure

Pulse rate

Buttons/Switches

Power

START/POWER Button (on / off)

Measurement Records

Memory Recall Button - MEM

Function

Date and Time Set Button – SET Mode (Alarm) Button - Mode

, ,

Analysis

N/A

Event Marking

N/A

Communication

N/A

Display/Symbols/Indicators

Preparation

N/A

Measurement Procedure

Inflation symbol

Deflation symbol

Heartbeat symbol during deflation

Irregular Heartbeat symbol

Post Measurement

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	Systolic blood pressure	Systolic blood pressure
	Diastolic blood pressure	Diastolic blood pressure
	Pulse rate	Pulse rate
	WHO indicator	WHO indicator
	Measurement Records	Measurement Records
	Memory recall number	Memory recall number
	Date and Time	Date and Time
	Date and Time	Date and Time
	Power	Power
	Low Battery detection symbol	Low Battery detection symbol
	Function	Function
	Average	Average
		Alarm
	Communication	Communication
	N/A	N/A
	Features	Features
	N/A	N/A
	Not described	Not described
	N/A	
		Algorithms
	Algorithms	Averages and Differences
	Averages and Differences	Average of the last 3 measurements
	Average of all measurement	
	Average morning values of the last seven days measurements	Diagnostic
	between 5:00AM and 9:00AM	N/A
	Average evening values of the last seven days measurements	
	between 6:00PM and 8:00PM	Functions
	between 0.00FW and 8.00FW	N/A
	Diagnostic	Communication
	N/A	N/A
	Functions	IN/C
	N/A	
	Communication	
	N/A	
Comparable Criteria		

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Comments		
Recommendation	Recommended	
Date	25 th September 2020	

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Page 5 of 5