Declaration of Equivalence Form

DECLARATION OF BLOOD PRESSURE MEASURING DEVICE EQUIVALENCE 2013

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

| I Bill Huar Name of a C | ng, Company Director | | a Director of AViTA Corporation,, Company name |
|---|---|--------------------|--|
| hereby state | e that there are no differences tha | at will aff | ect blood pressure measuring accuracy between the |
| Maker ^a | Medel | Address | Medel International, Via Villapizzone 26, 20156, Milan, Italy |
| Manufacturer ^b | Globalcare | Address | 7 th Building 39 Middle Industrial Main Road European Industrial Zone, Xiaolan Town, Zhongshan City Guangdong Province 52815 CHINA. |
| Brand ^c Medel Model ^d Medel Control 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 AVITA Corporation Address 9F, NO.78, SEC.1, KWANG-FU RD., SAN –Chung District, New Taipei City 24158 Taiwan R.O.C. | | | |
| Manufacturer ^b | AViTA Corporatio | Address | 9F, NO.78, SEC.1, KWANG-FU RD. , SAN –Chung District, New Taipei City 24158 Taiwan R.O.C. |
| Brand^c Existing validated | AVITA d blood pressure measuring device. | Model ^d | BPM63S |

which has previously passed the ESH-2010 protocol, the results of which were published as follows:

Kang Y-Y,Zeng W-F, Liu M, Li Y,and Wang J-G. Validation of the AVITA BPM63S upper arm blood pressure monitor for home blood pressure monitoring according to the European Society of Hypertension International Protocol revision 2010.Blood Pressure Full reference

The only differences between the devices involve the following components:

Tick one box for each item 1–18.

| | | | | | 0 |
|---------|----|---|-------|------|--------------------|
| Part I | 1 | Algorithm for Oscillometric Measurements | Yes 🗌 | No 🖂 | N/A ^e 🗌 |
| | 2 | Algorithm for Auscultatory Measurements | 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 🗌 | |
| | 12 | Carrying/Mounting Facilities | Yes 🖂 | No 🗌 | |
| | 13 | Software other than Algorithm | Yes 🖂 | No 🗌 | |
| | 14 | Memory Capacity/Number of stored measurements | Yes 🖂 | No 🗌 | |
| | 15 | Printing Facilities | Yes 🗌 | No 🗌 | $N/A^g \boxtimes$ |
| | 16 | Communication Facilities | Yes 🗌 | No 🗌 | $N/A^g \boxtimes$ |
| | 17 | Power Supply | Yes 🖂 | No 🗌 | |
| | 18 | Other Facilities | Yes 🖂 | No 🗌 | N/A ^g |
| | | | | | |

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

Notes: Provide the name and address of the actual maker of the device

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

Provide the name of the brand under which it is sold, even if it is the same as that of the manufacturer or maker. С

Provide the model name. If alternative or internal model names are used, include all. Each device must be uniquely identifiable. d

Only tick N/A (Not Applicable) if neither device measures blood pressure using the oscillometric method. е

f Only tick N/A (Not Applicable) if neither device measures blood pressure using the auscultatory method.

Only tick N/A (Not Applicable) if neither device provides printing, communication or other facilities, as appropriate. g

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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.

| 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 st | |
| | * 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 capy of this form, together with the manuals and images for both devices, to info@dableducational.org.

| Signature of Director | <u> </u> | amp/Seal |
|-----------------------|---|--|
| Name | Bill Hueng | |
| Date | 2016.03.15 | |
| Signature of Witness | Jonathan Chen | |
| Name | Jonathan Chen | |
| Address | 9F, NO.78, SEC.1, KWANG-FU RD. , SANChung Distr | ict, New Taipei City 24158 Taiwan R.O.C. |

Device Equivalence Evaluation Form

| Devices | Medel Control | AVITA BPM63S |
|------------|--|--|
| Pictures | readdal 1 creating to the static 1 creatin | Nolox Participation Participat |
| Display | < #38:88 < 38/38 < 38/38 < 38/38 < 38/38 < 38/38 < 38/38 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 6 6 7 < 6 7 | IS-38 Q IS:88 AM IS-38 Q IS IS-38 |
| Validation | | ESH 2010 |
| Category | Arm Type Blood Pressure Monitor | Arm Type Blood Pressure Monitor |
| Comparison | Dimension 112 * 110 * 58 mm (W * H *D) Weight 219g(Excluding batteries) Cuff Size 22-42cm | Dimension 113 * 140 * 57 mm (W * H *D) Weight 275g(Excluding batteries) Cuff Size 22-33cm |

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| | Measurement | Measurement | |
|---------------|--|--|--|
| | Accuracy Blood Pressure Accuracy \pm 3 mmHg | Accuracy Blood Pressure Accuracy ± 3 mmHg | |
| | Pulse Accuracy \pm 5% | Pulse Accuracy \pm 4% | |
| | | | |
| Same Criteria | Measurement | Measurement | |
| | Method Oscillometric | Method Oscillometric | |
| | Oscillometric | Oscillometric | |
| | Ranges | Ranges | |
| | Cuff pressure 0 -300 mmHg | Cuff pressure 0 -300 mmHg | |
| | Systolic 50 mmHg – 280 mmHg | Systolic 50 mmHg – 280 mmHg | |
| | Diastolic 30 mmHg – 200 mmHg | Diastolic 30 mmHg – 200 mmHg | |
| | Inflation | Inflation | |
| | Automatic inflation by internal pump | Automatic inflation by internal pump | |
| | Deflation | Deflation | |
| | Automatic speed deflation system | Automatic speed deflation system | |
| | Cuffs (Please state sizes and materials used) | Cuffs(Please state sizes and materials used) | |
| | 22-42 cm | 22-33 cm | |
| | Bladder dimension: 120x232mm | Bladder dimension: 120x232mm | |
| | Sensors | Sensors | |
| | US-9111-006-S | US-9111-006-S | |
| | Measurement Records | Measurement Records | |
| | 4*30 times with date and time | 1*60 times with date and time | |
| | Measurements other than Blood Pressure | Measurements other than Blood Pressure | |
| | Pulse rate | Pulse rate | |
| | Buttons/Switches | Buttons/Switches | |
| | Power | Power | |
| | START/POWER Button (on / off) | START/POWER Button (on / off) | |
| | Measurement Records | Measurement Records | |

| Memory Recall Button - MEM | Memory Recall Button - MEM |
|---|--|
| | |
| Function | Function |
| Date and Time Set Button – SET | Date and Time Set Button – SET |
| +/- BUTTONS TO ROLLS THE MEMORIES AND SETTING TIME Analysis | Mode (Alarm) Button - Mode Analysis |
| N/A | N/A |
| | |
| Event Marking | Event Marking |
| N/A | N/A |
| Communication | Communication |
| N/A | N/A |
| | |
| Display/Symbols/Indicators | Display/Symbols/Indicators |
| Preparation | Preparation |
| N/A | N/A |
| Measurement Procedure | Measurement Procedure |
| Inflation symbol | Inflation symbol |
| Deflation symbol | Deflation symbol |
| Heartbeat symbol during deflation | Heartbeat symbol during deflation |
| Irregular Heartbeat symbol | Irregular Heartbeat symbol |
| Post Measurement | Post Measurement |
| 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 N/A | Features N/A |
| N/A | N/A |
| Not described | Not described |
| Algorithms | |
| Averages and Differences | Algorithms |
| Average morning values of the last seven days measurements | Averages and Differences |
| between 5:00AM and 9:00AM | Average of the last 3 measurements |
| Average evening values of the last seven days measurements | |
| between 6:00PM and 8:00PM | |
| Diagnostic | Diagnostic |
| N/A | N/A |
| N/A | |
| Functions | Functions |
| N/A | N/A |
| | Communication |
| Communication | N/A |
| N/A | |
| Casing | Casing |
| Display | Display |
| LCD | LCD |
| | Ports |
| Ports | Cuff Port |
| Cuff Port | DC Jack *AC adapter is optional |
| | De Jack Ae adapter is optional |
| Power | Power |
| 4 * AA Batteries | 4 * AA Batteries |
| | |
| Features | Features |
| N/A | N/A |
| | |
| | |

| Comparable Criteria | |
|---------------------|--|
| | |

| Comments | |
|----------------|--------------|
| Recommendation | ecommended |
| Date | 4 April 2016 |