# dabl®Educational Trust

# **Validation Study Registration Form**

**Foracare** 

This form should be completed by the manufacturer or on behalf of the manufacturer.

Study Investigator Details

1401

Study Ref.

Do Not Fill

Name

**Dauyuan Ding** 

Address

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#### **Test Device Details**

Maker TaiDoc Manufacturer

**Diamond CUFF BP Brand** 

**Test N GO** Model Internal Model Number TD-3140 (P80)

Initiator Manufacturer Details if "Other"

If not initiated by the manufacturer, did the manufacturer agree to the study?

Select the correct option on each of the following or, if required, complete the explanation beside "Other".

Location

**Upper Arm** 

Details if "Other"

Method

Oscillometry

Details if "Other"

Purpose

Self/Home Measurement

Details if "Other"

Operation

Automatic

Details if "Other"

Automatic: Cuff inflation, deflation and blood pressure determination are fully performed by the device automatically; Semi-automatic: Blood pressure determination is performed automatically but cuff inflation and deflation need manual operation; Manual: Cuff pressure control and blood pressure determination are all performed by manual operation.

Cuff details including arm circumference ranges (as recommended by the device manufacturer).

Cuffs Small Adult: cm to cm Standard Adult: cm to

> Larae Adult: cm to Other Wide Ranged: 24 cm to 43 cm

Wrist Cuff cm to Wrist Support Method cm

#### Other features of the device (about 100 words).

The device of Foracare Diamond CUFF BP Test N GO blood pressure monitoring system automatically inflates the cuff and displays the readings. The device of Foracare Diamond CUFF BP Test N GO blood pressure monitoring system designs one button for easy to operate, and make sure patient can easily to understand the readings of blood pressure from the display of the monitor.

## Agreement

I agree to the publication of the results regardless of whether or not they are favourable to the device.

Signed

Company Stamp or Seal

Name

Jim Jan

ForaCare Suisse AG Neugasse 55, 9000 St. Gallen

2014/03/20 Date

Switzerland

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# dabl®Educational Trust

# Validation Study Result Form

Please complete Section 1 to Section 3 of this form and return it to dabl\*Educational Trust with copies of the validation plots and a digital photograph of the device used for the study showing its front face. Please follow the instructions for each section. The requirements for each table entry are described, by box number, under the respective table.

Study Ref. 1401 Do Not Fill

Maker	TaiDoc	Manufacturer	Foracare		
Brand	Diamond CUFF BP	Model	Test N GO		
Investigator	Dauyuan Ding				
Signed	Darth (1.			Date	2014/03/:

Section 1: Methodology

# **Familiarisation**

A brief description of the familiarisation session should be provided. Any difficulties should be reported.

Several pre-test were performed in house before clinical test started. No problems were encountered.

#### Recruitment

The population should be outlined and the method of selecting the sample should be described. Difficulties in recruitment should be described and how they were overcome.

Population Details if "Other"

#### Procedure

children is included.

 $\times$ Two observers with an independent supervisor X Observers blinded from each other's readings and from the device readings The European Society of Hypertension International Protocol revision 2010 for the validation of blood pressure X measuring devices in adults was followed precisely. Enter protocol adjustments, as necessary, when the study population is not general with sex, age and blood pressure distribution stated in detail. These adjustments should be justified, with references where possible. Because children and

adolescents have wide range of body size and blood pressure levels, the sample size for a validation study should depend on the study inclusion criteria. Thus, for example, a 33-subject study would be appropriate only if a narrow age range of

This form is intended for use only in connection with blood pressure monitor validation studies carried out in accordance with the protocol of the European Society of Hypertension: O'Brien E et al. 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.

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Form 05T28 130410

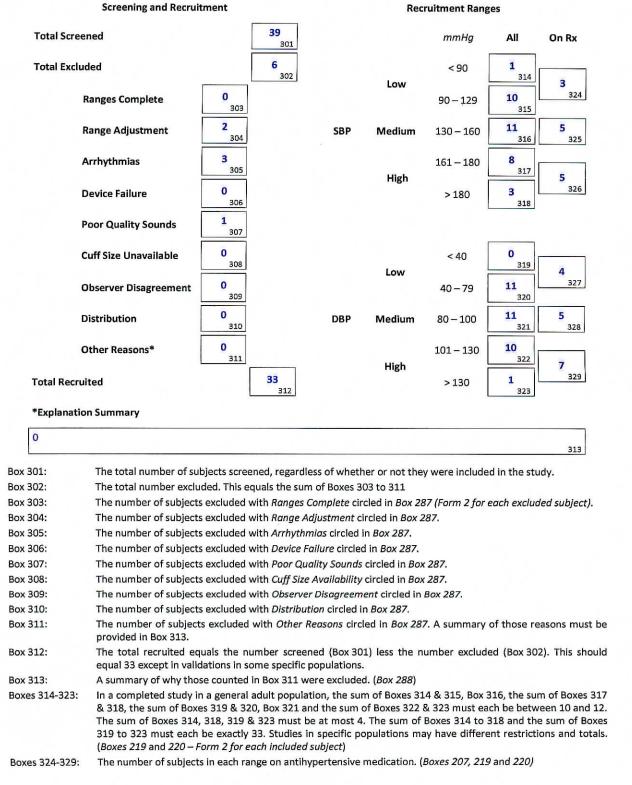
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## Section 2: Results

Note 1: The data from Form 2 – Subject Data for each subject should be analysed so that the results on this form can be completed. All references to boxes 201-289 refer to values obtained from all of the Forms 2 from the relevant subjects.

**Table 1: Screening and Recruitment Details** 



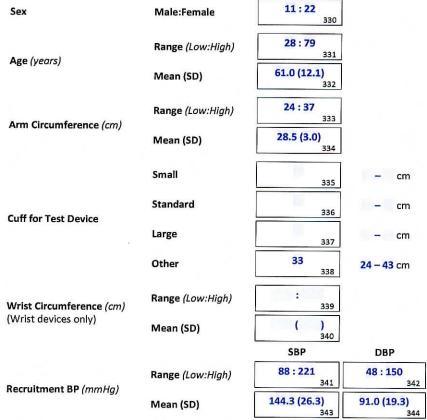
Box 340:

Boxes 341-342:

Boxes 343-344:

(Boxes 217 and 218)

**Table 2: Subject Details** 



Note 2:	The values in Boxes 314–380 refer only to the final recruited subjects, each of whom contributes SBP and DBP measurements for analysis. Excluded subjects are not included in any of this analysis.
Box 330:	Enter the number of males, a colon and the number of females. They should total 33 except in validations in some specific populations. If the minimum requirements (10 for a general population) are not met, subjects must be replaced as necessary. (Box 206)
Box 331:	Enter the age of the youngest subject, a colon and the age of the oldest subject e.g. 31:74. Subjects outside the required range (25 and over for a general population) are not permitted. (Box 205)
Box 332:	Enter the mean and, in parentheses, the SD of the subject ages. Values should be rounded to one decimal place e.g. 52.3 (11.9). (Box 205)
Box 333:	Enter the smallest arm circumference, a colon and the largest arm circumference e.g. 24:34. (Box 208)
Box 334:	Enter the mean and, in parentheses, the SD of the subject arm circumferences. Values should be rounded to one decimal place e.g. 29.0 (3.1). (Box 208)
Box 335:	If a small cuff was supplied, enter the number of subjects on whom it was used. If it was not supplied, enter an "X". Enter the arm sizes for which it is recommended beside it. (Box 209)
Box 336:	Enter the number of subjects on whom a standard (or medium) cuff was used. Enter the arm sizes for which it is recommended beside it. (Box 209)
Box 337:	If a large cuff was supplied, enter the number of subjects on whom it was used. If it was not supplied, enter an "X". Enter the arm sizes for which it is recommended beside it. (Box 209)
Box 338:	If a different size cuff was supplied, enter the number of subjects in whom it was used. If no such cuff was supplied, enter an "X". Enter the arm sizes for which it is recommended beside it. ( <i>Box 209</i> )
Box 339:	Enter the smallest wrist circumference, a colon and the largest wrist circumference e.g. 15:22. (Applicable only for wrist devices) (Box 210)

one decimal place e.g. 18.1 (2.3). (Applicable only for wrist devices) (Box 210)

should be rounded to one decimal place e.g. 140.4 (20.3). (Boxes 217 and 218)

Enter the mean and, in parentheses, the SD of the subject wrist circumferences. Values should be rounded to

Enter the lowest pressure, a colon and the highest pressure from BPA measurements only e.g. 104:180.

Enter the mean and, in parentheses, the SD of the subject pressures from BPA measurements only. Values

#### Table 3: Distribution

This section analyses the distribution of comparative measurements.

SBP		DBP	
Overall Range (mmHg) Low:High	<b>88 : 221</b> 345	Overall Range (mmHg) Low:High	<b>48 : 150</b> 350
Low (< 130 mmHg)	<b>28</b> 346	Low (< 80 mmHg)	<b>32</b> 351
Medium (130 mmHg – 160 mmHg)	<b>42</b> 347	Medium (80 mmHg – 100 mmHg)	<b>41</b> 352
High (> 160 mmHg)	<b>29</b> 348	High (> 100 mmHg)	<b>26</b> 353
Maximum Difference	<b>14</b> 349	Maximum Difference	<b>15</b> 354

Box 345: Enter the lowest pressure, a colon and the highest SBP from the observer measurements. (Boxes 281, 283 and 285)

Boxes 346-348: The observer measurements (three per subject) for SBP are categorised similarly to the recruitment ranges. Enter the counts of measurements falling into each range. These must total 99. (Boxes 281, 283 and 285)

Box 349 Subtract the smallest value from Boxes 346 to 348 from the largest one and enter the result.

Box 350: Enter the lowest pressure, a colon and the highest DBP from the observer measurements. (Boxes 282, 284 and

The observer measurements (three per subject) for DBP are categorised similarly to the recruitment ranges. Boxes 351-353: Enter the counts of measurements falling into each range. These must total 99. (Boxes 282, 284 and 286)

Box 354: Subtract the smallest value from Boxes 351 to 353 from the largest one and enter the result.

Note 3: In order to ensure a uniform distribution, there must be at least 22 measurements and at most 44 measurements (Boxes 346 to 348 and 351 to 353) in each of the low, medium and high ranges and the maximum differences (Boxes 349 and 354) must be at most 19. If not, further recruitment will be necessary. Subjects to be excluded will be those whose pressures drifted from recruitment pressures.

Note 4: The overall SBP range must be from ≤ 100 mmHg to ≥ 170 mmHg and the overall DBP range must be from ≤ 50 mmHg to ≥ 120 mmHg. If not, further recruitment will be necessary. Subjects to be excluded will be the last recruited within the relevant ranges.

Note 5: The minimum number of replacements should take place. If a subject is replaced for either of these reasons, circle Distribution in Box 287 of Form 2 for that subject.

In validations carried out in specific populations requiring more than 33 subjects but with similar blood pressure Note 6: distributions, similar proportions should be used. If the blood pressure distribution in the specific population differs from the standard distribution, ignore this table but comment on the distribution in the discussion.

## **Table 4: Observer Differences**

This section is for the differences in pressures between the two observers.

		SBP (mmHg)	DBP (mmHg)	Repeated	
Observer 2 – Observer 1	Range Low:High	<b>-4 : 4</b> 355	<b>-4:4</b> 356		
Observer 2 – Observer 1	Mean (SD)	- <b>0.1 (2.1)</b>	- <b>0.2 (2.2)</b> 358	0 359	

Boxes 355-356 Enter the lowest difference, a colon and the highest difference between the observers. Include the signs e.g. -3:+4. (Boxes 247, 249, 251 and 253 and Boxes 248, 250, 252 and 254). If the range is outside -4:+4, then this is a violation. Relevant subjects should be excluded, by reason of Observer Disagreement, and replaced.

Boxes 357-358 Enter the mean and, in parentheses, the SD of the observer differences. Values should be rounded to one decimal place e.g. 0.3 (1.2). (Boxes 247, 249, 251 and 253 and Boxes 248, 250, 252 and 254)

Enter the number of measurements that were repeated in the included subjects because observers were more Boxes 359 than 4 mmHg apart.

## **Table 5: Validation Results**

Part 1		≤ 5 mmHg	≤ 10 mmHg	≤ 15 mmHg	Grade 1	Mean (mmHg)	SD (mmHg)
Pass	Two of	73	87	96			
Requirement	All of	65	81	93			
Achieved	SBP	<b>75</b> 360	<b>95</b>	99 362	Pass 363	<b>1.6</b> 364	<b>4.8</b> 365
	DBP	<b>76</b> 366	<b>92</b> 36	<b>99</b> 368	Pass 369	<b>1.8</b> 370	<b>4.8</b> 371
Part 2		2/3 ≤ 5 mm	Hg 0	/3 ≤ 5 mmHg	Grade 2		Grade 3
Pass Requirement		≥ 24		≤3			
	SBP	26	372	<b>2</b> 373	Pass 374		Pass 375
Achieved	DBP	29	376	<b>2</b> 377	Pass 378		Pass 379
Part 3							Result
							Pass 380

	360
Note 7: In o	order for the device to pass, <u>all</u> requirements must be fulfilled. A fail in any part will result in an overall fail.
Box 360:	Enter the number of SBP differences (at most 99) between observer and device measurements falling within 5 mmHg. (The total number of <i>Boxes 273</i> , 275 and 277 circled A in the 33 subjects)
Box 361:	Enter the number of SBP differences (at most 99) between observer and device measurements falling within 10 mmHg. (The total number of <i>Boxes 273, 275</i> and <i>277</i> circled A or B in the 33 subjects)
Box 362:	Enter the number of SBP differences (at most 99) between observer and device measurements falling within 15 mmHg. (The total number of <i>Boxes 273, 275</i> and <i>277</i> circled A, B or C in the 33 subjects)
Box 363:	If Boxes 360, 361 and 362 fulfil the Pass requirements, then this is "Pass"; otherwise, it is "Fail".
Boxes 364-365	Enter the mean and standard deviation respectively of the 99 SBP differences between observer and device measurements. (Use data from circled <i>Boxes 261</i> or <i>267</i> , <i>263</i> or <i>269</i> and <i>265</i> or <i>271</i> )
Box 366:	Enter the number of DBP differences (at most 99) between observer and device measurements falling within 5 mmHg. (The total number of <i>Boxes 274, 276</i> and <i>278</i> circled A in the 33 subjects)
Box 367:	Enter the number of DBP differences (at most 99) between observer and device measurements falling within 10 mmHg. (The total number of <i>Boxes 274, 276</i> and <i>278</i> circled A or B in the 33 subjects)
Box 368:	Enter the number of DBP differences (at most 99) between observer and device measurements falling within 15 mmHg. (The total number of <i>Boxes 274, 276</i> and <i>278</i> circled A, B or C in the 33 subjects)
Box 369:	If Boxes 366, 367 and 368 fulfil the Pass requirements, then this is "Pass"; otherwise, it is "Fail".
Boxes 370-371	: Enter the mean and standard deviation respectively of the 99 DBP differences between observer and device measurements. (Use data from circled <i>Boxes 262</i> or <i>268</i> , <i>264</i> or <i>270</i> and <i>266</i> or <i>272</i> )
Box 372:	Enter the number of subjects (at most 33) with two or three of the absolute differences between observer and device SBP measurements within 5 mmHg. (Box 279 is 2 or 3)
Box 373:	Enter the number of subjects (at most 33) with none of the absolute differences between observer and device SBP measurements within 5 mmHg. ( <i>Box 279</i> is 0)
Box 374:	If Boxes 372 and 373 fulfil the Pass requirements, then this is "Pass"; otherwise, it is "Fail".
Box 375:	If Boxes 363 and 374 are both "Pass", then this is "Pass"; otherwise, it is "Fail".
Box 376:	Enter the number of subjects (at most 33) with two or three of the absolute differences between observer and device DBP measurements within 5 mmHg. (Box 280 is 2 or 3)
Box 377:	Enter the number of subjects (at most 33) with none of the absolute differences between observer and device DBP measurements within 5 mmHg. (Box 280 is 0)
Box 378:	If Boxes 376 and 377 fulfil the Pass requirements, then this is "Pass"; otherwise, it is "Fail".
Box 379:	If Boxes 369 and 378 are both "Pass", then this is "Pass"; otherwise, it is "Fail".
Box 380	If Boxes 375 and 379 are both "Pass", then this is "Pass"; otherwise, it is "Fail".
Note 8: In	validations carried out in specific populations requiring more than 33 subjects, proportionally equivalent passing

criteria should be used.

# Section 3: Closeout

#### **Plots**

Include the plots with this document. Confirm that they comply with the requirements

X X-axis: Range 80 mmHg to 190 mmHg Reference lines at 130 mmHg and 160 mmHg  $\times$  $\times$ Y-axis: Range -30 mmHg to 30 mmHg  $\times$ Reference lines every 5 mmHg from -15 mmHg to 15 mmHg X DBP X-axis: Range 30 mmHg to 140 mmHg X Reference lines at 80 mmHg and 100 mmHg Y-axis: Range -30 mmHg to 30 mmHg  $\times$ Reference lines every 5 mmHg from -15 mmHg to 15 mmHg  $\bowtie$ 

# **Image**

Include a digital photograph, of the device used in the study, with this document. The photograph should show the front face of the device. Use a plain background.

A photograph, of the device used in the study, is included  $\times$ 

#### Discussion

For tight blood pressure control, people with high blood pressure choose a convenient device provides the reliable measurement is important. The device of Foracare Diamond CUFF BP Test N GO automatically inflates the cuff and displays the readings. The display on the monitor is easy to read and understand. The clinically outcomes show an accurate and consistent results of Foracare Diamond CUFF BP Test N GO blood pressure monitoring system to detect hypertension.

Recruitment of subjects with high DBP range was difficult and accounted more screened subjects. However, the distribution conditions from 48 mmHg to 150 mmHg of DBP ranges were fulfilled the requirement of EHS protocol.

#### Conclusion

The conclusion as to whether the device is accurate for use in the population should be stated. If the results are particularly sensitive to correct use (e.g. most wrist devices) then this caution must be stated.

The device has passed to the criteria of ESH 2010 protocol, it is recommended for home use. The device of Foracare Diamond CUFF BP Test N GO blood pressure monitoring system provides an accurate and consistent results for most patients can more easily to detect hypertension.

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# **Bubbles Plot**

