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DECLARATION OF BLOOD PRESSURE MEASURING DEVICE EQUIVALENCE 2013

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

SECTION A	- Please complete all items.		
I Kevin Ta Co.,Ltd ,	n, Company Director		a Director of Guangdong Transtek Medical Electronics
		ıt will aff	ect blood pressure measuring accuracy between the
, Maker ^a	Greater Goods, LLC.	Address	4427 Chouteau Ave. St. Louis, MO 63110
Manufacturer⁵	Transtek	Address	Zone A, No.105 ,Dongli Road, Torch Development District,
			Zhongshan, 528437, Guangdong, China
Brand^c Blood pressure n	Greater goods neasuring device for which validation is claimed.	Model ^d If alternativ	0040 e model names are used, include all.
blood pressure measuring device and the validated blood pressure measuring device			
Maker ^a	Guangdong Transtek Medical	Address	Zone A, No.105 ,Dongli Road, Torch Development District,
	Electronics Co.,Ltd		Zhongshan, 528437, Guangdong, China
Manufacturer ^b	Guangdong Transtek Medical	Address	Zone A, No.105 ,Dongli Road, Torch Development District,

Brand^c TRANSTEK Existing validated blood pressure measuring device.

Electronics Co.,Ltd

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

Modeld

Title: Validation of Transtek blood pressure monitor TMB-1491 for self-measurement according to the European Society of Hypertensio

TMB-1491

Zhongshan, 528437, Guangdong, China

Authors: Huiyong Tian, Sijian Zeng, Xiaoyan Zhong, Wei Gong and Wenju Liu

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 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 🔀
	16	Communication Facilities	Yes 🗌	No 🗌	N/A ^g 🔀
	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: a Provide the name and address of the actual maker of the device.

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

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Declaration of Equivalence Form

- 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.

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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.

See attached document

SECTION C	Please o	Please check that the following are included with the application		
	A man	ual for the validated device	\square	
	A man	manual for the device for which equivalence is being sought		
	An ima	age of the validated device		\boxtimes
	An ima	age of the device for which equivalence	e is being sought	\boxtimes
	An ima	age of the screen layout of validated de	evice*	\boxtimes
	An ima	age of the screen layout of the device f	or which equivalence is being sought*	\boxtimes
	* Screer	n layouts shown complete, and without obscurin	g labels or lines, in manuals need not be included s	eparately.
SECTION D	•	signed copy of this form, together with the man	orint. Sign and seal it then send the original to our a uals and images for both devices, to info@dabledue	
Signature of Di	rector	Vigia Qi	Company Stamp/Seal	
Name		Kevin Tan		
Date		March 26,2021		
Signature of W	'itness	Kevin Tan		
Name		Viya Qi		

Address Zhongshan, 528437, Guangdong, China



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Device Equivalence Evaluation Form

Comparison of the Greater Goods 0040 with the TRANSTEK TMB-1491

Devices – Item 9	Greater Goods 0040	TRANSTEK TMB-1491
Pictures		
Display Image	TIME / DATE OPUL / MIN	
Validation	Upper arm device for self measurement of blood pressure	ESH 2010
Category	Upper arm device for self measurement of blood pressure	Upper arm device for self measurement of blood pressure
Casing – Item 10	Dimensions 140mm*130mm*55.6mm	Dimensions 110mm*100mm*41mm
	Ports Cuff port	Ports Cuff port and DC power port
	Features Cuff Trademark printing	Features Cuff and AC adaptor connectors Model name printing

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	Button printing	Button printing
Display – Item 11	LCD	LCD
Carrying/Mounting Facilities – Item 12	Dimensions 195*140*90mm	Dimensions 182*130*80MM
Software other than Algorithm – Item 13	Dual Users 60 sets memories/per user AHA indicator mmHg unit mmHg unit	Dual Users 60 sets memories/per user WHO indicator mmHg unit
Memory Capacity Item 14	60 sets memories/per user	60 sets memories/per user
Printing Facilities Item 15	N/A	N/A
Communication Facilities – Item 16	N/A	N/A
Power Supply Item 17	6V DC 4×AAA batteries	6V DC Jack 4*AAA batteries
Other differences	Other Details on Equivalent device that are different to Validated device N/A	Other Details on Validated device that are different to Equivalent device N/A
Same Criteria	Measurement Accuracy Pressurewithin±3mmHg(0.4kPa) Pulse value: ±5% Max Method Oscillographic testing mode Ranges SYS: 60mmHg~230mmHg (8.0kPa~30.7kPa) DIA: 40mmHg~130mmHg (5.3kPa~17.3kPa) Pulse value: (40-199)beat/minute	Measurement Accuracy Pressure:5°C-40°C within±3mmHg(0.4kPa) Pulse value: ±5% Method Oscillographic testing mode Ranges R SYS: 60mmHg~230mmHg (8.0kPa~30.7kPa) DIA: 40mmHg~130mmHg (5.3kPa~17.3kPa) Pulse value: (40-199) beat/minute
	Inflation Automatic inflation	Inflation Automatic inflation
	Deflation Automatic deflation	Deflation Automatic deflation

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	Sensors PSG010S Measurements other than Blood Pressure Pluse rate Buttons/Switches Up/Down Delete Buttons Start/Stop & Select Button	Sensors PSG010S Measurements other than Blood Pressure Pluse rate Buttons/Switches SYS botton DIA button Start/Stop & Select Button
	Display/Symbols/Indicators Preparation Automatic Zero setting	Display/Symbols/Indicators Preparation Automatic Zero setting
	Measurement Procedure Inflation symbol Pressure value indication Current time	Measurement Procedure Inflation symbol Pressure value indication Current time
	Measurement Records Systolic blood pressure (SYS) Diastolic blood pressure (DIA) Pulse rate Measurement time Memory Query symbol	Measurement Records Systolic blood pressure (SYS) Diastolic blood pressure (DIA) Pulse rate Measurement time Memory Query symbol
	Power Low power	Power Low power
	Features Measuring during inflation	Features Measuring during inflation
	Algorithms Equivalent device has the identical measurement algorithm as the validated device.	Algorithms Equivalent device has the identical measurement algorithm as the validated device.
Comparable Criteria	<i>Measurement</i> Cuffs (Please state sizes and materials used) About 22cm-42cm, large bore connector, Dacron material	Measurement Cuffs (Please state sizes and materials used) About22cm or 32 or 22cm-42cm, polyester
	Measurement Records 60 sets/per user Display/Symbols/Indicators Post Measurement Systolic blood pressure (SYS)	Measurement Records 60 sets/per user Display/Symbols/Indicators

Diastolic blood pressure (DIA) Pulse rate Measurement time	Post Measurement Systolic blood pressure (SYS) Diastolic blood pressure (DIA) Pulse rate
<i>Function</i> Measure blood pressure and heart rate Recall measurement records Delete measurement records	Measurement time Function Measure blood pressure and heart rate Recall measurement records Delete measurement records

Comments	
Recommendation	Recommended
Date	April 2021