



Home sphygmomanometers can help in the control of blood pressure: a nationwide field survey

Tekin Akpolat¹ · Mustafa Arici² · Sule Sengul³ · Ulver Derici⁴ · Sukru Ulusoy⁵ · Sehsuvar Erturk³ · Yunus Erdem²

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Abstract

Home blood pressure monitoring (HBPM), which integrates patients into their treatment program, is a self-management tool. The prevalence of home sphygmomanometer ownership and patient compliance with HBPM guidelines are not well known, especially in developing and underdeveloped countries. The aims of this study were to measure the prevalence of home sphygmomanometer ownership among hypertensive subjects through a nationwide field survey (Patent2), to investigate the validation of sphygmomanometers and consistency of the user arm circumference and cuff size of the upper-arm device owned, as well as to compare blood pressure (BP) readings between hypertensive subjects who have or do not have a sphygmomanometer. Sample selection was based on a multistratified proportional sampling procedure to select a nationally representative sample of the adult population ($n = 5437$). Of 1650 hypertensive subjects, 332 (20.1%) owned a device, but the percentage of patients who owned a sphygmomanometer was 28.8% among patients who were aware of their hypertension (260/902). The usage of wrist devices and nonvalidated devices is common, and selection of an appropriate cuff size is ignored. Linear-regression analysis showed that owning a BP monitor is associated with decreases of 3.7 mmHg and 2.8 mmHg for systolic and diastolic BPs, respectively. Many patients do not own a sphygmomanometer. The decrease of systolic and diastolic BPs among BP monitor owners is a striking finding. The implementation of a hypertension care program consisting of sphygmomanometer reimbursement and training of patients in its use for HBPM might be cost-effective.

Introduction

Home blood pressure monitoring (HBPM) integrates patients into their treatment program and is a common self-management tool, especially in developed countries [1–5]. Substantial evidence regarding the usefulness of HBPM in the management of hypertension (HT) has accumulated over the last two decades [6–9], and HBPM is recommended in HT guidelines [10–14]. Home blood pressure

(BP) monitoring can also be supported by pharmacy assistance, telemonitoring or self-management programs [15–17]. A recent systematic review found HBPM to be superior to office measurements in diagnosing uncontrolled HT, assessing the efficiency of antihypertensive agents, and improving patients' compliance and HT control [18].

An accurate and validated [19–22] sphygmomanometer, the correct measurement of BP and adherence to the current guidelines regarding when and how BP should be measured at home settings are the essentials of reliable HBPM. However, the prevalence of home sphygmomanometer ownership and patient awareness and compliance with the HBPM guidelines are not well known, especially in developing and underdeveloped countries. In Turkey, a developing country with a high prevalence of HT (30.3%), we conducted a survey (published in this journal) using computer-assisted telephone interviewing among 2747 hypertensive patients in 2011; of the respondents, 1281 (46.6%) had a home sphygmomanometer [23–26]. Computer-assisted telephone interviewing has some limitations; therefore, we were unable to investigate the

✉ Tekin Akpolat
tekinakpolat@yahoo.com

¹ Istinye University Liv Hospital, Istanbul, Turkey

² Hacettepe University Faculty of Medicine, Ankara, Turkey

³ Ankara University Faculty of Medicine, Ankara, Turkey

⁴ Gazi University Faculty of Medicine, Ankara, Turkey

⁵ Karadeniz Technical University Faculty of Medicine, Trabzon, Turkey

validation of the sphygmomanometers, the consistency of the arm circumference of the user and the cuff size of the upper-arm device owned and the possible beneficial effects of having a sphygmomanometer. The aims of this study were to measure the prevalence of home sphygmomanometer ownership among hypertensive subjects through a nationwide field survey [27], to investigate the validation of sphygmomanometers, to determine the consistency of user arm circumference and the cuff size of the upper-arm device owned and to compare BP readings between hypertensive subjects who have or do not have a sphygmomanometer.

Methods

Research population and sampling

The PatenT2 study [27] was designed, directed and fully supported by the Turkish Society of Hypertension and Renal Diseases. PatenT2 sample selection was based on a multistratified proportional sampling procedure to select a nationally representative sample of the adult population over 18 years of age ($n = 5437$). During the home visits, a standard face-to-face interview questionnaire was administered to collect data on social demographics and behavioral characteristics, and a minimum of three consecutive BP measurements were taken. A total of 5437 individuals [2704 men (49.7%) and 2733 women (50.3%), 74.3% urban and 25.7% rural residents] were randomly selected from 26 cities, and all completed the face-to-face interview questionnaire, and BP and anthropometric measurements were taken over 92 days.

Data collection

In total, 30 health professionals who attended a training course to familiarize themselves with the survey and measurement techniques before the survey participated. The health professionals were trained in BP measurement and anthropometric measurement protocols.

Each team was given the same model of an automatic oscillometric BP-measuring device (Omron M3 Intellisense, HEM-7051-E, Tokyo, Japan), two cuffs (22–32 cm and 33–42 cm), a weighing scale and a measuring tape. In addition, each team had an iPad 2, which was used to transfer the collected data electronically via the Internet to the main computer at the study center. In the study center, two supervisors were responsible for the quality control of the collected data.

Measurements and definitions

Systolic and diastolic blood pressure (SBP, DBP), heart rate, weight, height, and waist and arm circumferences were measured according to standard protocols. Body mass index (BMI) was also calculated. Obesity and overweight were defined as BMIs of at least 30 and 25–29.9 kg/m², respectively.

BP measurements were completed according to the recommendations at the time the study was performed [28]. None of the participants had alcohol or tea/coffee intake nor had smoked at least 30 min prior to the measurement. Initially, each participant's BP was measured using appropriately sized cuffs after 5 min of rest in a sitting position with his/her back supported. If the reading was higher in one arm, that arm was used for the following measurements. At least three consecutive BP measurements were obtained, with a time interval of at least 2 min between each measurement according to the recommendations of the European Society of Hypertension (ESH). If the difference between the last two measurements was <5 mmHg, the arithmetic mean of the second and third BP measurements was noted as the visit BP. A fourth or multiple additional measurements were obtained when there was a difference of ≥ 5 mmHg between the last two measurements. When the difference between the last two measurements was <5 mmHg, the arithmetic mean of the last two BP measurements was recorded as the visit BP.

Hypertension was defined as an average SBP of ≥ 140 mmHg and an average DBP of ≥ 90 mmHg, or previously diagnosed HT, and/or the use of antihypertensive medication regardless of BP readings. Awareness of HT was described as any previous diagnosis of HT by a health professional among the participants identified as having HT. Treatment of HT was defined as the use of antihypertensive medication at the time of the interview. Control of HT was defined as an SBP < 140 mmHg and a DBP < 90 mmHg [23]. The control rates among patients receiving antihypertensive medication were also recorded.

Arm circumference

Mid-arm circumference (the midpoint of the acromion and olecranon) was measured with a plastic tape on bare extremities. Arm circumference was measured from both arms, but only the measurements taken from the right arm were reported. The right mid-arm circumference for the PatenT study, conducted in 2003 [20], was calculated from the following formula derived from the linear-regression analysis in the PatenT2 study to identify independent predictors of right mid-arm circumference:

$$\text{Right-arm circumference} = 21.303 + ((-0.001)*\text{age}) + ((-1.245)*\text{gender}) + (0.379*\text{BMI}) + (0.137*\text{residence})$$

Cuff size and device validation

Cuff size was checked only for upper-arm devices including aneroid and mercury devices. Information regarding the cuff size of the upper-arm devices was taken from the device or box of the device, if it was available. Unless the owner gave specific information, the device was considered standard (22–32 cm). An automated upper-arm device listed and recommended on either or both of the dabl Educational Trust and the British Hypertension Society websites [29, 30] was considered “validated”.

Telephone survey in 2017

In 2017, 5 years after the field survey, 1318 hypertensive subjects who did not own a sphygmomanometer were called by telephone to ask whether they now owned a sphygmomanometer. If they had one, the type of the sphygmomanometer (nonautomated, automated upper arm or automated wrist) was recorded.

Statistical analysis

Predictive analytics software (PASW statistics 18, 2009) was used for the analysis. A type-I error level of less than 5% was used to infer statistical significance. The variables were investigated using visual (histograms, probability plots) and analytical (Kolmogorov–Smirnov/Shapiro–Wilk tests) methods to determine whether they were normally distributed. Univariate analyses were used to identify the variables associated with the possession of a blood-pressure measuring device of hypertensive patients who were aware of their HT; for this, the chi-square and Mantel–Haenszel tests were used as appropriate. The Mann–Whitney *U* test was used for the comparison of SBP and diastolic BPs both in the HT groups and the aware group in terms of the device type and device validation. For the multivariate analysis, the possible factors identified with univariate analyses were further entered into a logistic regression analysis with backward selection to determine the independent predictors of possessing a blood-pressure measuring device for hypertensive patients who were aware of their HT. Hosmer–Lemeshow and Pearson's goodness-of-fit statistics were used to evaluate the model fit. To investigate the factors affecting the baseline SBP and DBPs in the aware group, multivariate linear-regression analysis was performed.

Results

The basic demographic information of the participants was published previously [23]. A total of 1650 participants

(30.3%) of the PatenT2 study were hypertensive, and 54.7% of the hypertensive patients ($n = 902$) were aware of their diagnosis. The use of pharmacological treatment and control rates were 47.4% and 28.7%, respectively. Among the 5437 participants including normotensive and unaware hypertensive patients, 660 (12%) had a home sphygmomanometer (95% CI: 11.3–13.0%), and the percentage of patients owning a device was 8.7% (328/3787). Of 1650 hypertensive subjects, 332 (20.1%) had a device (95% CI: 18.2–22.1), but the percentage of patients owning a sphygmomanometer was 28.8% among patients who were aware of their disease (260/902).

Factors associated with the ownership of a sphygmomanometer among hypertensive patients who were aware of their HT

Various factors related to the ownership of a sphygmomanometer among the patients who were aware of their HT are presented in Tables 1 and 2. Sphygmomanometer ownership was significantly higher among participants living in urban areas, participants with higher education status, participants with higher income level and patients using antihypertensive medication. Multivariate logistic regression analysis demonstrated that the factors associated with owning a sphygmomanometer were female gender, older age, higher educational status, living in urban areas and antihypertensive drug use (Table 2).

Device information

Of 660 devices, 482 (73%) were automated, wrist devices being more common. Table 3 shows the types of sphygmomanometers owned among all participants and hypertensive subjects who were aware of their HT.

HBPM practice among participants

Of the device owners, 429 (65%) stated that they had not had any training regarding the operation of the device. The participants learned the usage of their devices mainly from their relatives and the sellers; the percentage of patients who learned to use the device from healthcare professionals, physicians or nurses was 29%, including relatives working in health care.

Among patients who were aware of their HT, 50% stated that they had not had any training regarding the operation of the device, and the percentage of patients who learned to use the device from healthcare professionals was 32%. Among the patients who were aware of their HT, 204 had automated devices, 74 (36%) of which were validated. Twenty-two (18%) of the 120 wrist devices and 52 (62%) of the 84 automated upper-arm devices were validated.

Table 1 Factors associated with possession of a blood-pressure measuring device in the hypertensive population who were aware of their hypertension

Variable	Patients possessing a blood-pressure measuring device (%)	Statistical test used for analysis	<i>p</i>
Gender		Pearson chi-square	
Female	175 (29.7)		0.446
Male	85 (27.2)		
Age groups		Mantel–Haenszel test (linear-by-linear association)	
18–29 years	0 (0.0)		0.267
30–39 years	6 (17.1)		
40–49 years	35 (26.5)		
50–59 years	82 (32.0)		
60–69 years	78 (29.0)		
≥70 years	59 (28.9)		
Residence		Pearson chi-square	
Urban	227 (33.1)		<0.001
Rural	33 (15.2)		
Educational status		Mantel–Haenszel test (linear-by-linear association)	
Illiterate	20 (20.6)		<0.001
Literate	27 (21.8)		
Primary school graduate	127 (27.9)		
Middle school graduate	24 (32.9)		
High school graduate	33 (36.3)		
University graduate	29 (47.5)		
Monthly income level		Mantel–Haenszel test (linear-by-linear association)	
<1001 TL ^a	142 (24.8)		<0.001
≥1000 TL	102 (37.0)		
Blood-pressure categories		Mantel–Haenszel test (linear-by-linear association)	
Optimal	60 (37.3)		0.001
Normal	52 (34.4)		
High normal	36 (22.2)		
Stage 1 hypertension	78 (28.4)		
Stage 2 hypertension	27 (24.8)		
Stage 3 hypertension	7 (15.9)		
Body mass index		Mantel–Haenszel test (linear-by-linear association)	
Underweight (<18.5 kg/m ²) +normal weight (18.5–24.9 kg/m ²)	22 (22.2)		0.045
Overweight (25–29.9 kg/m ²)	75 (26.9)		
Obese (≥30 kg/m ²)	163 (31.2)		
Antihypertensive drug usage		Pearson chi-square	
Yes	239 (30.7)		0.002
No	21 (17.1)		

^a 1 Euro=2.4 TL March 2012

Table 2 Results of multivariate logistic regression analyses for ownership of a blood-pressure measuring device

Variables ^a	Multivariate (<i>p</i>)	Odds ratio	95% CI
Gender (female)	0.054	1.390	0.995–1.943
Age			
18–39 years		1.0 (reference)	1.0 (reference)
40–49 years	0.138	2.088	0.788–5.528
50–59 years	0.025	2.903	1.144–7.369
60–69 years	0.031	2.805	1.100–7.149
70 years and above	0.021	3.109	1.190–8.124
Living in urban areas	<0.001	2.609	1.729–3.936
Educational status			
Illiterate		1.0 (reference)	1.0 (reference)
Literate	0.841	1.071	0.548–2.091
Primary school graduate	0.060	1.740	0.976–3.101
Middle school graduate	0.049	2.153	1.003–4.620
High school graduate	0.006	2.733	1.332–5.607
University graduate	<0.001	3.991	1.862–8.555
Antihypertensive drug usage	0.012	1.922	1.153–3.205

CI confidence interval.

^a The variables included in the multivariate logistic regression analysis were gender, age, educational status, living in urban areas, income level, blood-pressure categories, awareness of hypertension, antihypertensive drug usage, and body mass index. Only variables having significant differences are shown.

Table 3 Types of sphygmomanometers among all participants (including aware hypertensive subjects) and hypertensive subjects who were aware of their hypertension in 2012

Type of device	All participants (<i>n</i>)	Aware hypertensive subjects (<i>n</i>)
Automated wrist	294	120
Automated upper arm	188	84
Aneroid	132	50
Mercury	4	1
Not at home	42	5
Total	660	260

Arm circumference and cuff size

The mean right-arm circumference was 30.0 cm among all participants and was 29.6 cm for the participants of the PATENT study. The mean right-arm circumference was

wider in hypertensive patients than that in hypertensive patients in the PATENT study (31 cm vs. 30.6 cm). The frequency of requiring a large cuff was 32% among aware hypertensive patients having an upper-arm device, but only one had a cuff ≥ 33 cm.

The association between possessing a home sphygmomanometer and BP levels in hypertensive patients who were aware of their disease

The mean SBP and DBPs among aware patients who had a BP monitor were 5.8 mmHg and 3.4 mmHg lower than those of patients not having a device. Validated and/or upper-arm devices were associated with lower SBP and DBPs (Table 4).

The results of linear-regression analysis

While the baseline SBP was 143.9 mmHg in the HT awareness group, linear-regression analysis indicated that being male resulted in a 3.125-mmHg increase, and every 1-year increase in age resulted in a 0.291-mmHg increase (an ~ 3 -mmHg increase with every 10-year increase in age); furthermore, living in rural areas resulted in a 2.369-mmHg increase in the baseline SBP ($p = 0.003$, $p < 0.001$ and $p = 0.027$, respectively). On the other hand, each categorical increase in income level resulted in a 2.401-mmHg decrease, antihypertensive drug use resulted in a 14.386-mmHg decrease and the presence of a device at home resulted in a 3.744-mmHg decrease in the baseline SBP

Table 4 Mean systolic blood pressure (SBP) and diastolic blood pressure (DBP) according to the device type and validation status (only automatic devices are shown)

	SBP mmHg	DBP mmHg
Device type		
Upper arm	133	73
Wrist	141	76
	$P=0.010$	$P=0.069$
Validation status		
Validated (upper arm+wrist)	133	72
Nonvalidated (upper arm+wrist)	140	77
	$P=0.011$	$P=0.014$
Validated (upper arm)	132	72
Nonvalidated (upper arm)	135	76
	$P=0.293$	$P=0.093$
Validated (wrist)	135	75
Nonvalidated (wrist)	142	77
	$P=0.161$	$P=0.484$

Table 5 Linear-regression analysis of factors affecting systolic blood pressure

Variables	Multivariate (<i>p</i>)	β
Constant coefficient	<0.001	143.923
Gender (male)	0.003	3.125
Age	<0.001	0.291
Living in rural areas	0.027	2.369
Educational status	0.214	-0.578
Income level	0.002	-2.401
Euro-stat region	0.550	0.085
BMI	0.342	-0.087
Antihypertensive drug usage	<0.001	-14.386
Ownership of a blood-pressure measuring device	0.002	-3.744

($p = 0.002$, $p < 0.001$ and $p = 0.002$, respectively) (Table 5).

While the baseline DBP was 91.5 mmHg in the HT awareness group, linear-regression analysis indicated that being male resulted in a 1.384-mmHg increase and living in rural areas resulted in a 2.335-mmHg increase in the baseline DBP ($p = 0.031$ and $p < 0.001$, respectively). On the other hand, every 1-year increase in age resulted in a 0.145-mmHg decrease (~1.5-mmHg decrease with every 10-year increase in age), antihypertensive drug use resulted in a 6.206-mmHg decrease and the presence of a device at home resulted in a 2.837-mmHg decrease in the baseline DBP ($p < 0.001$ for each) (Table 6).

Linear-regression analysis of factors affecting SBP and DBP after stratification by antihypertensive drug use is shown in Table 7 (the method and variables are the same as those in Tables 5 and 6; only the ownership of a blood-pressure measuring device variable is shown). The association between possessing a BP device and BP decrease remained after stratification by antihypertensive drug use.

Table 6 Linear-regression analysis of factors affecting diastolic blood pressure.

Variables	Multivariate (<i>p</i>)	β
Constant coefficient	<0.001	91.537
Gender (male)	0.031	1.384
Age	<0.001	-0.145
Living in rural areas	<0.001	2.335
Educational status	0.674	-0.121
Income level	0.083	-0.849
Euro-stat region	0.474	0.063
BMI	0.497	0.038
Antihypertensive drug usage	<0.001	-6.206
Ownership of a blood-pressure measuring device	<0.001	-2.837

Table 7 Linear-regression analysis of factors affecting systolic blood pressure (SBP) and diastolic blood pressure (DBP) after stratification by the antihypertensive drug use (method and variables same as Tables 5 and 6, only ownership of a blood-pressure measuring device variable is shown)

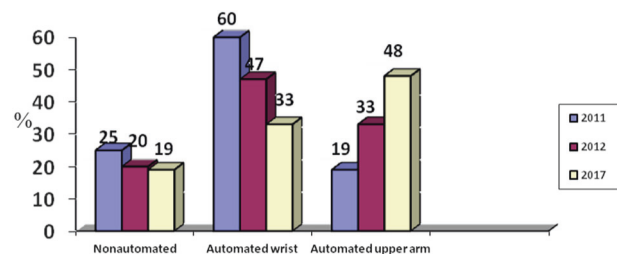
	Multivariate (<i>p</i>)	β
Antihypertensive drug (+)		
SBP	0.019	-4.107
DBP	0.004	-2.880
Antihypertensive drug (-)		
SBP	0.160	-2.566
DBP	0.030	-2.758

Telephone survey in 2017

In total, 1318 hypertensive Patent2 participants did not have a BP monitor in 2012, and 437 were reached by telephone in 2017. After 2012, 254 (58.1%) of 437 patients bought a new device. The types of devices were automated upper arm ($n = 122$, 48%), automated wrist ($n = 82$, 33%) and nonautomated ($n = 48$, 19%). The device type of two owners could not be determined. Figure 1 shows the change in sphygmomanometer type with time.

Discussion

The present study confirmed the findings of the previous study. Only the new findings will be discussed. Among 902 hypertensive patients who were aware of their disease, 260 (28.8%) had a home sphygmomanometer, a lower rate than that found in our first survey in 2011 (28.8 vs. 46.6%). As mentioned in our paper [23], the first survey using computer-assisted telephone interviewing had some limitations. The findings of the 2017 telephone survey supported a possible Hawthorne effect and showed that the frequency of ownership of a sphygmomanometer had increased. Forty two (6%) of the 660 devices were not in the houses of the participants during our visit in this study, which indicated

**Fig. 1** Comparison of sphygmomanometer types in three surveys by time (the results are expressed as percentages)

that BP monitors can be shared among neighbors and relatives.

The most striking finding of our study was the demonstration of an association between sphygmomanometer ownership and BP decrease [31, 32] in a nationwide field survey. The SBP and DBPs among aware hypertensive patients possessing a home BP monitor were 5.8 and 3.4 mmHg lower, respectively, than those of patients that did not have one. The linear-regression analysis showed that possessing a BP device is associated with a decrease of 3.7 mmHg and 2.8 mmHg on SBP and DBPs, respectively. To our knowledge, similar nationwide data are not available. The findings of this study support previous data regarding the usefulness of HBPM [33, 34]. Moreover, it should be kept in mind that nonadherence to the recommended guidelines for BP measurement is a common problem [35], and the consecutive measurement of BP (two times) may be more useful [36].

The cause of lower BP in patients possessing a sphygmomanometer is not known, and the current study was not designed to find the answer to this question. It can be argued that compliant patients willing to learn, measure and control their BP bought a sphygmomanometer, as reported by Ayala et al. [37]; regardless, the association between possessing a BP monitor and lower BP supports the notion that HBPM is a supplementary tool in the control of BP. Ostchega et al. [38] reported that hypertensive patients who received providers' recommendations to perform HBPM were more likely to do so than those who did not receive recommendations. Home BP monitoring and home BP monitor reimbursement have been recommended by many guidelines [10–14]. Our data support the previous evidence suggesting that HBPM may be cost-effective [2, 18, 34]; therefore, home BP monitor reimbursement should be encouraged. Lifestyle changes are the effective measures in the management of HT. Home BP monitoring is a lifestyle change that can increase treatment adherence [14]. Dietary salt reduction is an effective lifestyle change in the management of HT and can save billions of healthcare-related payments [39]. Similarly, owning a home BP monitor may play a role in improving BP control and could save HT-related treatment expenses, including the cost of medication. In addition, HBPM is a good supplementary tool for the improvement of BP control in low-income countries [40].

There were problems in the patients' choice of a BP monitor, such as device validation and the use of an appropriate cuff bladder for nonstandard arm circumference sizes in addition to the dominance of wrist devices. It is encouraging to see the increasing number of upper arm automated sphygmomanometers (Fig. 1). Since obesity and arm circumference are increasing, problems related to cuff

size may also increase. Many nonvalidated devices are being introduced into the market [41–43], and only 36% of the devices in this study were validated. Blood pressure was lower among hypertensive patients possessing a validated and/or upper-arm BP monitor (Table 4). Since the use of an appropriate cuff size is essential for the accurate measurement of BP [3, 10, 11], overweight and obese patients often require sphygmomanometers with large or extra-large cuffs. Although most regular cuff sizes are appropriate for patients having an arm circumference of 22–32 cm, more than 30% of the patients having an upper-arm device required large cuffs, but only one (less than 1%) patient had a large cuff, indicating ignorance of the problem.

This was the first study investigating the arm circumference of a Turkish population, and the results demonstrated that arm circumference and the frequency of patients requiring a large cuff were increased compared to 2003 [24]. Since obesity continues to increase and arm circumference is closely associated with obesity [44], it is expected that the number of patients requiring a large cuff will increase.

In conclusion, despite market prices being reasonable (cheaper than 30–50 Euros; most patients can afford this in Turkey), many patients do not have a sphygmomanometer. The use of wrist devices and nonvalidated devices is common, and the selection of an appropriately sized cuff is ignored in obese patients. The demonstration of an association between sphygmomanometer ownership and BP decrease was a striking finding. As demonstrated in several studies, the implementation of a HT care program consisting of sphygmomanometer reimbursement, training patients on how to use the device and HBPM can be cost-effective.

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Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

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