

Detection of peripheral arterial disease with an improved automated device: comparison of a new oscillometric device and the standard Doppler method

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Abstract: In occidental countries, peripheral arterial disease (PAD) is an important health issue; however, most subjects are asymptomatic (~50%) and therefore undiagnosed and untreated. Current guidelines recommend screening for PAD in primary care setting using ankle brachial index (ABI) in all patients with cardiovascular risks. This is, however, not performed strictly because the standard Doppler method is cumbersome and time-consuming. Here, we evaluate the accuracy and reproducibility of ABI measurements obtained by an improved automated oscillometric device, the MESI ABPI MD[®] device, and the standard Doppler method. ABI was measured in random order in a general practice with Doppler probes by two operators separately (ABI_{dop}) and twice with the MESI ABPI MD device (ABI_{mesi}). ABI_{dop} was calculated dividing the highest systolic blood pressure from both tibial and dorsalis pedis arteries by the highest systolic blood pressure of both brachial arteries. ABI_{mesi} was obtained automatically with simultaneous measurements on three extremities. According to ABI_{dop}, PAD was present in 10% of the 136 screened subjects (68.2±7.4 years). Interoperator coefficient of variation was 5.5% for ABI_{dop}, while the intrasubject coefficient of variation for ABI_{mesi} was 3.0%. ABI_{mesi} was correlated with ABI_{dop} ($R=0.61$, $P<0.0001$). The difference between the two techniques was 0.06 ± 0.14 with ABI_{mesi} providing slightly higher values ($P<0.0001$) and negligible bias across the range ($R=0.19$, $P<0.0001$). Therefore, ABI_{mesi} ≤1 had a sensitivity of 85% and specificity of 96% to detect ABI_{dop} ≤0.9 and hence PAD. Doppler measurements took seven times longer than MESI ABPI MD measurements to be performed. In conclusion, MESI improved automated oscillometric method and offered a faster and repeatable measurement of ABI with only a small, clinically irrelevant overestimation of ABI value. The tested MESI ABPI MD-improved oscillometric system can be used as a screening tool for patients in general practice and would enable family doctors to comply with current guidelines for PAD.

Keywords: ankle brachial index, peripheral arterial disease, automated oscillometric method, Doppler method, screening tool

Background

The incidence of the peripheral arterial disease (PAD) in lower extremities is increasing with age in the population. In Western European countries, more than 20% of people older than 65 years have PAD in lower extremities either in symptomatic or asymptomatic phase.¹ More than 50% of patients with present PAD are asymptomatic² and are therefore untreated.³ Progress of the PAD in the lower extremities can lead to onset of symptoms.

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First, leg pain during activity (intermittent claudication) occurs. If not treated, symptoms worsen and progress to critical limb ischemia, ulcer, or gangrene. Even more importantly, patients with PAD have four- to six-fold higher risk of cardiovascular or cerebrovascular events in comparison to the healthy population.⁴⁻⁶ Progression and severity of PAD can be altered by risk factor management and preventive medical treatment.⁷

The measurement of ankle brachial index (ABI) is a reliable noninvasive diagnostic tool with high sensitivity and specificity to diagnose symptomatic and asymptomatic PAD.^{5,8} Current guidelines for management of PAD advise to measure ABI in all subjects >65 years, >50 years if they have hypertension, diabetes, and/or are currently smokers, and in subjects with clinical symptoms of PAD.^{3,7}

ABI obtained by Doppler method is usually regarded as the gold standard,⁹ where the operator detects the return of blood flow with Doppler probe during the deflation of a proximal cuff. But there are some drawbacks of this Doppler method: it requires an experienced operator, is time-consuming,¹⁰⁻¹³ and does not allow measurement of transient state such as after a strandness test during which it is important to measure ABI just after walking.

MESI d.o.o. (Ljubljana, Slovenia) developed the MESI ABPI MD[®] system based on an improved oscillometric technology to provide an easy solution for PAD screening in clinical settings. The aim of this study was to evaluate the reproducibility and the accuracy of the MESI ABPI MD device from MESI compared with standard Doppler method to measure ABI in general practice population.

Methods

Subjects were approached at the general practitioner's office during their regular visits. They were invited to participate in the study if they were 1) >65 years or 2) >50 years with smoking habit or diabetes or 3) had pain in the lower extremities during exercise. Subjects who fulfilled at least one of these criteria were included in the study, while subjects with known arrhythmia, upper extremity arteriovenous fistulas or symptomatic critical limb ischemia, ulcers or major lower leg amputations were excluded. All subjects provided written informed consent and the study protocol was approved by the Ljubljana University ethics committee and was in accordance with the Declaration of Helsinki. Permission was also obtained from the subject for the use of Figure 1.

Subject's medical history was recorded by the general practitioner. After a few minutes of rest in supine position, four ABI measurements were obtained for each subject: one with the Doppler probe method and operator 1, one with the

Doppler probe and operator 2, and two with the automatic oscillometric device (MESI ABPI MD). The order of the ABI method was chosen randomly, and the ABI results were hidden and not visible to the subjects or operators. There were two skilled operators who performed Doppler probe measurements on each subject separately and a third operator who performed both oscillometric measurements.

Measurement of ABI by means of Doppler probe method (ABI_dop) was performed according to the standard protocol⁹ with a calibrated sphygmomanometer and 8 MHz Doppler probes (Dopplex SD2, Huntleigh Healthcare Ltd, Cardiff, UK). ABI_dop was calculated as the ratio of the highest systolic blood pressure (SBP) obtained from both tibial and dorsalis pedis arteries at one ankle to the highest SBP of both brachial arteries. ABI_dop_L refers to ABI values obtained on the left leg, ABI_dop_R on the right leg. Measurement of ABI (ABI_mesi) with the improved automated oscillometric device (MESI ABPI MD; Figure 1) was performed according to the manufacturer's instructions. In short, MESI ABPI MD system is based on simultaneous plethysmographic measurements through three cuffs, one conical cuff placed on the right or left arm, and two conical cuffs placed on each of the ankles. Precisely regulated system inflated and then deflated the three cuffs simultaneously at the same pressure level. Heart beats create oscillations on the plethysmographic signals, which are used to calculate blood pressure values from the three limbs using specific proprietary algorithms for the ankle blood pressure determination. The ankle cuffs measure oscillometric signals over the whole ankle and hence the highest SBP of all arteries within the ankle is detected.



Figure 1 Picture of the MESI ABPI MD[®] (MESI d.o.o., Ljubljana, Slovenia) device.

Values of ABI_{dop} <0.90 were regarded as being a significant sign of PAD, whereas values >1.40 indicated incompressible ankle arteries and values in between were regarded as normal.

Time taken by operator 1, operator 2, and MESI ABPI MD to assess ABI was also measured.

Statistical analysis

Numerical data are presented as a mean \pm standard deviation, while categorical data are expressed as percentage. Differences between measurements were tested with bilateral Student's *t*-test. Coefficient of variation (CV) was defined as the mean standard deviation of measurements in a same subject divided by average of these measurements in a same patient. Relationship between variables was measured with the Pearson's correlation coefficient and Bland–Altman plots were used to quantify the differences between measurements. Limits of agreement were determined by 95% confidence interval. Assuming that ABI values in a typical primary care should be between 0.6 and 1.6, we calculated that we needed at least 132 subjects to detect a difference of 0.04 with 95% confidence level and 90% power. All statistical analysis was performed using SPSS version 18 (SPSS Inc., Chicago, IL, USA).

Results

Out of 150 screened subjects, six had atrial fibrillation and were hence excluded from the final analysis. In 14 (9.3%) subjects, it was not possible to assess ABI with Doppler probe and/or oscillometric device due to critical limb ischemia or incompressible arteries. Patients' characteristics of the 136 remaining subjects are presented in Table 1.

Results from Doppler method

Both operators were experienced in measuring SBPs using Doppler probe. However, they had a strong tendency to round numbers as 90% of the pressure values reported ended by 0 or 5. Interoperator differences were small with highest interoperator difference for the right posterior tibial artery being 3.5 ± 15.9 mmHg. There was no trend in the interoperator differences across patients confirming that both operators were experienced in measuring ABI_{dop} and that neither operator experienced a learning curve. ABI_{dop} values varied from 0.5 to 1.7 with a mean ABI_{dop} of 1.1 ± 0.2 for both operators. The difference between operator was not significant (0.01 ± 0.1 , $P=0.107$) and the interoperator CV for ABI_{dop} was 5.9% for each leg.

Table 1 Patients' characteristics

	N=136
Age (years), mean \pm SD	64.0 \pm 7.8
Office SBP (mmHg), mean \pm SD	128 \pm 16
Office DBP (mmHg), mean \pm SD	75 \pm 11
Office HR (bpm), mean \pm SD	67 \pm 11
Hypertension, n (%)	66 (49)
Dyslipidemia, n (%)	58 (43)
Diabetes mellitus, n (%)	19 (14)
Smoking, n (%)	
Smoker	22 (16)
Former smoker	39 (29)
Nonsmoker	75 (55)
Physical activity, n (%)	
Active	80 (59)
Moderately active	44 (32)
Inactive	12 (9)

Notes: Hypertension was defined as treated with antihypertensive drugs or office blood pressure >140/90 mmHg. Dyslipidemia was defined as treated with lipid-lowering drugs. Diabetes was defined as undertreatment for diabetes. Physically active was defined as when a subject exercised at least 30 minutes a day more than three times a week, moderately active was when a subject exercised at least 30 minutes once a week, and inactive was no activity over a week.

Abbreviations: DBP, diastolic blood pressure; HR, heart rate; SBP, systolic blood pressure; SD, standard deviation.

Results from MESI device

Inpatient differences between repeated measurements with the MESI ABPI MD device were small: 1.8 ± 6.6 , 3.0 ± 8.8 , and 2.8 ± 8.0 mmHg for right arm, left ankle, and right ankle, respectively. ABI_{mesi} varied from 0.57 to 1.46 with mean values of 1.19 ± 0.15 and 1.20 ± 0.14 on left and right side, respectively. CV of repeated measurements was 3.5% on left side and 3.2% on the right side, showing excellent repeatability.

We compared the average of the two measurements of ABI_{mesi_L} with the average of ABI_{dop_L} from operators 1 and 2 as well as ABI_{mesi_R} with ABI_{dop_R}. As results were similar, they were pooled together, each subject having two data points (Figure 2A and B).

ABI_{mesi} was strongly correlated to ABI_{dop} ($R=0.61$, $P<0.0001$, Figure 2A). The error between the two methods exhibited a normal distribution and the Bland–Altman plot showed a mean difference of 0.06 ± 0.14 (limits of agreement: -0.211 to 0.332 , Figure 2B) representing a small overestimation (5.1%, $P<0.0001$) of ABI_{mesi} compared with ABI_{dop}. A small but significant bias across ABI range was observed with lower ABI value being slightly more overestimated than higher ABI value ($R=0.19$, $P<0.0001$).

The difference between Doppler and MESI ABPI MD device did not vary across recruitment (data not shown) confirming that MESI system does not drift with time.

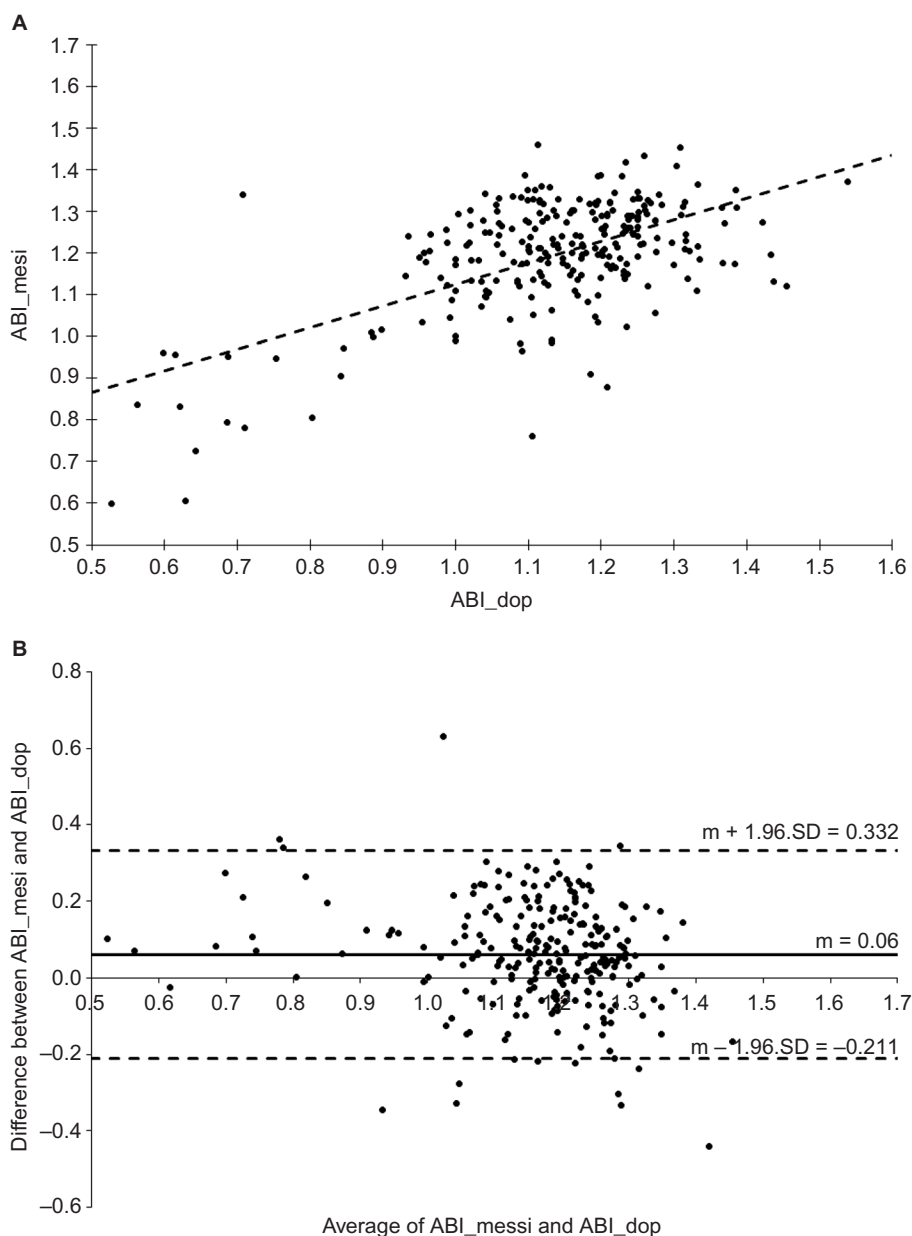


Figure 2 Correlation and Bland–Altman plot of the difference between MESI ABPI MD[®] device and Doppler method

Notes: (A) Correlation of ABI_mesi versus ABI_dop and (B) Bland–Altman plot of the difference between ABI_mesi and ABI_dop.

Abbreviations: ABI, ankle brachial index; ABI_dop, ABI by means of Doppler probe method; ABI_mesi, MESI ABPI MD device; SD, standard deviation.

To further evaluate the accuracy of the MESI ABPI MD device to screen for PAD, we looked at the number of patients with an ABI_{dop} ≤ 0.9 on at least one leg and found that 10% of the subjects had PAD on at least one leg according to the Doppler method. Using a threshold of 0.9 on ABI_{mes}, MESI ABPI MD device diagnosed 6.8% of the subjects with a sensitivity of 57% and a specificity of 99%. However, several publications have advised to use ABI ≤ 1 for oscillometric techniques.^{14–17} In that case, MESI device reached 85% of sensitivity and 96% specificity to detect PAD in at least one leg.

Only 3.8% of the screened patients had an ABI_{dop} > 1.4 (3% for ABI_{mes}). This number is too small to calculate the specificity and sensitivity to detect arterial calcification.

Time spent to assess ABI

Time spent to measure ABI is presented in Figure 3. Confirming the experience of the operators, Doppler method took 14 ± 1.54 minutes for operator 1 and 14 ± 1.90 minutes for operator 2 ($P=0.58$) to measure arm and both left and right dorsalis pedis and tibial systolic pressures. Measurements with MESI ABPI MD took exactly 2 minutes whoever the

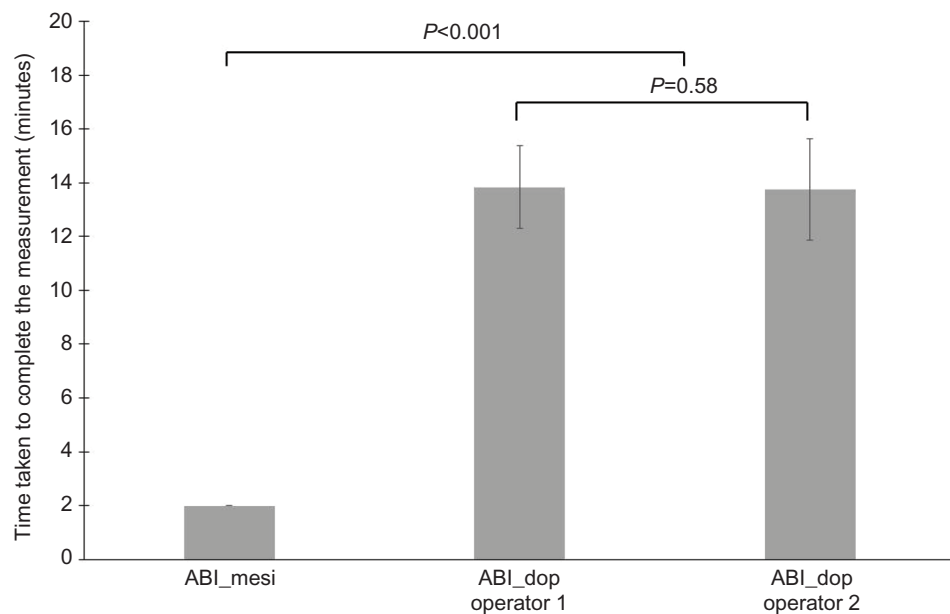


Figure 3 Time spent in minutes to measure ABI.

Note: Error bars show standard deviation.

Abbreviations: ABI, ankle brachial index; ABI_dop, ABI by means of Doppler probe method; ABI_mesi, MESI ABPI MD device.

patient and were hence complete seven times faster than the standard Doppler ABI measurement ($P < 0.001$).

Discussion

This is the first study comparing the improved automated oscillometric MESI ABPI MD device from MESI d.o.o with the standard Doppler method. This device offers the opportunity to assess ABI easily from a proprietary method based on the analysis of oscillations from plethysmographic signals recorded through conical cuffs. Hence, there is no bias due to operator error. Even with experienced operators such as the ones we used in our study, there is a strong tendency to round up pressure values, which could lead to inaccuracy in ABI assessment. This is one of the various sources of errors which makes Doppler method having higher variability than oscillometric measurements.¹⁸ Our data confirm this as MESI variability was small (highest CV = 3.5% on left ankle) compared with Doppler variability. Due to time constraints, we did not measure the intraoperator variability for Doppler measurements, but others¹⁹ have reported 8% for intraoperator and 9% for interoperator variability regardless of operator's medical background.

Unlike for validation of brachial blood pressure electronic devices, there is no recognized protocol to test the accuracy of ABI devices. However, most publications reports: 1) the coefficient of correlation between the two techniques, 2) the difference between the two techniques, and 3) in a smaller proportion of studies, the sensitivity and specificity values of the oscillometric method in diagnosing PAD. In 2012,

Verberk et al found 25 studies comparing oscillometric ABI determination with Doppler determination.¹⁸ Sixteen of them reported the coefficient of correlation which was on average 0.71 ± 0.05 , which is relatively similar to our coefficient of correlation of 0.61 ($P < 0.001$). Eighteen studies reported the difference in ABI values assessed by the two methods with an average error of 0.02 indicating that in general, oscillometric determination of ABI gave slightly higher values than the Doppler method. Our results are also in agreement with this result as we observed a small overestimation of 0.06 ± 0.14 . This small overestimation of ABI with the MESI ABPI MD device is due to higher ankle systolic pressure values recorded on the ankles (5.8 ± 14.8 mmHg on average compared with Doppler tibial or dorsalis pedis artery pressure, $P < 0.001$). This could be explained by observer error: the time for the operator between hearing the Doppler signal and recording the pressure value from the sphygmomanometer might induce a lower systolic pressure value,¹⁸ however, we did not observe any differences between brachial SBP recorded with the MESI ABPI MD device and the Doppler method on the same arm.

The average sensitivity and specificity found by the meta-analysis by Verbeck et al were $69\% \pm 6\%$ and $96\% \pm 0.8\%$, respectively.¹⁸ However, several authors have reported that using a cutoff value closer to 1 was better to diagnose PAD surely because of the systematic small overestimation of oscillometric method.¹⁴⁻¹⁷ When using 1 as a cutoff value, the MESI ABPI MD device sensitivity was 85% and 96% for specificity showing excellent detection of PAD. In our study, PAD diagnosis was

solely based on Doppler ABI results. To truly evaluate sensitivity and specificity of MESI ABPI MD system, ABI results should be compared with PAD diagnosis from imagery techniques.

Both operators of our study took approximately 14 minutes to complete full measurement of ABI. Indeed, they had to successively inflate the cuff at least once on the arm and twice per leg for the tibial and dorsalis pedis arteries. In comparison, MESI ABPI MD took 2 minutes to perform measurement on the three cuffs simultaneously. This important discrepancy is intuitive and has been reported previously.¹⁸ What is more important is that our measurement did not take into account the required resting time. Indeed, with successive Doppler measurements, it is important to make sure all hemodynamic variables are stable before assessing ABI. While with MESI ABPI MD, systemic variations of blood pressure do not matter as both legs and arms are measured simultaneously.

There are some controversies regarding the use of oscillometric devices for ABI measurements; however, several dedicated ABI oscillometric devices have shown comparable results to the Doppler method.¹⁸ Some complain that if ankle pressure is particularly low, oscillometric might not be able to calculate ABI. This is, however, by itself, an important clinical diagnosis. It is also crucial to use specific ankle oscillometry technique and not a standard blood pressure designed for brachial pressure. Indeed, as lower leg geometry and anatomy are different from the upper arm, specific cuffs and specific pressure detection algorithms are necessary at the ankle level. Simultaneous measurements are also a plus in order to reduce measurements variability and “white coat” effect.¹⁸ MESI ABPI MD device has been designed in that respect and offers an accurate, easy, and rapid solution to avoid drawbacks of the Doppler method. In addition, MESI ABPI MD offers a comprehensive detection of critical limb ischemia (blood pressure <70 mmHg) by displaying a specific message (message E1) particularly relevant for patients with severe PAD.

Limitations

The main limitation of our study resided in the studied ABI range. Subjects recruitment was performed in general practice settings and we only diagnosed 10% of subjects with PAD. The distribution of ABI values is hence centered around normal ABI values with fewer data available <0.9 and >1.4. While further studies are required to verify MESI’s accuracy <0.9, our cohort represents typical primary care population and hence shows that the MESI device is suitable for such settings.

Conclusion

MESI ABPI MD device provides fast, accurate, and precise measurements of ABI in general practice. As ABI values are

slightly overestimated and sensitivity compared with Doppler value is slightly lower, patients with borderline oscillometric results between 0.9 and 1 should be considered at very high risk of PAD. MESI ABPI MD can be accepted as an alternative to the Doppler method and be used as a large-scale screening tool for family doctors in order to comply with current guidelines for the diagnosis of PAD.

Acknowledgments

Part of this paper will be presented as a poster presentation with interim findings at the European Society of Hypertension congress on June 11, 2016. Organizers of the congress intend to publish the poster abstract in the August supplement issue of *Journal of Hypertension*.

Disclosure

SCM received financial compensation for her work on the analysis, the interpretation of the data, as well as editing this manuscript. MŠ has some shares in MESI d.o.o. The other authors report no conflicts of interest in this work.

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