Inter-limb blood pressure difference, ankle-brachial index and its association with border zone cerebral infarcts: A prospective single-blind study from central India

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Abstract

Background: Border zone infarcts (BZI) are ischemic lesions occurring between two non-anastomosing arterial territories. BZI patients have higher incidence of recurrent strokes (RS). The predictors of peripheral arterial disease (PAD), low ankle-brachial index (ABI) and high inter-arm blood pressure difference (IABD) are associated with intracranial atherosclerosis and thus, could also have an association with BZI. Our single blind prospective study explores the predictive value of these clinical variables, ABI and IABD for BZI in patients with acute ischemic stroke.

Methods: The subjects fulfilling the inclusion criterion (183 of 335, 78.14% men) were divided into BZI (n=89) and pure core arterial territory infarct (CATI, n=94) groups, based upon MRI characteristics of infarcts. The value of ABI (<1 or ≥1) and IABD (<10 or ≥ 10 mmHg) was deduced in all the subjects. The groups were compared using the clinical features, ABI and IABD values as variables.

Results: Odds for BZI were high in men (OR 1.78, 95% CI 0.87-3.66), hypertension (p=0.003, OR 2.79, 95% CI 1.39 to 5.6), history of ischemic heart disease (OR 2.06, 95% CI 0.73-5.8), prior stroke (OR 1.92, 95% CI 0.54-6.8), smoking (OR 1.85, 95% CI 0.96-3.56) and dyslipidemia (OR 2.5, 95% CI 0.83-7.58). ABI mean (ABI-M) <1 was significantly associated with BZI (p=0.022) as compared to CATI. ABI-M of <0.96 was able to diagnose BZI over CATI (sensitivity 56.2%, specificity 63.8%, p<0.05). IABD ≥10 mm Hg showed no association with BZI. Logistic regression analysis showed that HTN (p<0.001) and low ABI-M (p<0.033) were a significant predictor of BZI.

Conclusion: Low ABI-M could be of value as a screening tool to identify persons at risk for BZI in stroke population.

Keywords: Border-zone infarct; ankle-brachial index; inter-arm blood pressure difference; stroke; intracranial arterial stenosis; peripheral arterial disease; hypertension

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Introduction

Stroke is potentially preventable yet a devastating cause of mortality and morbidity worldwide [1]. Despite big advancements in the secondary prophylaxis strategies the ischemic strokes continue to recur (RS), at times with alarmingly high rate [2]. Furthermore, some ethnic groups seem to be at higher risk for RS. Recently, a group of Chinese researchers has cited a threefold increased incidence of RS between 2006 and 2012 [3].

Three patterns of ischemic infarcts have been described: (a) confined to the perforator territory; (b) lying within the core vascular territory of a major intracranial artery (CATI) and (c) located at the distal part of two non-anastomosing arterial systems, the border-zone infarcts (BZI) [4]. Later are further divided into internal BZI (IBZ) and external BZI (EBZ). Of these two the former is located between the deep and superficial arterial system of anterior circulation and the later between the cortical territories of the three major vessels, viz. middle (MCA), anterior (ACA) and posterior cerebral (PCA) arteries [5].

It is noteworthy that preventive strategies often yield poor results in BZI patients. Despite aggressive medical management, compared to CATI persons with BZI continue to have a higher risk of RS (8% vs. 2.6%) [2]. Therefore, extensive research has been done to understand the mechanisms of BZI stroke [6]. It is known that the BZI differs from CATI not only in the etiopathogenesis but also in their course and prognosis [7]. In experimental studies systemic hypotension has produced BZI [8]. BZI has been linked with clinical situations leading to systemic hypotension, including iatrogenic [9]. Recent research has linked BZI with hypoperfusion due to higher degree of intracranial arterial stenosis [2,10,11]. It has been suggested to study the hemodynamics of the cerebral blood flow before individualizing the therapy for RS prevention [2].

In countries like India, where high-tech medical care is not accessible to a significant proportion of the population, there is an urgent need to find some easy way of identifying those at risk of RS. Recording blood pressure in all four limbs and observing the inter-arm blood pressure difference (IABD) and ankle-brachial index (ABI) is an inexpensive screening test for PAD [12, 13].

The association of IABD of ≥10mmHg and ABI of <1 or ≤0.9 with intracranial atherosclerosis and RS and/or stroke is already known [13-15]. Although the relationship of ABI and IABD with ischemic stroke is well established, to the best of our knowledge their association with BZI alone is not known.

We, therefore, set to study the association of ABI and IABD with BZI in the subjects with acute ischemic stroke. We also planned to look into the clinical and demographic factors associated with BZI.

Material, methods and data management

The study was conducted in the neurology department of a medical college hospital. After obtaining informed consent, we prospectively enrolled patients aged >18 years admitted with acute ischemic stroke (n=355, 71% male) from March 2018 through July 2019. Subjects with recent head trauma (n=2), vasculopathy (n=5), stroke mimic (n=4) and those who could not undergo MRI of the brain (n=141) were excluded (Figure 1). Remaining 183 subjects (78% male) formed the study population. Every patient was thoroughly examined by one neurology resident (KT, PR) and a senior neurologist, later was not associated with the study.

The gender and age were noted. History of prior stroke, ischemic heart disease (IHD), hypertension (HTN), diabetes (T2DM), alcohol use and smoking were sought for. Old medical records, medications were reviewed for the evidence of HTN, IHD, smoking, T2DM.

Supine four limb BP was measured in all the subjects by a trained and supervised team of nurses using an automated device (PHILIPS Intellivue MP 20).
according to the available guideline [15]. The upper extremity BP was recorded over the brachial artery and that in the lower extremity was measured over the posterior tibial artery at the ankle. Three readings of the systolic BP (SBP) and diastolic BP (DBP) were noted at each site.

ABI, defined as the ratio of SBP measured over posterior tibial and brachial arteries, was calculated as described [17]. We thus, calculated the ABI using SBP at the ankle as numerator while the highest value of the SBP, of either arm, remained the denominator. Thereby obtaining three separate ABI values- a) ABI high (ABI-H) calculated using the highest recorded SBP b) ABI mean (ABI-M) calculated with the mean of the three SBP readings and c) ABI low (ABI-L) calculated by utilizing the lowest SBP recorded. We thus calculated the ABI using SBP at the ankle as numerator while the highest value of the SBP, of either arm, remained the denominator. Thereby obtaining three separate ABI values- a) ABI high (ABI-H) calculated using the highest recorded SBP b) ABI mean (ABI-M) calculated with the mean of the three SBP readings and c) ABI low (ABI-L) calculated by utilizing the lowest SBP recorded. We thus calculated the ABI using SBP at the ankle as numerator while the highest value of the SBP, of either arm, remained the denominator.

Inter-arm BP difference for systolic (IASBD) and diastolic BP (IADBD) was calculated as per Chang et al.[13]. A difference of ≥10mm Hg was taken as significant.

Complete blood and biochemical profile, electrocardiogram (EKG), lipid profile, HBA1C were done in each subject and values were recorded in the study proforma.

MRI brain was performed using a 1.5 T, 18 channel system (Magnetom Symphony, Siemens Medical Solutions, Germany). The raw data was stored in the hospital server and was discretely retrieved later by two authors, blinded for the clinical features and BP recordings (AKS and RJ). The diffusion-weighted (DWI), T2 weighted and fluid-attenuated inversion recovery (FLAIR) images were studied. DWI images were correlated with the apparent diffusion coefficient (ADC) maps [19]. The CATI, IBZ, and EBZ were identified and recorded on a template as described previously [20]. Peri-ventricular confluent, white matter hypertensive (leukoaraiosis) changes and discrete, focal, small T2WI hyper-intense lesions were ignored. Any discrepancy in the observation was sorted out with the help of a neuroradiologist (MB).

Subjects were divided according to age (<60, ≥60 yr), gender (male/ female) and presence or absence of the above-mentioned risk factors. Persons with a history of HTN, or those with SBP >140, DBP>90 on repeated measurement and patients with LVH on EKG or 2D Echo were considered to be hypertensive [21]. Patients with glycosylated hemoglobin (HBA1C) level of ≥6.5 were considered to have T2DM [21]. Dyslipidemia was defined as total cholesterol ≥220 mg/dl or LDL≥160 mg/dl [21].

The subjects with acute or chronic EBZ and IBZ, with or without CATI, were grouped as BZI (n=89). Those with pure CATI (and no acute/chronic BZI) were grouped as CATI patients (n=94). The groups were compared for the associated risk factors and the values of the IABD and ABI.

Statistical technique

The statistical analysis was done using software like Medcalc [22], Graphpad [23], Excel 2007 [24] and SPSS version 17.0 Trial. Results of continuous data are presented as Mean ± Standard deviation (SD) and the results of the categorical measurements are given as a percentage (%).

As and when required, Pearson’s Chi-Square test, two-tailed Fisher exact test and comparison of proportions were deployed to know the association of various risk factors with border zone infarcts. Multivariate binary logistic regression was used to predict the border zone infarcts (dependent variable) among patients by employing selected continuous and non-continuous predictors (independent variables).

The probability value p ≤ 0.05 and OR >1 (at 95% CI) was considered significant. The study was approved by the Institutional Scientific and Research Committee.

Results

The clinical and demographic data are presented in Table 1.

Out of 183 subjects, BZI was detected in 89 (48.63%) while 94 (51.37%) had CATI. The difference was non-significant (p=0.69). The values of DBP, mean age and HbA1C did not differ significantly between BZI and CATI groups.

As compared to the CATI group the mean SBP at right arm was significantly high in the BZI group. (138 ±23.34 vs. 145 ±23.18 mm Hg, p=0.04; table-1, section 1).
### Table 1: Demographic and clinical profile of study population (n=183).

| Variable | Borderzone infarcts (BZI) | Core Arterial Territory Infarcts (CATI) | OR (95% CI) | p-value (LOS) |
|----------|---------------------------|----------------------------------------|-------------|---------------|
| **Section-I** |
| Age in yr (mean ± SD) | 57.02±13.15 | 58.37±14.57 | - | 0.51 **ns,§** |
| SBP in mmHg (mean ±SD) | 145±23.18 | 138±23.34 | - | 0.04 **s,§** |
| DBP in mmHg (mean ±SD) | 83.87±12.84 | 80.06±14.26 | - | 0.10 **ns,§** |
| HBA1C (mean ±SD) | 6.85±2.24 | 6.68±2.01 | - | 0.58 **ns,§** |

| **Section-II** |
| Gender | Male n=143 (100) | 74 (51.74) | 69 (48.26) | 1.78 (0.87-3.66) | 0.15 **ns** |
| | Female n=40 (100) | 15 (37.5) | 25 (62.5) | - | - |
| Age | ≥60 86(100) | 41 (47.67) | 45 (52.33) | 0.93 (0.52-1.66) | 0.88 **ns** |
| | <60 97(100) | 48 (49.48) | 49 (50.52) | - | - |
| T2DM | Yes n=78 (100) | 37 (47.43) | 41 (52.57) | 0.91 (0.51-1.65) | 0.78 **ns** |
| | No n=105 (100) | 52 (49.52) | 53 (50.48) | - | - |
| HTN | Yes n=134 (100) | 74 (55.22) | 60 (44.78) | 2.79 (1.39-5.6) | 0.003 **p** |
| | No n=49 (100) | 15 (30.61) | 34 (69.39) | - | - |
| History of IHD | Yes n=17 (100) | 11 (64.70) | 6 (35.30) | 2.06 (0.73-5.8) | 0.16 **ns** |
| | No n=166 (100) | 78 (46.98) | 88 (53.02) | - | - |
| Prior Stroke | Yes n=11(100) | 7 (63.63) | 4 (36.37) | 1.92 (0.54-6.8) | 0.30 **ns** |
| | No n=172 (100) | 82 (47.67) | 90 (52.33) | - | - |
| History of Smoking | Yes n=52 (100) | 31 (59.61) | 21 (40.39) | 1.85 (0.96-3.56) | 0.061 **ns** |
| | No n=131 (100) | 58 (44.27) | 73 (55.73) | - | - |
| Alcohol use | Yes n=38 (100) | 22 (57.89) | 16 (42.11) | 1.60 (0.77-3.29) | 0.19 **ns** |
| | No n=145 (100) | 67 (46.20) | 78 (53.80) | - | - |
| Dyslipidemia | Yes n=16 (100) | 11 (68.75) | 5 (31.25) | 2.5 (0.83-7.58) | 0.11 **ns** |
| | No n=150 (100) | 70 (46.66) | 80 (53.34) | - | - |

BZI- Border zone infarct, CATI- Core Arterial Territory infarct, **§** Comparisons of means, ^calculated as per Altman, 1991, *2 tailed Fischer Exact test, †Pearson Chi-Square test, ‡Comparison of proportion as per Altman, 1991, ϕcalculated in 166 subjects, data missing n=17, CI-confidence interval, OR - odds ratio, SBP-systolic blood pressure, DBP-Diastolic blood pressure, LOS-Level of significance, s-significant, ns-non significant.
A summary of the demographic and clinical profile of the study population is given in table 1 section II. We found that males have higher odds (1.98, 95%CI 0.87-3.66) for having BZI, although the probability did not reach statistical significance (p=0.15). Similarly, those with prior stroke and a history of IHD showed higher odds for having BZI but the difference did not reach the statistical significance.

The results suggest that HTN (n=134) has a strong association with the occurrence of BZI (OR 2.79, CI 1.39-5.6; p=0.003). Similarly, smokers (n=52) showed a trend for higher risk for BZI (OR 1.85, CI 0.96-3.56; p=0.06).

The data of the lipid profile of 17 patients were missing. Results were thus calculated for the remaining 166 subjects, which depicts higher odds for BZI in patients with dyslipidemia (n= 16; OR 2.5, CI 0.83-7.58). Odds for BZI occurrence were slightly increased (OR 1.6, CI 0.77-3.29) in those who consumed excess alcohol (n= 38).

We did not find any association of IASBD (OR 0.53 CI 0.27-1.03, p=0.06) and IADBD (OR 0.96 CI 0.49-1.86, p=0.89) with BZI. The relationship of ABI with stroke pattern is given in the Table 2.

An abnormal ABI (<1) for three different techniques of calculation, ABI-H, ABI-M, and ABI-L, was found in 50.2, 64.48 and 80.87% respectively. Out of them, no association of BZI with ABI-H (p=0.089) and ABI-L (p=0.88) was seen. But ABI-M <1 showed significant association with BZI (p=0.022). A receiver operating characteristic (ROC) curve was drawn to know the value of low ABI-M in the diagnosis of BZI over CATI. We found that ABI-M of <0.96 was able to diagnose BZI (sensitivity 56.2%, specificity 63.8%, p<0.05) over CATI. (Figure 2).

Although the highest abnormality was detected in ABI-L, but compared to the CATI, the risk of having BZI was associated only with low ABI-M (p=0.022). To understand this phenomenon we looked into systolic BP of individual patients grouped under above mentioned three ABI sub-groups and calculated the mean SBP at right arm for each group. Results showed that compared to the other two ABI groups the difference in the mean SBP of low (<1) and high (≥1) ABI-M group was significant (p=0.05, Table-3).

Multivariate binary logistic regression was performed to identify the significant predictor for BZI at 95% CI. The Table 4 summarizes the binary logistic regression model. Out of the incorporated risk factors only HTN (p<0.001) and ABI-M (p<0.033) are found to be the most significant predictor of BZI, the probability of the correct prediction being 67.2% (sensitivity 67.4% and specificity 67%). The presence of dyslipidemia and the male gender had weak predictive value for BZI (p<0.06).

**Discussion**

The presence of a severe degree of ICAS and hypoperfusion has been implicated in the etiopathogenesis of the BZI [8-10, 25]. Patients with cerebral vascular stenosis with poor collateral circulation when exposed to aggressive BP management, the presence of ICAS predisposes them to BZI and RS. Further, increasing BP improves the prognosis of BZI patients [26, 27]. Therefore, knowledge of ICAS seems essential to prevent and manage BZI [2]. The diagnosis of ICAS is possible with sophisticated medical gadgets but they are not of much use as a screening tool because of the high cost and poor availability.
### Table 2: Interlimb blood pressure difference vs. Borderzone infarcts.

| Variable | Borderzone infarcts (BZI) (n=89) | Core Arterial Territory Infarcts (CATI) (n=94) | OR (95% CI) | p-value (LOS) |
|----------|----------------------------------|-----------------------------------------------|-------------|---------------|
| ABI-H    | < 1.0 (92) 39 (42.95)            | 53 (57.60)                                    | 0.772       | 0.089         |
|          | ≥ 1.0 (91) 50 (53.84)            | 41 (46.16)                                    |             |               |
| ABI-M    | < 1.0 (118) 50 (42.38)           | 68 (57.62)                                    | 0.49        | 0.022*        |
|          | ≥ 1.0 (65) 39 (60.00)            | 26 (40.00)                                    |             |               |
| ABI-L    | < 1 (148) 69 (46.62)             | 79 (53.38)                                    | 0.65        | 0.88**        |
|          | ≥ 1 (35) 20 (57.14)              | 15 (42.86)                                    |             |               |
| IASBD    | ≥ 10 (53) 20 (37.73)             | 33 (62.26)                                    | 0.53        | 0.060**       |
|          | < 10 (130) 69 (53.07)            | 61 (46.92)                                    |             |               |
| IADBD    | ≥ 10 (46) 22 (47.82)             | 24 (52.18)                                    | 0.958       | 0.89**        |
|          | < 10 (137) 67 (48.90)            | 70 (51.10)                                    |             |               |

*calculated as per Altman, 1991, ± Pearson Chi-Square test, ABI-Ankle brachial index, OR-odds ratio, CI-confidence interval, LOS-Level of significance, IASBD- Interarm systolic blood pressure difference, IADBD- Interarm diastolic blood pressure difference, s-significant, ns-non significant.

### Table 3: ABI and mean systolic blood pressure at right arm.

| ABI ratio | Systolic BP mmHg mean ± SD |
|-----------|-----------------------------|
|           | ABI-H | ABI-M | ABI-L |
| < 1       | 144.98 ±24.9                | 144 ±23.5 | 142.35±23.51 |
| ≥ 1       | 138.71±21.56               | 137±22.88 | 139.8±23.43 |
| p-value (LOS)* | 0.70** | 0.05* | 0.56** |

*Comparison of means, ns=nonsignificant, s= significant, LOS-Level of significance, ABI-Ankle brachial index.

A strong association of PAD with ICAS and its easy identification by measuring the IABD and ABI is well established [13, 15]. In most of the previous research, patients have been recruited retrospectively to study BZI either to identify the stroke mechanisms or cerebral vascular abnormalities or MRI characteristics of the BZI.

Through our single-blind study, we have attempted to identify predictors of BZI and explore the association of MRI discernible BZI and inter-arm BP difference in consecutive acute stroke patients by deploying IASBD, IADBD and ABI measurements as tools. To best of our knowledge, no such work has been done in the past.

Since our study design and patient population differ from the earlier works so do our demographic findings. For example, we recruited subjects consecutively and found BZI in 48.63% of them. Young et al. attempted to study the mechanism of IBZ and EBZ infarcts in MCA territory by retrospectively recruiting the stroke patients based upon vascular studies. They found BZI in only 12.7% of their patients [25]. In contrast to them, Bogousslavsky et al., recruited consecutive patients with acute stroke and found BZI in 53% of their subjects [28].
Table 4: Multivariate binary logistic regression analysis of cerebrovascular risk factors for borderzone infarcts.

| Predictors            | \( \beta \) | Std. Error | p-value (LOS) | Odds ratio | 95% CI        |
|-----------------------|-------------|------------|---------------|------------|---------------|
| Age (Year)            | -0.012      | 0.013      | p>0.05 \textsuperscript{ns} | 0.99       | 0.96 - 1.01   |
| Sex (Male)            | 0.676       | 0.448      | p<0.06 \textsuperscript{ns} | 1.97       | 0.82 - 4.73   |
| Hypertension          | 1.431       | 0.420      | p<0.001 \textsuperscript{s} | 4.18       | 1.84 - 9.53   |
| DM (Present)          | -0.353      | 0.355      | p>0.05 \textsuperscript{ns} | 0.70       | 0.35 - 1.41   |
| Smoking (Y)           | 0.451       | 0.442      | p>0.05 \textsuperscript{ns} | 1.57       | 0.66 - 3.74   |
| Alcohol (Y)           | -0.031      | 0.486      | p>0.05 \textsuperscript{ns} | 0.97       | 0.37 - 2.51   |
| Prior Stroke History (Y) | 1.184   | 0.784      | p>0.05 \textsuperscript{ns} | 3.27       | 0.70 - 15.19  |
| Past IHD Hist. (Y)    | 0.935       | 0.629      | p>0.05 \textsuperscript{ns} | 2.55       | 0.74 - 8.74   |
| Dyslipidemia (Y)      | 1.570       | 0.826      | p<0.06 \textsuperscript{ns} | 4.81       | 0.95 - 24.25  |
| IASBP (≥10 mmHg)      | -0.499      | 0.406      | p>0.05 \textsuperscript{ns} | 0.61       | 0.27 - 1.35   |
| IADBP (≥10 mmHg)      | 0.096       | 0.393      | p>0.05 \textsuperscript{ns} | 1.10       | 0.51 - 2.38   |
| ABI-Mean (<1.0)       | -0.755      | 0.354      | p<0.033 \textsuperscript{**} | 0.47       | 0.24 - 0.94   |

Y-Yes, N-No; LOS- Level of significance; CI- Confidence interval, Log odds of border zone infarcts = - 0.879 - 0.012 \times \text{Age (year)} + 0.676 \times \text{Sex (1 If male and 0 If female)} + 1.431 \times \text{Hypertension (1 If present and 0 If absent)} - 0.353 \times \text{Diabetes Mellitus (1 If present and 0 If absent)} + 0.451 \times \text{Smoking (1 If present and 0 If absent)} - 0.031 \times \text{Alcohol (1 If present and 0 If absent)} + 1.184 \times \text{History of Past Stroke (1 If present and 0 If absent)} + 0.935 \times \text{History of Ischemic Heart Disease (1 If present and 0 If absent)} + 0.255 \times \text{Dyslipidemia (1 If absent and 0 If present)} + 0.499 \times \text{Inter-arm systolic blood pressure difference (1 If ≥ 10 mmHg and 0 If < 10 mmHg)} + 0.096 \times \text{Inter-arm diastolic blood pressure difference (1 If ≥ 10 mmHg and 0 If < 10 mmHg)} - 0.755 \times \text{ankle-brachial index average (1 If < 1 and 0 If ≥ 1).}

The previous research on BZI can be divided into two broad groups: (a) those which recruited patients only of BZI and studied the mechanisms of IBZ and EBZ infarcts; (b) those which aimed to compare the CATI with BZI. Since the former type of studies made a comparison between the subgroups of BZI, hence did not find much significant difference in the demographic or clinical risk factor other than HTN [11, 24, 28]. In contrast to first the second type of studies has reported age, HTN, dyslipidemia, male gender, and T2DM to have higher odds for having BZI [30, 31]. We compared BZI with CATI and found that in addition to HTN, patients with male gender, dyslipidemia, IHD, prior stroke, and smoking have higher odds of having BZI.

Regarding inter-arm BP difference we have the following observations: (a) although the low ABI-M (<1) is associated with BZI (p=0.033) but it predicts BZI over CATI with rather low sensitivity and specificity (56.2 and 63.8 respectively). Low ABI is known to be associated with CATI also [15]. Because we compared BZI with the CATI patients the predictive strength probably appears low; (b) High IADBP and IASBP have no association with BZI as compared with CATI. At first glance, this observation might appear to be anomalous but is not unexpected. IASBD and IADBD have a strong association with PAD but not with ICAS [32]. In contrast, ABI has a strong association with ICAS [33]. (c) The predictive value for BZI of ABI-M (<1) is higher than ABI-H and ABI-L because the former incorporated the mean of SBP while later two take into consideration either the highest or the lower value of SBP. This observation is supported by the fact that the mean SBP in our patients with abnormal (<1) ABI-M was significantly higher than that in persons with ABI-M value of >1. (Table 3).
Our results suggest that low ABI-M, although not specific, is a sensitive method to identify persons at risk of BZI, particularly in the presence of HTN.

The collateral circulation is also a critical determinant of cerebral perfusion pressure in acute cerebral ischemia [34]. It is known that the development of BZI depends upon the status of the cerebral collateral circulation, which is known to be suboptimal in persons with BZI [2]. The development of collaterals is further impaired in presence of systemic HTN [35]. Catena et al. concluded that hypertension is a strong risk factor of atherosclerosis and carotid artery stenosis which in turn are risks of internal BZI [36]. Our results also suggest that ABI-M (<1) with hypertension could be a surrogate marker of ICAS and therefore is associated with BZI.

**Conclusion**

Low ABI-M (<1) could be a marker for the presence of ICAS in patients of ischemic stroke especially in patients with BZI. It is easy to calculate and could be used to identify patients at risk for BZI in the stroke population. Subset of stroke patients with HTN, if also have low ABI-M should be investigated further for presence of BZI and ICAS before aggressive BP control.

Our study has certain limitations arising due to the exclusion of a significant number of patients for want of the MRI. The predictive value of low ABI-M should be further studied in a multicentric study incorporating different ethnic groups and the normal population as control.

**Main points:**

1. Ankle-brachial index (ABI) is easy to calculate and can be used to identify patients at risk for BZI in the stroke population.
2. Low ABI-Mean (<1) can be a marker for the presence of ICAS in patients of ischemic stroke especially in patients with border zone infarcts (BZI).
3. Subset of stroke patients with HTN, if also have low ABI-Mean should be investigated further for presence of BZI and intracranial arterial stenosis before aggressive BP control.

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**Conflicts of interest**

Authors declare no conflicts of interest.

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