

A phantom study on the influence of viscosity of blood, percentage diameter stenosis, length of stenosis and cross-sectional area of blood vessel on blood flow dynamics in peripheral (brachial and femoral) artery models using Doppler ultrasound spectral analysis

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Abstract

Objectives: To study the influence of viscosity of blood, percentage diameter stenosis, length of stenosis and cross-sectional area of blood vessel on blood flow dynamics in peripheral (brachial and femoral) artery models using Doppler ultrasound spectral analysis. **Methods/Analysis:** An experimental study was carried out using an indigenously fabricated Doppler ultrasound compatible blood flow phantom. Doppler frequency spectral analysis was the method used for the study. To study the influence of various factors on blood flow dynamics, brachial and femoral artery models with 25%, 50%, 75% diameter stenosis of 5mm, 10mm lengths and 2cP, 3cP, 4cP viscosity blood mimicking fluids (BMF) were used. To study the influence of the parameters, the change in ratios of Pre and Post stenotic Peak Systolic Velocities (PSV ratio) obtained with respect to change in the parameter were assessed. **Findings:** The magnitude and pattern of change in the PSV ratio with respect to pre-destined values of viscosity, percentage diameter stenosis, length of stenosis and cross-sectional area of the blood vessel were analysed. **Novelty:** As the analysis of flow dynamics under such conditions of variability cannot be exercised clinically (as a patient-based study), a model-based experimental study was performed with a cost-effective indigenously fabricated Doppler ultrasound blood flow phantom.

Key Words: Blood flow dynamics, Blood viscosity, Doppler spectral analysis, Length of stenosis, Percentage diameter stenosis, Peripheral artery models.

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INTRODUCTION

Vascular resistance is one of the major factors influencing blood flow dynamics. The vascular resistance mainly depends on the cross-sectional area of blood vessel, blood viscosity and length of the blood vessel. Blood viscosity is the inherent resistance of blood to flow. An important factor which determines the blood flow dynamics is blood viscosity as it is the only biological parameter that has been correlated with all of the major cardiovascular risk factors². A normal human's whole blood viscosity is around 3 centipoise^{3,4}. Due to aging or under certain

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diseased conditions or under certain medications, blood viscosity may become lesser or greater than the normal value and this change may lead to changes in blood flow dynamics. The cross-sectional area ($\pi D^2/4$) of blood vessel is directly proportional to blood flow capacity and has the potential to modify blood flow dynamics. Blood flow capacity within two vessels of diameter D_1 and D_2 can be compared using a ratio of their diameter squared. $[(D_2)^2:(D_1)^2]$ Arterial stenosis is narrowing or constriction of lumen of the artery, usually caused by atherosclerosis. Two facets of stenosis are percentage diameter stenosis and length of stenosis. Out of the peripheral arteries, femoral artery is a common site for narrowing or obstruction that occurs as a result of the atherosclerotic process. The atherosclerotic stenosis in peripheral arteries may lead to peripheral artery diseases (PAD) which will affect the normal blood flow pattern and thereby reduce flow of blood to the corresponding parts. The peripheral artery diseases may be asymptomatic or symptomatic, but when there is occlusion they have proved to show close relationship with increased risk of coronary heart disease, heart attack, stroke and transient ischemic attack (mini stroke). Apart from imaging the sections of patient's body and helping in taking accurate measurements of structures within, ultrasound is also being used for performing blood flow studies using the Doppler Effect. Doppler spectral analysis is a quantitative method which shows the distribution of Doppler frequency signals within the operator selected sample volume of the blood vessel over a period of time and hence is useful to do blood flow studies.

MATERIALS AND METHODS

Doppler ultrasound machine - GE logic P5, linear array ultrasound probe of 12 MHz frequency, instead of patients / volunteers a Doppler ultrasound compatible blood flow phantom which induces pulsatile flow of blood mimicking fluid within brachial and femoral artery models presented with various percentage diameter stenotic conditions and lengths of stenosis kept embedded in tissue equivalent material¹. The average inner diameters (ID) of brachial artery and femoral artery of normal healthy young adults were found to be around 4mm and 8mm respectively¹. Ultrasound compatible blood vessel equivalent tubes with these specifications were used for the study¹. The analysis of blood flow is done by Doppler spectrum. The spectral display is obtained using Fast Fourier Transformation (FFT) analyser which gets the distribution of Doppler frequency signals within the operator selected sample volume and in turn shows the velocity of blood flow. In this study the influence of parameters such as viscosity of blood ($\eta_1 = 2$ cP, $\eta_2 = 3$ cP and $\eta_3 = 4$ cP), percentage diameter stenosis (25%, 50% and 75% - symmetrical), length of stenosis (5mm and 10mm), and cross-sectional

area of blood vessel (12.57 mm^2 for 4mm ID brachial artery and 50.29 mm^2 for 8mm ID femoral artery) on blood flow dynamics in brachial artery models and femoral artery models were analysed by varying only the influential parameter under study and keeping all other factors constant. The analysis of blood flow dynamics under such conditions of variability cannot be exercised clinically (as a patient-based study), hence a model-based experimental study was performed with a cost-effective indigenously fabricated Doppler ultrasound blood flow phantom¹. Ultrasound scanner is initially operated in B-scan mode to image the vessel model. The Doppler ultrasound spectral display was activated by switching over to Pulsed Doppler mode in ultrasound scanner. The sample volume (SV of 1mm) is set at a location within the vessel before the simulated stenosed region. The Peak Systolic Velocities (PSV) of the Doppler Spectrum at the pre-stenotic region and also at the post-stenotic region of the blood vessel model noted. The same is repeated by using various other simulated conditions. The ratio of Post-stenotic PSV to Pre-stenotic PSV known as PSV ratio is calculated [Table - 1 and Table - 2]. The PSV ratio denotes the change in the flow dynamics due to the influence of the parameter under analysis. PSV ratio is taken into consideration rather than absolute PSV as there is a greater correlation observed between PSV ratio and stenosis than between absolute PSV and stenosis⁴. The various pre-destined flow conditions were carried out and the influences of each parameter on the blood flow dynamics were analysed.

OBSERVATIONS AND RESULTS

Doppler Spectral analysis and flow data acquired using brachial and femoral artery models:

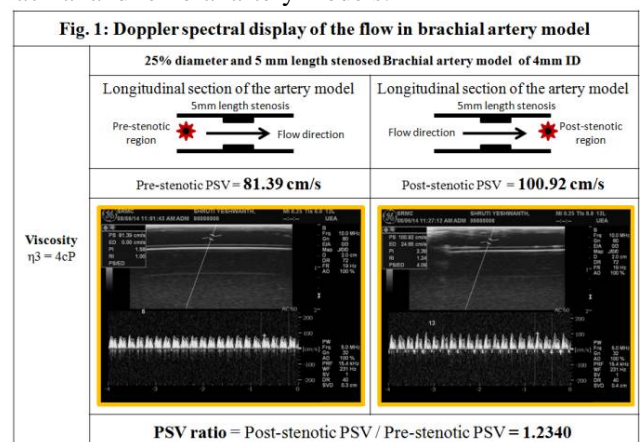


Figure 1:

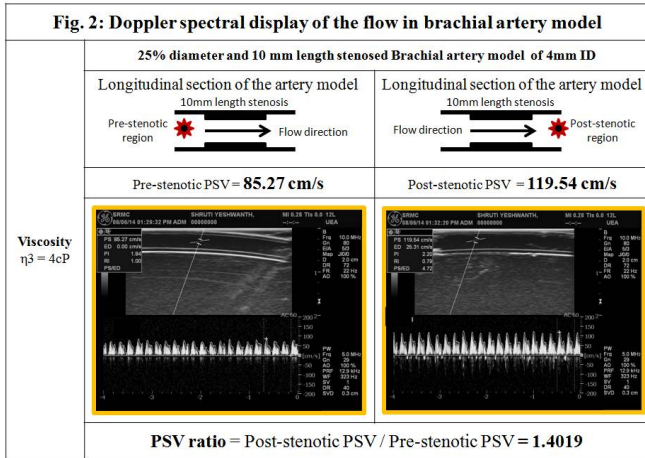


Figure 2:

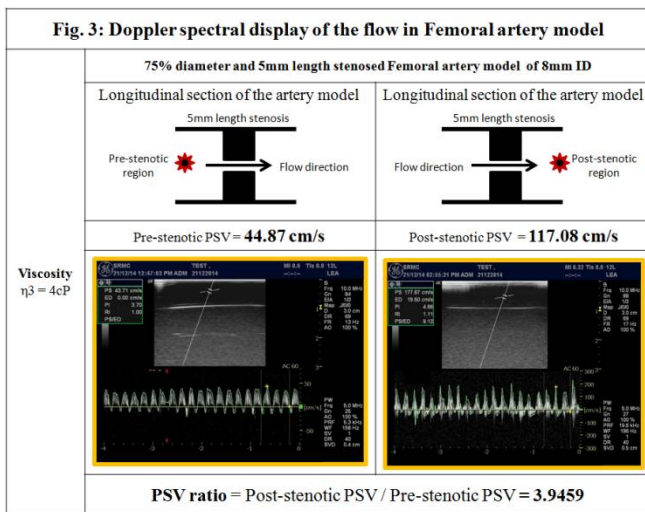


Figure 3:

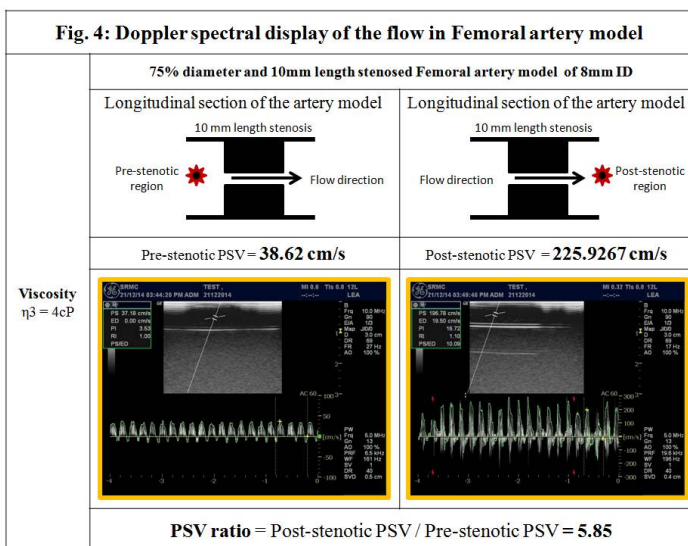


Figure 4:

Table 1: Flow data of Brachial Artery model

Viscosity of BMF (cP)	Length of Stenosis (mm)	% Diameter Stenosis	PSV ratio
$\eta_1 = 2$	5	25	1.1956
		50	1.5792
		75	1.7743
$\eta_2 = 3$	10	25	1.3032
		50	1.3744
		75	1.9564
$\eta_3 = 4$	5	25	1.2085
		50	1.5843
	10	50	2.0716
		75	1.3686
	10	50	1.5501
		75	2.3353
25		1.2340	
50		1.7374	
75		2.6245	
10	10	25	1.4019
		50	1.7328
		75	2.4859

BMF – Blood Mimicking Fluid, cP – centipoise, PSV – Peak Systolic Velocity

Table 2: Flow data of Femoral Artery model

Viscosity of BMF (cP)	Length of Stenosis (mm)	% Diameter Stenosis	PSV ratio
$\eta_1 = 2$	5	25	1.1300
		50	1.2776
		75	2.6887
$\eta_2 = 3$	10	25	1.0361
		50	1.4559
		75	2.8014
$\eta_3 = 4$	5	25	1.1367
		50	1.9790
	10	50	3.9459
		75	1.0996
	10	50	1.5031
		75	4.7863
25		1.1790	
50		2.2289	
75		5.3239	
10	10	25	1.1025
		50	2.3566
		75	5.8500

BMF – Blood Mimicking Fluid, cP – centipoise, PSV – Peak Systolic Velocity

INTERPRETATION AND DISCUSSION:

BRACHIAL ARTERY MODEL - 5mm length of stenosis

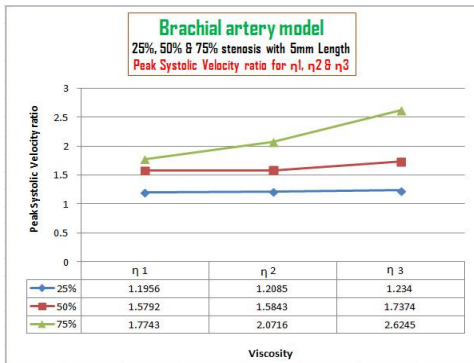


Figure 5: PSV ratio with respect to the variation in viscosity in Brachial artery models with stenosis of 5mm length having 25%, 50% and 75% diameter stenosis

FEMORAL ARTERY MODEL - 10mm length of stenosis

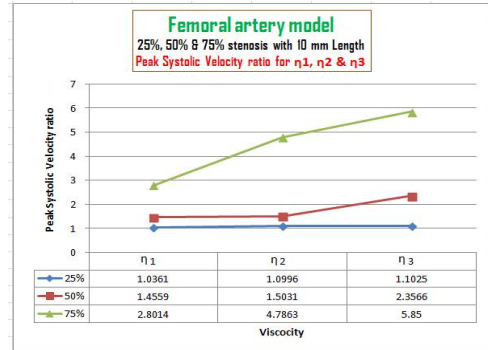


Figure 8: PSV ratio with respect to the variation in viscosity in Femoral artery models with stenosis of 10mm length having 25%, 50% and 75% diameter stenosis

BRACHIAL ARTERY MODEL - 10mm length of stenosis

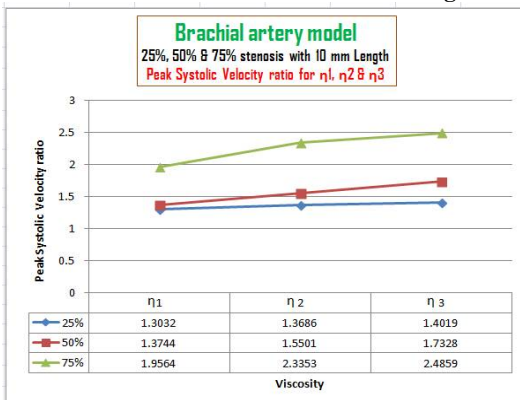


Figure 6: PSV ratio with respect to the variation in viscosity in Brachial artery models with stenosis of 10mm length having 25%, 50% and 75% diameter stenosis

BRACHIAL ARTERY MODEL - 5mm length of stenosis

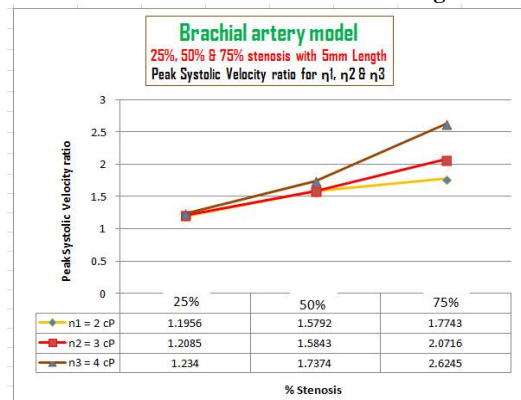


Figure 9: PSV ratio with respect to the percentage diameter stenosis in Brachial artery model with 5mm length of stenosis for various blood mimic viscosities

FEMORAL ARTERY MODEL - 5mm length of stenosis

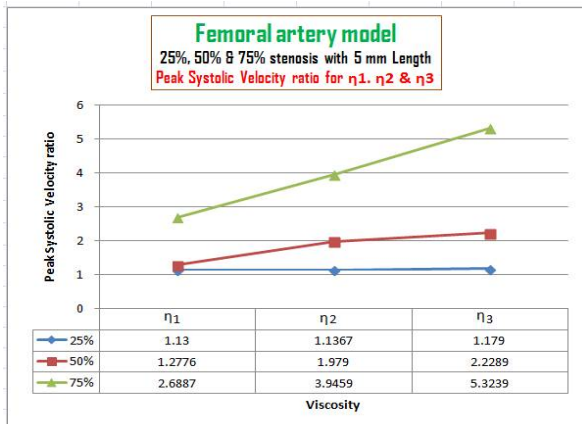


Figure 7: PSV ratio with respect to the variation in viscosity in Femoral artery models with stenosis of 5mm length having 25%, 50% and 75% diameter stenosis

BRACHIAL ARTERY MODEL - 10mm length of stenosis

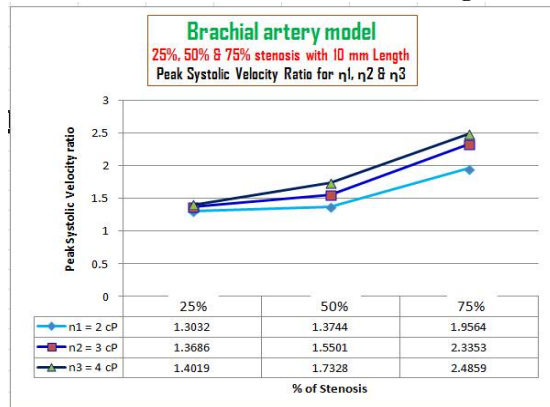


Figure 10: PSV ratio with respect to the percentage diameter stenosis in Brachial artery model with 10mm length of stenosis for various blood mimic viscosities

FEMORAL ARTERY MODEL - 5mm length of stenosis

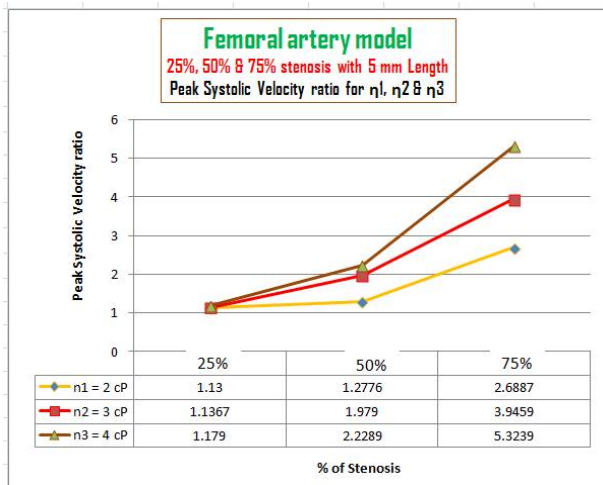


Figure 11: PSV ratio with respect to the percentage diameter stenosis in Femoral artery model with 5mm length of stenosis for various blood mimic viscosities

FEMORAL ARTERY MODEL - 10mm length of stenosis

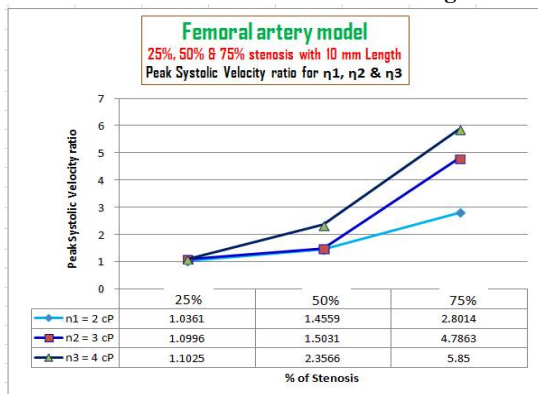


Figure 12: PSV ratio with respect to the percentage diameter stenosis in Femoral artery model with 10mm length of stenosis for various blood mimic viscosities

Influence of Viscosity of BMF: With variation in viscosity of the blood mimic, the PSV ratio of pre-and post-stenotic regions under various simulated conditions (Figures 5, 6, 7 and 8) are as follows. With increased viscosity of the blood mimic, for both 5mm and 10mm lengths of stenosis,

- Under 25% diameter stenotic conditions in the Brachial artery and Femoral artery models, there is trivial increase in PSV ratio of pre-and post-stenotic regions.
- Under 50% diameter stenotic conditions in the Brachial artery and Femoral artery models, there is notable and gradual increase in PSV ratio of pre-and post-stenotic regions.

- Under 75% diameter stenotic conditions in Brachial artery and Femoral artery models, there is significant increase in PSV ratio of pre-and post-stenotic regions.

With increased viscosity condition, the higher percentage diameter stenosis seems to show increased variation in PSV ratio.

Influence of Percentage diameter Stenosis: With the variation in percentage diameter stenosis, there will be a variation in the cross-sectional area of the lumen. This variation affects the flow capacity and hence percentage diameter stenosis is expected to be a more prominent factor to significantly change the flow pattern.

Table 3: Flow capacity variation due to percentage diameter stenosis in Brachial artery model

BRACHIAL ARTERY MODEL				
% Diameter stenosis	0%	25%	50%	75%
Diameter of the vessel (mm)	4	3	2	1
Cross-sectional area (mm ²) = πD ² /4	12.57	7.07	3.14	0.786
Relative Flow capacity to 0% stenosis - (D ₂) ² : (D ₁) ²	-	4 ² :3 ² 1.78 times lesser	4 ² :2 ² 4 times lesser	4 ² :1 ² 16 times lesser

Table 4: Flow capacity variation due to percentage diameter stenosis in Femoral artery model

FEMORAL ARTERY MODEL				
% Diameter stenosis	0%	25%	50%	75%
Diameter of the vessel (mm)	8	6	4	2
Cross-sectional area (mm ²)	50.29	28.29	12.57	3.14
Relative Flow capacity to 0% stenosis - (D ₂) ² : (D ₁) ²	-	8 ² :6 ² 1.78 times lesser	8 ² :4 ² 4 times lesser	8 ² :2 ² 16 times lesser

With increased percentage of stenosis, under constant viscosity flow in the Brachial artery and Femoral artery models, there is significant and also steep increase in PSV ratio which makes the curve look like a top-opening parabola (Figures 9, 10, 11 and 12), for which the equation being $y = mx^2 + b$. This suggests of uniform acceleration. This shows when there is more increase in percentage diameter stenosis (as high as 75%), the flow gets accelerated.

Influence of length of stenosis: For both 5mm length stenosed Brachial and Femoral arteries, with increased viscosity and for 75% diameter stenotic conditions, there is a steep increase in PSV ratio which makes the curve look like a top-opening parabola, for which the equation being $y = mx^2 + b$.

$= mx^2 + b$ (Figures 5 and 7). This suggests of uniform acceleration. This shows when the high viscous flow is through 75% stenosis of lesser length (as less as 5mm), the flow get accelerated. Whereas, for both 10mm length stenosed Brachial and Femoral arteries, with increased viscosity and for 75% diameter stenotic conditions, there is a steep decrease in PSV ratio which makes the curve look like a side-opening parabola for which the equation being $y^2 = mx + b$ (Figures 6 and 8). This suggests of uniform retardation. This shows when the high viscous flow is through 75% stenosis of more length (as high as 10mm), the flow gets retarded.

Influence of cross-sectional area of the blood vessel

Table 5: Range of PSV ratio with respect to cross-sectional area of the blood vessel (from Tables 1 and 2)

Artery model	Range of PSV ratio under various simulated conditions		
	25% diameter stenosis	50% diameter stenosis	75% diameter stenosis
Brachial artery model	1.1956 to 1.4019	1.3744 to 1.7374	1.7743 to 2.6245
Femoral artery model	1.0361 to 1.179	1.2776 to 2.3566	2.6887 to 5.85

Brachial artery model with a cross-sectional area of 12.57mm^2 showed notable variation in flow dynamics under lesser percentage of stenotic conditions itself but Femoral artery model with a cross-sectional area of 50.29mm^2 showed good variation in flow dynamics only under higher percentage of stenotic conditions. This shows that to make significant variation in flow dynamics, the percentage of stenosis has to be comparable with the size of the vessel. In Brachial artery models, there is a gradual and less steep increase in PSV ratio with respect to % diameter stenosis. Whereas, in Femoral artery models, more steep increase in PSV ratio noted with respect to % diameter stenosis. This shows that with increased cross-sectional area, the flow gets more accelerated.

CONCLUSION

The experimental study using the Doppler blood flow phantom shows the strong impact of viscosity of blood, stenotic conditions and cross-sectional area of the blood vessel on flow dynamics. Various studies have been published to prove the potential role of whole blood viscosity in haemodynamics and to support the need for the assessment of Whole blood viscosity in order to rule out various blood flow related diseased conditions^{10,11}. In a study done by Ajmani RS *et al*, whole blood viscosity measurement with respect to aging and age-related diseases was discussed². In spite of these, the measurement of whole blood viscosity of patients is still not a common practice in clinical laboratories. This study

suggests that the measurement of whole human blood viscosity of the patients also shall be carried out along with other routine blood tests as this will prove to be a predictor or an indicator of blood flow related ailment in patients. The study also suggests that the peripheral artery diseases especially the stenotic conditions shall be noted as a sign of alarm for the upcoming serious morbidity and hence appropriate step has to be taken to avoid or manage the severity of the consequential problem.

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REFERENCES

1. Sudha Pattan, P.M. Venkata Sai, Santhosh Joseph, Aruna P, Ravishankar P: An indigenously fabricated Doppler ultrasound blood flow phantom: International Journal of Recent Trends in Science and Technology, April 2015; 14(3); 587-595.
2. Ralph E. Holsworth, Jr. And Jonathan V.Wright: Blood viscosity - the unifying parameter in Cardiovascular disease risk: Holistic primary care, 18:14; Vol.13, 2012.
3. Ajmani RS *et al*: Hemodynamic changes during aging associated with cerebral blood flow and impaired cognitive function: Neurobiol Aging, 2000 Mar-Apr; 21(2); 257-69.
4. K. Rammnarine *et al*: Validation of a New Blood-Mimicking Fluid for Use in Doppler Flow Test Objects: Ultrasound in Medicine and Biology, Vol. 24. No. 3, 454.
5. Gerhard-Herman *et al*: Guidelines for Noninvasive Vascular Laboratory Testing: A Report from the American Society of Echocardiography and the Society of Vascular Medicine and Biology: Journal of the American Society of Echocardiography, August 2006; 955-972.
6. Ronald O. Bude, Jonathan M. Rubin: Relationship between the Resistive Index and Vascular Compliance and Resistance: Radiology, 1999; 211; 411-417.
7. D H Evans, W W Barrie, M J Asher, S Bentley and P R Bell: The relationship between ultrasonic pulsatility index and proximal arterial stenosis in a canine model: Circulation Research, 1980; 46; 470-475.

8. Tohoku: Exercise related time course of Pulsatility Index in brachial artery following forearm exercise assessed by Doppler ultrasound: *J. Exp. Med.* 2004; 203; 241-252.
9. Hoskins PR, Loupas T, McDicken WN: A comparison between the Doppler signals from human blood and artificial blood used in a flow phantom: *Ultrasound in Medicine and Biology*, 1990; 16 (2); 141-7.
10. G.A.M. Pop, D.J. Duncker, M. Gardien, P. Vranckx, S. Versluis, D. Hasan, and C.J. Slager: The clinical significance of whole blood viscosity in (cardio)vascular medicine: *Neth Heart J.* 2002 Dec; 10(12); 512–516.
11. Elif Hande Ozcan et al: The Forgotten Variable of Shear Stress in Mitral Annular Calcification - Whole Blood Viscosity: *Med Princ Pract* 2015; 24; 444–450.

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