



RESEARCH ARTICLE

QUANTITATIVE ASSESSMENT OF PORTAL VEIN BY COLOUR DOPPLER IMAGING IN PATIENTS WITH VARIOUS LIVER DISEASES IN WEST BENGAL AND REFLECTION OF PATHOPHYSIOLOGICAL HAEMODYNAMICS AND EVALUATION OF PROGNOSIS

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ABSTRACT

We conducted a study in West Bengal in K.P.C. medical college on patients with Liver disease with the help of Duplex Doppler Ultrasound and assessed quantitative and semi quantitative indices of portal vein. We found that it is a safe, noninvasive, inexpensive, excellent investigation of choice in assessing pathophysiological haemodynamics, judging the severity of the disease, effects of several drugs and prediction of prognosis of the disease. Before conducting the study we took permission from the ethical committee of our college.

Key words:

Ultrasound, Portal vein, duplex
doppler, congestion index,
pulsatility index, resistance index.

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INTRODUCTION

Colour Doppler Sonography is the most effective primary investigation of choice in elucidation of Liver pathology. It is a superb, noninvasive inexpensive method and plays a crucial role in diagnosis and management of various Liver diseases. So as per guideline we conducted a Doppler flowmetry study in K.P.C. Medical College over 235 people with taking into account quantitative and semi quantitative data.

MATERIAL AND METHOD

We started our original and honest study after taking permission from our ethical committee over a period of (2012 – 2014) over 235 people. The patients are coming from all over West Bengal like, Kolkata, North and South 24 Parganas, Howrah, Hooghly, Burdwan, Birbhum, Bankura, Midnapore, Nadia, Malda, Murshidabad. After taking consent from all the patients and collecting demographic data like age, sex, religion following questions were asked –

1. For how long they are suffering from this disease.
2. Are they taking any medicines.
3. Any past history of operation or blood transfusion.

4. Any other medical condition associated like – hypertension, diabetes etc.
5. Drug, alcohol addiction, HIV infection.
6. We have taken into account – normal subjects after taking consent from them, to serve as a reference scale.

We studied total 235 people – ^{4, 15, 16, 22}

1. Normal subjects 100 (N)
2. Patients with acute hepatitis 10 (AH)
3. 40 Patients with chronic active hepatitis (CAH)
4. 80 Patients with liver cirrhosis (LC)
5. 5 Patients with Idiopathic portal hypertension (IPH)

METHOD

We studied the patient as per guidelines – [Sabba et al 1995](#).

1. Measure in suspended normal respiration.
2. Longitudinal scan of portal vein.
3. Sample volume at centre of vessel at level of hepatic artery covering 50 percent of vessel diameter.
4. Doppler angle of 55° or less.
5. Pulse repetition frequency (PRF) = 4 KHZ wall filter = 100 Hz

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6. Doppler and B mode tracing recorded simultaneously.
7. PV diameter measures from inner anterior to inner posterior wall.
8. Values result from measuring of 3 consistent measurement. ^{3, 8, 9, 11, 13, 14, 20}

Other semi quantitative assessment were taken into account.

$$\text{Pulsatility Index PI} = \frac{A - B}{A}$$

Mean

A - Peak systolic velocity

B - Peak diastolic velocity

Mean - Mean of velocity in a set interval of time.

$$\text{Resistance Index RI} = \frac{A - B}{A}$$

DISCUSSION

In the study by [F. Moriyasu et al](#) they found –

A positive correlation between congestion index CI and portal venous pressure.

They found congestion index reflects the pathophysiological haemodynamics of portal venous system in portal hypertension. They also conducted a comparative study of congestion index of normal subject and patients with liver disease.

These results were like this

cross sectional area –

- Normal subjects – (0.99 ± 0.28 cm²) (n=85)
- Acute hepatitis - (1.05 ± 0.22 cm²) (n=11)
- Chronic active hepatitis - (1.22 ± 0.41 cm²) (n=42)
- Cirrhosis - (1.49 ± 0.49 cm²) (n=72)
- Idiopathic portal hypertension (1.56 ± 0.45 cm²) (n=11)

Congestion Index –

- Normal – 0.070 ± 0.029 cm x sec
- Acute Hepatitis – 0.071 ± 0.014 cm x sec
- Chronic active hepatitis – 0.119 ± 0.084 cm x sec
- Liver Cirrhosis – 0.171 ± 0.075 cm x sec
- Idiopathic portal hypertension – 0.180 ± 0.107 cm x sec

In our study with colour doppler flowmetry we also found that–

1. Patient with chronic liver disease, cirrhosis and idiopathic portal hypertension have higher value of cross sectional area.

Whereas people with liver cirrhosis have altered value of mean blood flow velocity and blood flow volume.

2. Congestion index is higher in patients with chronic active hepatitis, Liver cirrhosis, Idiopathic portal hypertension.
3. P.I. of superior mesenteric artery is reduced in patients with Liver cirrhosis.
4. P.I. of portal vein is more important indicator for clinical evaluation of patients with right heart failure.
5. Finally the compliance curve of portal vein – i.e portal vein pressure volume – curve is not linear.

Congestive Index –

$$\text{CI} = \frac{\text{Cross sectional area of portal vein}}{\text{Blood flow velocity of portal vein}}$$

$$= \frac{A \times B \times \cos \theta}{4 \times \text{Vdmax}} \div 0.57 \text{ Vdmax (cm X sec)}$$

A= Short axis of portal vein

B=Long axis of portal vein

Vdmax=maximum velocity obtained from Doppler

θ = angle between ultrasound beam and blood vessels.

Coefficient 0.57 is the ratio of mean velocity to maximum velocity obtained. ^(5 – 7, 10, 12, 17 – 21)

RESULTS

Observation table – 1 Cross sectional area of portal vein

1	Normal n=100	0.99 ± 0.22 cm ²
2	Acute Hepatitis n= 10	1.08 ± 0.24 cm ²
3	Chronic Active Hepatitis n= 40	1.32 ± 0.38 cm ²
4	Liver Cirrhosis n= 80	1.55 ± 0.50 cm ²
5	Idiopathic Portal Hypertension n=5	1.59 ± 0.48 cm ²

Table – 2 Mean Blood Flow Velocity

1	Normal n=100	15.5 ± 4.0 cm/sec
2	Acute Hepatitis n= 10	15.1 ± 2.2 cm/sec
3	Chronic Active Hepatitis n= 40	12.5 ± 3.3 cm/sec
4	Liver Cirrhosis n= 80	9.8 ± 2.8 cm/sec
5	Idiopathic Portal Hypertension n=5	11.0 ± 3.5 cm/sec

Table 3 Blood Flow Volume of Portal Vein

1	Normal n=100	900 ± 275 ml/min
2	Acute Hepatitis n= 10	1000 ± 350 ml/min
3	Chronic Active Hepatitis n= 40	850 ± 250 ml/min
4	Liver Cirrhosis n= 80	850 ± 300 ml/min
5	Idiopathic Portal Hypertension n=5	1050 ± 350 ml/min

Table – 4 Congestion Index

1	Normal n=100	0.070 ± 0.028 cm x sec
2	Acute Hepatitis n=10	0.072 ± 0.015 cm x sec
3	Chronic Active Hepatitis n=40	0.122 ± 0.088 cm x sec
4	Liver Cirrhosis n=80	0.180 ± 0.075 cm x sec
5	Idiopathic Portal Hypertension n=5	0.188 ± 0.110 cm x sec

Increased flow volume is compensated by deformity of blood vessel cross-sections from elliptical to round, thereby elevation of portal pressure is minimized.

CONCLUSION

Doppler flowmetry with quantitative data, are excellent, effective, noninvasive, inexpensive method in assessing severity of hepatic disease, evaluating the role and effectiveness of drugs and predicting haemorrhagic risk and elucidating the prognosis.

This study also plays special role in disease affecting portal tract, like Budd-Chiari syndrome, schistosomiasis, malaria, drug toxicity etc.

In post-operative case (like liver transplantation) on day 1,3,5 and 7 and weekly there after, the quantitative flow- study plays excellent role.

References

1. Bolondi L, Gandolfi L, Arienti V, Caletti GC, Corcioni E, Gasbarrini G, *et al.* Ultrasonography in the diagnosis of portal hypertension: Diminished response of portal vessels to respiration. *Radiology* 1982;142:167-72.
2. Vilgrain V, Lebrech D, Menu Y, Scherre A, Nahum H. Comparison between ultrasonographic signs and the degree of portal hypertension in patients with cirrhosis. *Gastrointest Radiol* 1990;15:218-22 [PUBMED]
3. Ralls PW. Color Doppler sonography of the hepatic artery and portal venous system. *AJR Am J Roentgenol* 1990;155:517-25.
4. Moriyasu F, Nishida O, Ban N, Nakamura T, Sakai M, Miyake T, *et al.* "Congestion index" of the portal vein. *AJR Am J Roentgenol* 1986;146:735-9
5. Haag K, Rossle M, Ochs A, Huber M, Siegerstetter V, Olschewski M, *et al.* Correlation of duplex sonography findings and portal pressure in 375 patients with portal hypertension. *AJR Am J Roentgenol* 1999;172:631-5.
6. Kok T, van der Jagt EJ, Haagsma EB, Bijleveld CM, Jansen PL, Boeve WJ. The value of Doppler ultrasound in cirrhosis and portal hypertension, *Scand J Gastroenterol Suppl* 1999;230:82-8 [PUBMED]
7. Zoli M, Dondi C, Marchesini G, Cordiani MR, Melli A, Pisi E. Splenic vein measurements in patients with liver cirrhosis: A case-control study. *J Ultrasound Med* 1985;4:641-6 [PUBMED]
8. Subramanyam BR, Balthazar EJ, Madamba MR, Raghavendra BN, Horii SC, Lefleur RS. Sonography of portosystemic venous collaterals in portal hypertension. *Radiology* 1983;146:161-6. [PUBMED]
9. Kedar RP, Merchant SA, Malde HH, Patel VH. Multiple reflective channels in the spleen: A sonographic sign of portal hypertension. *Abdom Imaging* 1994;19:453-8 [PUBMED]
10. Dodd GD 3rd, Memel DS, Baron RL, Eichner L, Santiguada LA. Portal vein thrombosis in patients with cirrhosis: Does sonographic detection of intrathrombus flow allow differentiation of benign and malignant thrombus? *AJR Am J Roentgenol* 1995;165:573-7. [PUBMED]
11. Richter J, Zwingenberger K, Ali QM, Lima Wde M, Dacal AR, de Siqueira GV, *et al.* Hepatosplenic schistosomiasis: Comparison of sonographic findings in Brazilian and Sudanese patients-correlation of sonographic findings with clinical symptoms. *Radiology* 1992;184:711-6.
12. Forsberg L, Holmin T. Pulsed Doppler and B-mode ultrasound features of interposition meso-caval and porta-caval shunts. *Acta Radiol [Diagn] (Stockh)* 1983;353-357
13. Wetzner SM, Kiser LC, Bezreh JS. Duplex ultrasound imaging: vascular applications. *Radiology* 1984;150:507-514
14. Gill RW. Pulsed Doppler with B-mode imaging for quantitative blood flow measurement. *Ultrasound Med Biol* 1979;5:223-235
15. Moriyasu F, Ban N, Nishida O, *et al.* Quantitative measurement of portal blood flow in patients with chronic liver diseases using an ultrasonic duplex system consisting of a pulsed Doppler flowmeter and B-mode electroscanner. *Gastroenterol Jpn* 1984;19:529 – 536
16. Moriyasu F, Nishida O, Ban N *et al.* Measurement of portal vascular resistance in patients with portal hypertension, *Gastroenterology* (in press)
17. Subramanyam BR Balthazar EJ, Raghavendra BN, Lefleur RS. Sonographic evaluation of patients with portal hypertension. *Am J Gastroenterol* 1983;78:369-373
18. Niederau C, Sonnenberg A, Muller J, Erckenbrecht JF, Scholten T, Fritsch WP. Sonographic measurements of the normal liver, spleen, pancreas and portal vein. *Radiology* 1983;149:537-540
19. Lafortune M, Marlean D, Breton G, Viallet A, Lavoie P, Huet P. Portal venous system measurements in portal hypertension. *Radiology* 1984;151:27-30
20. Ueda H, Kitani K, Kameda H, *et al.* Splenic blood flow in idiopathic portal hypertension in Japan measured by Kr clearance method. *Acta Hepatosplenol* 1971;18:28-40
21. Groszmann R, Kotelanski B, Cohn JN, Khatri JM. Quantitation of port systemic shunting from the splenic and mesenteric bed in alcoholic liver disease. *Am J Med* 1972;53:715-722
22. Sherlock DS. The portal venous system and portal hypertension. In: *Diseases of the liver and biliary system*, 6th ed. Oxford: Blackwell Scientific, 1982:135-176.
