

RAPID COMMUNICATION

Portal vein pulsatility index is a more important indicator than congestion index in the clinical evaluation of right heart function

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CONCLUSION: Our data show that RI is a more significant indicator than CI in the clinical evaluation of high RA ≥ 10 mmHg, whereas CI is better than PI in the assessment of left heart function.

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Key words: Portal blood flow; Heart failure; Ultrasonic Doppler; Congestion index; Portal vein pulsatility index

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Abstract

AIM: To study the changes of portal blood flow in congestive heart failure.

METHODS: We studied the congestion index (CI) and portal vein pulsatility index (PI) in patients with varied degrees of congestive heart failure using ultrasonic Doppler. Ten patients with mean right atrial pressure (RA) < 10 mmHg were classified as group 1 and the remaining 10 patients with RA ≥ 10 mmHg as group 2.

RESULTS: There were no difference on cardiac index (HI, $P=0.28$), aortic pressure (AO, $P=0.78$), left ventricular end-diastolic pressure (LVED, $P=0.06$), maximum portal blood velocity (V_{max} , $P=0.17$), mean portal blood velocity (V_{mean} , $P=0.15$) and portal blood flow volume (PBF, $P=0.95$) between the two groups. Group 2 patients had higher pulmonary wedge pressure (PW, 29.9 ± 9.3 mmHg vs 14.6 ± 7.3 mmHg, $P=0.002$), pulmonary arterial pressure (PA, 46.3 ± 13.2 mmHg vs 25.0 ± 8.2 mmHg, $P=0.004$), RA (17.5 ± 5.7 mmHg vs 4.7 ± 2.4 mmHg, $P < 0.001$), right ventricular end-diastolic pressure (RVED, 18.3 ± 5.6 mmHg vs 6.4 ± 2.7 mmHg, $P < 0.001$), CI (8.7 ± 2.4 vs 5.8 ± 1.2 , $P=0.03$), and PI ($87.8 \pm 32.3\%$ vs $27.0 \pm 7.4\%$, $P < 0.001$) than Group 1. CI was correlated with PI ($P < 0.001$), PW ($P < 0.001$), PA ($P < 0.001$), RA ($P=0.043$), RVED ($P=0.005$), HI ($P < 0.001$), AO ($P < 0.001$), CO ($P < 0.001$), LVED ($P < 0.001$), V_{max} ($P < 0.001$), V_{mean} ($P < 0.001$), cross-sectional area of the main portal vein ($P < 0.001$) and PBF ($P < 0.001$). CI could be as high as 8.3 in patients with RA < 10 mmHg and as low as 5.9 in those with RA ≥ 10 mmHg.

INTRODUCTION

Congestive heart failure increases the pressure in the inferior vena cava and hepatic veins^[1-3]. Ultrasonic Doppler is a safe and non-invasive method in the clinical evaluation of portal blood flow and portal hypertension^[4-9]. Portal vein pulsatility index (PI) is calculated by the percentage of peak-to-peak maximum portal vein velocities^[10,11]. In our earlier study^[10], patients with right heart failure developed transient reduced, stagnant, or hepatofugal portal blood flow with increased PI. However, the change of portal flow pattern and PI did not correlate with left heart function.

The congestion index (CI) has been used to assess the pathophysiological hemodynamics of portal venous system in different forms of liver diseases^[12-14]. The correlation between CI and PI and the role of CI on right heart function remain uncertain. Therefore, we have studied the changes of portal blood flow in patients with different degrees of heart failure using non-invasive ultrasonic Doppler^[15,16].

MATERIALS AND METHODS

We studied the portal hemodynamic profiles in 20 patients (9 males, 11 females, mean age: 49 ± 13 years) who received cardiac and Swan-Ganz catheterizations for cardiovascular disorders (16 rheumatic heart disease, 4 atherosclerotic heart disease) to compare with 20 healthy volunteers. All

Table 1 Clinical and biochemical data in patients with heart failure (mean \pm SD)

	Controls	RA < 10 mmHg	RA \geq 10 mmHg
Gender (M/F)	10/10	4/6	5/5
Age (yr)	46 \pm 12	50 \pm 13	47 \pm 19
Total protein (g/dL)	7.5 \pm 0.6	7.1 \pm 0.8	6.9 \pm 1.1
Albumin (g/dL)	4.3 \pm 0.2	3.9 \pm 0.7	3.7 \pm 0.5
Total serum bilirubin (mg/dL)	0.9 \pm 0.4	1.3 \pm 0.8	1.4 \pm 0.8
AST (U/L)	21 \pm 6	31 \pm 11	59 \pm 29 ^{1b}
ALT (U/L)	24 \pm 6	23 \pm 8	35 \pm 20
Prolonged prothrombin time (s)	-	1.1 \pm 0.9	1.4 \pm 0.8

RA, right atrial pressure; ¹ $P=0.009$ vs patients with RA < 10 mmHg and ² $P<0.001$ vs controls.

Table 2 Cardiac profiles in patients with congestive heart failure (mean \pm SD)

	RA < 10 mmHg	RA \geq 10 mmHg	<i>P</i>
HI (L/min/m ²)	3.0 \pm 0.9	2.4 \pm 0.4	0.28
AO (mmHg)	89.0 \pm 9.6	87.3 \pm 12.8	0.78
LVED (mmHg)	12.2 \pm 6.7	22.1 \pm 10.9	0.06
PW (mmHg)	14.6 \pm 5.6	29.1 \pm 7.7	0.002
PA (mmHg)	25.0 \pm 6.8	42.4 \pm 12.0	0.004
RA (mmHg)	4.7 \pm 2.1	16.8 \pm 4.9	<0.001
RVED (mmHg)	6.4 \pm 2.1	17.8 \pm 4.4	<0.001

RA, right atrial pressure; HI, cardiac index; LVED, left ventricular end-diastolic pressure; AO, mean aortic pressure; PW, pulmonary wedge pressure; PA, mean pulmonary arterial pressure; RA, mean right atrial pressure; RVED, right ventricular end-diastolic pressure.

patients had medications affecting the hemodynamics such as isosorbide dinitrate and furosemide, and their systemic blood pressure and body weight were measured to be constant for more than 48 h prior to the study. Patients with fever, infection, and shock were excluded. All patients had no history of liver disease, alcoholism or other metabolic disorders. None of the patients received transfusion, inotropic agents or dopamine. All patients had an abdominal sonography to exclude chronic liver disease or splenomegaly. Patients with severe orthopnea were excluded if they were not able to remain in the supine position for the study of ultrasonic Doppler.

Cardiac profiles including cardiac index (HI), left ventricular end-diastolic pressure (LVED), mean aortic pressure (AO), pulmonary wedge pressure (PW), mean pulmonary arterial pressure (PA), mean right atrial pressure (RA), right ventricular end-diastolic pressure (RVED) were recorded during the cardiac and Swan-Ganz catheterizations. Ten patients with RA < 10 mmHg (range: 1-7 mmHg) and without right heart failure were classified as Group 1. The remaining 10 patients with right heart failure and RA \geq 10 mmHg (range: 10-28 mmHg) were classified as Group 2.

The portal profiles were assessed using an ultrasonic Doppler composed of a real-time mechanical sector scanner and a 3.5 MHz pulsed Doppler flowmetry (Aloka Echo Camera, Model SSD-1700, Tokyo) within 12 h of cardiac catheterization. After more than 8 h of fasting, portal pro-

files were measured in the supine position for more than 30 min. Portal blood flow was measured from the main portal vein at a site just entering or immediately after entering the liver with the patient in expiratory apnea. The flow angle formed by the directions of ultrasonic beam and the portal blood flow below 55 degree was corrected to minimize the variation caused by the angle of insonation. The Doppler signal could be viewed on the screen and heard through a build-in speaker. Portal blood flow was measured by the same physician (SY) to avoid interobserver variation^[17].

For each measurement, at least three reproducible spectral patterns were recorded for calculating the mean maximum portal blood velocity (V_{max}) over a period of 3-4 s to ensure accuracy. Mean portal blood velocity (V_{mean}) was calculated by the equation " $V_{mean}=0.57 \times V_{max}$ " as described by Moriyasu *et al.*^[18]. Cross-sectional area (area, cm²) was also recorded at the site of main portal vein where portal blood velocity was measured. The direction of portal blood velocity, antegrade or retrograde, was also measured. Positive velocity indicates the blood flow towards the transducer and vice versa. Portal blood flow volume (PBF, mL/min) was obtained by the equation " $PBF=area \times V_{mean} \times 60$ "^[17,18]. PI was calculated by the equation " $PI=(\text{maximum}-\text{minimum})/\text{maximum frequency shift}$ "^[6,15,17]. The waveforms were classified as continuous ($PI \leq 40\%$), decreased ($PI 41-99\%$), stagnant ($PI = 100\%$), or retrograde ($PI > 100\%$)^[10,19]. CI was calculated by the equation " $CI=(area/V_{mean}) \times 100$ "^[12].

The study protocol was reviewed and approved by the Institutional Review Committee under the guidelines of the 1975 Declaration of Helsinki. Statistical analysis was performed using Student's *t*-test and simple linear regression as appropriate.

RESULTS

The biochemical data of the 20 patients (Table 1) showed total protein 7.0 \pm 0.8 g/dL, albumin 3.8 \pm 0.5 g/dL, total bilirubin 1.3 \pm 0.6 mg/dL, AST 49.5 \pm 23.4 IU/L, ALT 28.7 \pm 10.4 IU/L, and prolonged prothrombin time 1.2 \pm 0.9 s (normal < 3 s). All controls had normal blood chemistries. Gender ($P=0.11$), age ($P=0.61$), total protein ($P=0.85$), albumin ($P=0.62$), total bilirubin ($P=0.83$), ALT ($P=0.15$) and prolonged prothrombin time ($P=0.19$) were not different between those with RA < 10 mmHg and \geq 10 mmHg. Patients with RA \geq 10 mmHg had higher serum AST activities ($P=0.009$), which were related to ischemic hepatitis.

HI (3.0 \pm 0.9 L/min/m²; range: 1.6-5.3 L/min/m² vs 2.4 \pm 0.4 L/min/m²; range: 1.7-2.9 L/min/m²; $P=0.28$), AO (89.0 \pm 9.6 mmHg; range: 85-100 mmHg vs 87.3 \pm 12.8 mmHg; range: 65-115 mmHg; $P=0.78$), and LVED (12.2 \pm 6.7; range: 4-34 mmHg vs 22.1 \pm 10.9 mmHg; range: 10-40 mmHg; $P=0.06$) were not statistically different between Groups 1 and 2 (Table 2).

For all Group 1 patients, the values of PW (mean: 14.6 \pm 5.6 mmHg; range: 5-28 mmHg), PA (mean: 25.0 \pm 6.8 mmHg; range: 16-38 mmHg), RA (mean: 4.7 \pm 2.1 mmHg; range: 1-7 mmHg), and RVED (mean: 6.4 \pm 2.1 mmHg; range: 2-11 mmHg) were within the nor-

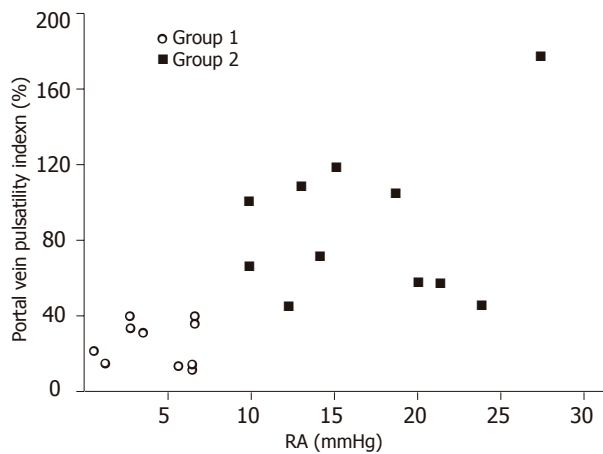


Figure 1 Portal vein pulsatility index of patients with right atrial pressure < 10 mmHg (Group 1) and ≥ 10 mmHg (Group 2).

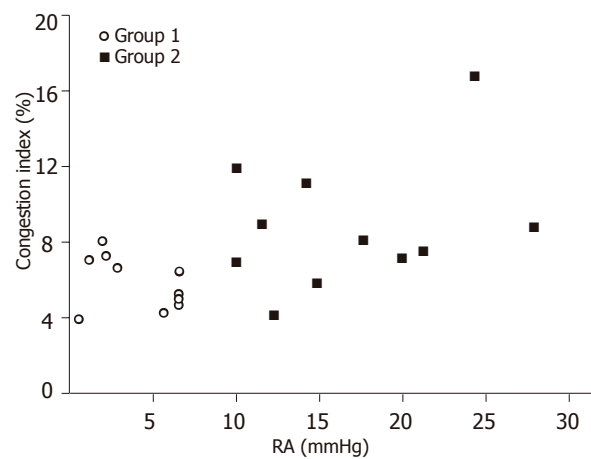


Figure 2 Congestion index of patients with right atrial pressure (RA) < 10 mmHg (Group 1) and ≥ 10 mmHg (Group 2).

Table 3 Portal profiles in patients with congestive heart failure (mean \pm SD)

	Controls (n = 20)	RA ≤ 10 mmHg (n = 10)	RA > 10 mmHg (n = 10)
V_{\max} (cm/s)	20.1 \pm 3.1	24.5 \pm 3.9	21.1 \pm 4.8
V_{mean} (cm/s)	11.2 \pm 1.9	14.0 \pm 2.3	12.0 \pm 2.7
Area (cm ²)	1.01 \pm 0.20	0.80 \pm 0.13	0.96 \pm 0.13
PBF (mL/min)	685 \pm 136	678 \pm 172	672 \pm 162
PI (%)	23.3 \pm 6.3	27.0 \pm 7.4	87.8 \pm 32.3 ^b
Congestion index	5.3 \pm 1.2	5.8 \pm 1.2	8.7 \pm 2.4 ¹

RA, right atrial pressure; V_{\max} , maximum portal velocity; V_{mean} , mean portal velocity; PBF, portal blood flow; PI, portal vein pulsatility index. ^b $P < 0.001$, ¹ $P = 0.03$ vs controls and RA ≤ 10 mmHg.

mal limits.

All Group 2 patients had higher PW (mean: 29.1 \pm 7.7 mmHg; range: 13-40 mmHg; $P = 0.002$), PA (mean: 42.4 \pm 12.0 mmHg; range: 25-65 mmHg; $P = 0.004$), RA (mean: 16.8 \pm 4.9 mmHg; range: 10-28 mmHg; $P < 0.001$), and RVED (mean: 17.8 \pm 4.4 mmHg; range: 9-26 mmHg; $P < 0.001$) than Group 1 patients.

The healthy controls had V_{\max} 20.1 \pm 3.1 cm/s, V_{mean} 11.2 \pm 1.9 cm/s, area 1.01 \pm 0.20 cm², PBF 685 \pm 136 mL/min, PI 23.3 \pm 6.3%, and CI 5.3 \pm 1.2. The mean values of V_{\max} (24.5 \pm 3.9 cm/s; range 17-33 cm/s vs 21.1 \pm 4.8 cm/s; range 14-33 cm/s; $P = 0.17$), V_{mean} (14.0 \pm 2.3 cm/s; range: 9.7-18.8 cm/s vs 12.0 \pm 2.7 cm/s; range: 8.6-18.8 mmHg; $P = 0.15$) and PBF (678 \pm 172 mL/min; range: 373-1 120 mL/min vs 672 \pm 162 mL/min; range: 432-922 mL/min; $P = 0.95$) between Groups 1 and 2 did not show any statistical difference (Table 3). Group 2 patients had a larger area of portal vein than that of Group 1 (0.80 \pm 0.13 cm²; range: 0.64-1.13 cm² vs 0.96 \pm 0.13 cm²; range: 0.79-1.33 cm²; $P = 0.04$).

All the 10 patients in Group 1 had a continuous antegrade portal flow with a mean PI 27.0 \pm 7.4% (range: 17-40%) (Figure 1). The mean PI of the 10 patients in Group 2 was 87.8 \pm 32.2% (range: 43-194%). In Group 2, all the patients had a PI > 40%. Six of them had transient reduced portal blood flow, one had stagnant flow, and

three had hepatofugal flow.

Group 2 patients (mean: 8.7 \pm 2.4, range: 5.9-16.7) had a higher CI than that of Group 1 patients (mean: 5.8 \pm 1.2, range: 3.9-8.3; $P = 0.03$). Although Group 2 had a higher mean CI than Group 1, the CI could be as low as 5.9 in Group 2 and as high as 8.3 in Group 1 (Figure 2).

Using linear regression, CI showed a good correlation with PI ($P < 0.001$), PW ($P < 0.001$), PA ($P < 0.001$), RA ($P = 0.043$), RVED ($P = 0.005$), HI ($P < 0.001$), AO ($P < 0.001$), CO ($P < 0.001$), LVED ($P < 0.001$), V_{\max} ($P < 0.001$), V_{mean} ($P < 0.001$), area ($P < 0.001$) and PBF ($P < 0.001$).

DISCUSSION

It is well known that the passive "backward" congested liver develops into hepatomegaly, synchronous pulsation, engorged and dilated terminal hepatic veins, atrophy of hepatocytes and eventually cardiac cirrhosis. The high hepatic vein pressure can transmit through the liver to cause post-sinusoidal portal hypertension, cardiac ascites and change of portal vein flow patterns^[12,13]. Therefore, the changes of portal flow may help the assessment of heart function.

Prolonged right heart failure may result in atrophy of hepatocytes and eventually cardiac cirrhosis^[3]. In the present study, we have strived to exclude those patients with chronic liver disease. The abdominal sonographies showed no splenomegaly or coarse liver echogenicity and the peripheral blood showed no abnormal reduction of leukocyte, hemoglobin or platelet account, which were common in cirrhosis. Furthermore, the portal flow pattern did not show reduced fluctuation, which was common in cirrhosis with portal hypertension^[4]. Our patients were not likely to develop obvious cardiac cirrhosis.

In the present study, all patients with RA ≥ 10 mmHg had a PI > 40% and all patients with RA < 10 had a PI < 40% or less. The findings were consistent with our prior study^[10] that PI showed a good correlation with PW, PA, RA, and RVED. The waveform changes of portal blood flow correlate well with right heart function, and the measurement of PI change is a simple and non-

invasive method to identify right heart failure^[10]. Our data also demonstrated that PI had no any correlation with HI, AO, CO, LVED, V_{max} , V_{mean} and PBF. Furthermore, the waveform changes of portal blood flow correlated well with right heart function; and the PI is helpful for the diagnosis of stagnant or hepatofugal portal blood flow but not by the CI^[10]. Therefore, CI is better than PI in the assessment of left heart function.

In addition to the assessment of left heart function, the CI correlated with all PBF, V_{max} , V_{mean} , area, PI, HI, PW, PA, RA, AO, CO, LVED, and RVED. These results suggest that CI also correlates well with right heart profiles. Our findings were consistent with earlier studies^[12,20,21]. However, the CI values could be as high as 8.3 in patients with RA < 10 mmHg and as low as 5.9 in those with RA \geq 10 mmHg. If the CI value is between 5.9 and 8.3, it is difficult to predict whether or not the RA values \geq 10 mmHg. Therefore, RI is a more significant indicator than CI in the clinical evaluation of high RA \geq 10 mmHg.

The occurrence of congestive liver is not uncommon in patients with congestive heart failure. In addition to the occurrence of congestive hepatomegaly and dilatation of inferior vena cava and hepatic veins during abdominal sonography, the measurement of both CI and PI is helpful for the indirect non-invasive evaluation of cardiac function.

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