



The Most Useful Method to Evaluate the Volume Status of Critical Patients in The Emergency and Intensive Care Units: Point of Care Ultrasound

Acil Servis ve Yoğun Bakım Ünitelerindeki Kritik Hastaların Volüm Durumunu Değerlendirmede En Kullanışlı Yöntem: Yatak Başı Ultrasonu

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Abstract

Background: Accurate and rapid assessment of intravascular volume status of the patients in emergency services and intensive care units at diagnosis, treatment and follow-up stages is crucial yet rather difficult. The purpose of hemodynamic monitoring is to determine cardiovascular insufficiency and to provide the most suitable treatment for unstable patients in critical condition.

Aim: The study aims to compare vena cava inferior diameter, vena cava inferior- collapsibility index (for spontaneously breathing patients) and vena cava inferior- distensibility index (for patients breathing on mechanical ventilation support) measurement by ultrasonography to central venous pressure measurement by placing invasive catheter for assessment of the intravascular volume status and making an accurate volume replacement in emergency service and intensive care units and to determine the correlation between them.

Material and Method: The study was carried out prospectively on the patients above the age of 18 who applied to the emergency service clinic between the dates of 01.06.2014 and 01.04.2015 or who stayed in the emergency intensive care unit between these dates. Measurements were taken from vena cava inferior in both the inspirium and expirium phases by using M mode and they were recorded in millimeter. Simultaneous central venous pressure measurements were performed on the patients by using manometric devices and the results were recorded in cm H₂O.

Results: 43.3% of the patients were female (n: 26) and 56.7% were male (n: 34), and the mean age is 70.58±14.86. The study found high degree of positive correlation between central venous pressure and vena cava inferior diameters and high degree of negative correlation between vena cava inferior- collapsibility index. The study also found that there is a high degree of negative correlation between vena cava inferior- distensibility index and central venous pressure in patients receiving mechanical ventilatory support.

Conclusion: Measurement of respiratory variation in vena cava inferior diameter by using ultrasonography is a quick, reliable, easily applicable, cost-efficient and non-invasive method in critical patients receiving mechanical ventilatory support or have spontaneous respiration in emergency services and intensive care units and it can be useful in assessing the volume status and estimating central venous pressure.

Keywords: Central venous pressure, vena cava inferior, ultrasonography, volume status, collapsibility

Öz

Giriş: Acil servis ve yoğun bakımdaki hastaların; tanı, tedavi ve takibinde intravasküler volüm durumunun, doğru ve hızlı şekilde tespiti oldukça önemli ve bir o kadar da zordur. Hemodinamik izlemin amacı kardiyovasküler yetmezliği belirlemek ve stabil olmayan kritik düzeydeki hastalara (septik şok, hipovolemi, kardiyojenik şok v.b.) en uygun tedaviyi sağlamaktır.

Amaç: Bu çalışmada amacımız; acil servis ve yoğun bakım ünitesinde, intravasküler volüm durumunu değerlendirmede ve doğru volüm replasmanına yönelmede ultrasonografi ile vena cava inferior çapı, vena cava inferior - kollapsibilite indeksi (spontan solunumu olan hastalar için) ve vena cava inferior distensibilite indeksi (mekanik ventilasyon desteğinde soluyan hastalar için) ölçümünün invaziv kateter yerleştirilerek yapılan santral venöz basınç değeri ile karşılaştırılması ve arasındaki ilişkinin saptanmasıdır.

Gereç ve Yöntem: Bu çalışma acil tıp kliniğine 01.06.2014-01.04.2015 tarihleri arasında başvuran veya bu tarihler arasında acil yoğun bakım ünitesinde yatmakta olan, herhangi bir endikasyon ile santral venöz kateter takılan 18 yaş üstü hastalar üzerinde prospektif olarak yürütüldü. Vena cava inferior ultrasonografik M mod kullanılarak, hem inspiryum hem de ekspiryum fazında ölçümler alındı ve milimetre cinsinden kaydedildi. Hastalardan eş zamanlı santral venöz basınç ölçümü monometrik cihazlar ile yapıldı ve sonuçlar cm H₂O cinsinden kayıt altına alındı.

Bulgular: Hastaların %43.3'ü kadın (n:26), %56.7'si erkek (n:34) olup, yaş ortalamaları 70.58±14.86 yıl idi. Santral venöz basınç ile vena cava inferior çapları arasında pozitif yönde, vena cava inferior- kollapsibilite indeksi arasında negatif yönde yüksek derecede korelasyon tespit edildi. Çalışmamızda ayrıca mekanik ventilatör ile solunumu sağlanan hastalarda da vena cava inferior- distensibilite indeksi ve santral venöz basınç arasında negatif yönde yüksek derecede korelasyon olduğu saptandı.

Sonuç: Acil servis ve yoğun bakım ünitelerinde mekanik ventile ya da spontan solunuma sahip olan kritik hastalarda; hızlı, güvenilir, kolay uygulanabilir, maliyeti düşük ve noninvaziv bir yöntem olan ultrasonografi ile vena cava inferior çapındaki respiratuar değişkenlik ölçümü; volüm durumunu değerlendirmede ve santral venöz basıncı tahmin etmede kullanılabilir.

Anahtar Kelimeler: Santral venöz basınç, vena cava inferior, ultrasonografi, volüm durumu, kollapsibilite



INTRODUCTION

Accurate and rapid assessment of intravascular volume status of the patients in emergency services and intensive care units at diagnosis, treatment and follow-up stages is crucial yet pretty difficult. The purpose of hemodynamic monitoring is to determine cardiovascular insufficiency and to provide the most suitable treatment for unstable patients in critical condition (septic shock, hypovolemia, cardiogenic shock, etc.). Studies show that either of hypovolemia or volume overload may lead to serious clinical consequences including prolonged mechanical ventilation, higher mortality rates, renal dysfunction and disruption in oxygenation.^[1] Rivers et al. state that decrease in organ damages and increase in survival rates can be achieved in patients with serious sepsis and septic shock through target-specific treatment protocols including aggressive volume treatment aimed at central venous pressure (CVP) and physiological variables.^[2]

Invasive and non-invasive methods are employed in order to determine the volume requirement in patients who applied to the emergency service. In clinical practice, the non-invasive methods such as physical examination (mental state, skin turgor pressure and skin drying, capillary refill time, mucous membrane hydration, temperature of the extremities, peripheral pulse palpation, heart rate and blood pressure, their orthostatic changes, urine volume), chest x-ray and laboratory parameters, and the invasive methods such as monitorization of cardiac output or central venous pressure are used in determining intravascular volume and directing the volume treatment.^[3]

Emergency ultrasonography is also defined as Point of Care Ultrasound (POCUS). It is an imaging method that can be used by the physicians working in emergency services as well as radiologists and it is modified, limited to or focused on emergency conditions. After 1999, diagnostic ultrasound (USG) came into use in many hospitals providing emergency medicine education in USA, and more than 70% of these hospitals included USG training in their education program until 2001.^[4,5] The first POCUS study was carried out by emergency physicians upon clinical use of echocardiography in an emergency service in 1988.^[6] In consequence of the studies which indicate that POCUS can be used for hemoperitoneum and hemopericardium in particular in trauma patients, focused abdominal sonography for trauma (FAST) came out.^[7] POCUS must be made in life-threatening cases such as abdominal aortic aneurysm, thoracoabdominal trauma, traumatic hemoperitoneum, pericardial tamponade, massive pulmonary embolism, tension pneumothorax, ruptured aortic aneurysm, cardiogenic shock and hypovolemia.^[8]

In people with spontaneous breathing, intrathoracic pressure decreases during inspiration, the pressure difference between the right atrium and vena cava inferior (VCI) increases, blood flow accelerates and causes VCI-diameter collapse. During expiration, the intrathoracic pressure rises again and the

pressure difference between the right atrium and vena cava inferior decreases, leading to an increase in the diameter of the vena cava inferior. This respiratory variability in VCI diameter is known as the caval index (collapsibility index= $VCI_{max}-VCI_{min} / VCI_{max}$) and is considered to reflect the intravascular volume state.^[9] In mechanically ventilated patients, the increase in intrathoracic pressure during inspiration decreases the venous return gradient and the diameter of the vena cava inferior increases while its diameter decreases on expiration. This variability in the diameter of the vena cava inferior during mechanical ventilation (distensibility index= $VCI_{max}-VCI_{min} / VCI_{min}$) indicates the patient's response to fluid therapy.^[10]

In differential diagnosis of hypotensive patient, assessment of VCI collapsibility and distensibility is crucial in providing quick and proper treatment. It is important to decide if urgent volume replacement, vasopressor support, pericardiocentesis, thoracostomy, thrombolytic treatment, surgical examination or a combination of them is necessary for a hypotensive patient. The decision can be made by using POCUS in a fast and cheap way.^[9] Assessment of vena cava inferior diameter measurement, collapsibility and distensibility by using POCUS helps making the right volume replacement. It enables to prevent such complications as pulmonary edema, cerebral edema, electrolyte disturbances and dilutional coagulopathy that are caused by volume over-replacement.^[11,12]

The purpose of this study is to compare VCI diameter, vena cava inferior- collapsibility index (VCI-CI) and vena cava inferior- distensibility index (VCI-DI) measurement by POCUS to CVP value obtained by inserting invasive catheter in assessing the volume status and making the right volume replacement in emergency service and intensive care unit and to find the correlation between them. CVP measurement has lost popularity over time. However; being noninvasive, easy, cheap and repeatable makes POCUS indispensable for detecting and monitoring the volume status of critical patients in the emergency room and intensive care units.^[9] One aim of this study is to encourage emergency medicine and intensive care clinicians to use POCUS more frequently.

MATERIAL AND METHOD

The study was carried out prospectively on the patients above the age of 18 who applied to the Emergency Service Clinic between the dates 01.06.2014-01.04.2015 or stayed in the emergency intensive care unit between these dates, and to whom central venous catheter was inserted for any indication upon obtaining the approval of Ondokuz Mayıs University Medical Faculty Clinical Research Ethics Committee dated 30.05.2015 (OMU CREC decree no: 2014/675). A total of 60 people, 26 women and 34 men, were included in this study. The patients were included in the study after necessary and adequate information on the study was given to the conscious patients themselves and the immediate relatives of the unconscious patients and their written consent was taken.

The patients with cor pulmonale, pulmonary hypertension, severe tricuspid insufficiency, severe left heart failure (EF< 40%), cardiac tamponade, and tension pneumothorax and those who are pregnant were excluded from the study. The patients from whom image cannot be taken due to intense gas superposition and obesity were also excluded from the study.

A data collection form was prepared in order to standardize data collection in the study. The data recorded on the form are as follows: name-surname, age, gender, systolic and diastolic tension arterial values, average arterial blood pressure, heart rate, respiratory rate of the patients, their admission diagnosis in application or emergency service intensive care unit, laboratory results (full blood count, electrolyte, kidney function tests, blood gas values) for CVP monitoring of the patients, respiration of the patients (spontaneous or mechanical ventilation), CVP measurement catheter type, CVP measurement values, VCI diameter measured by POCUS in expiration and inspiration, VCI-CI values, VCI-DI values and three-month survival rates. VCI-CI and VCI-DI were calculated using the formula below.^[9,10]

$VCI-CI = (\text{maximum VCI diameter} - \text{minimum VCI diameter}) / \text{maximum VCI diameter} \times 100$

$VCI-DI = (\text{maximum VCI diameter} - \text{minimum VCI diameter}) / \text{minimum VCI diameter} \times 100$

Ultrasonographic imaging was performed at the bedside by the clinician certified for basic and advanced emergency ultrasonography by using Philips HD3 ultrasonography device according to american echocardiography association guidelines. Measurements; It was performed by only one clinician (Meltem Ince) in order not to affect the objectivity of the study. All measurements were made in supine position by a 3,5 MHz sector probe. Vena cava inferior and aorta were detected by transverse examination using 3,5 MHz probe. Then, the probe was turned to the longitudinal plane on the vena cava inferior. On this plane, the heart was detected by cranial angulation first. Then, the junction point of hepatic veins with vena cava inferior was found by reducing the angulation. In the meantime, the liver was used as an acoustic window. In order to obtain optimal and standardized measurements in this position, measurements were taken in both inspiration and expiration phases by using M mode and directing the probe to 20 mm distal to the junction point of hepatic veins and vena cava inferior and they were recorded in millimeter (**Figure 1a, 1b**). Simultaneous vital findings were obtained from the patients and simultaneous CVP measurements were made. CVP measurement was made on the reference point on the midaxillary line on a level with the fourth costal cartilage assuming the right atrium level as reference (zero) level. CVP measurement was made by trained emergency service and intensive care nurse by manometric devices and CVP results were recorded in cm H₂O. Ultrasonographic VCI measurement was made by the ultrasound operator without knowing invasive CVP value of the patients measured centrally

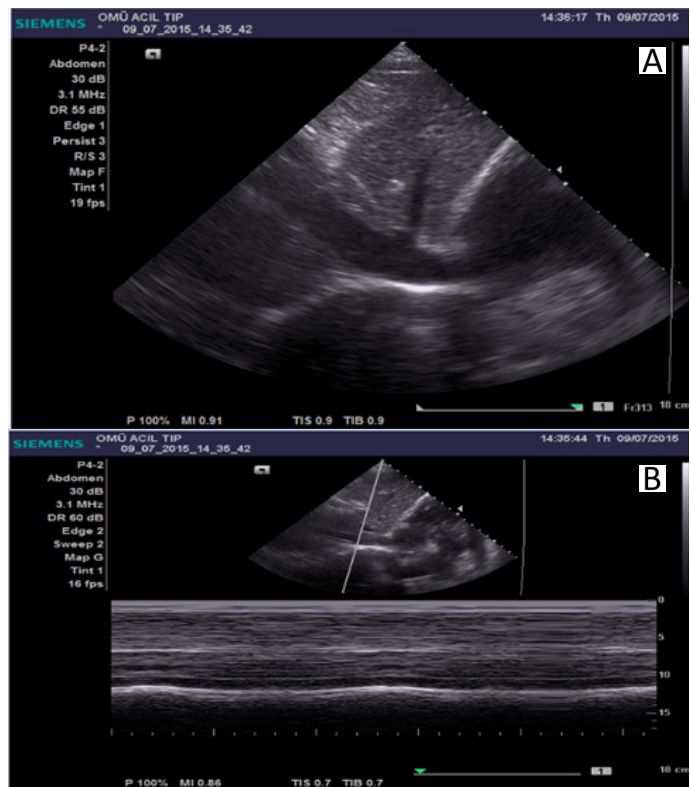


Figure 1. a) View of VCI in B mode in longitudinal section **b)** View of VCI in M mode in longitudinal section

The study data were uploaded to computer and assessed by means of SPSS (Statistical Package For Social Sciences For Windows v.18.0 SPSS Inc. Chicago, IL). Shapiro-Wilk test was made for test of normality of the data. In the table; the fact that the values on Assymp Sig. (significance) line are greater than 0.05 which is assumed as the limit value in statistical significance calculations indicates that distributions of the factors examined are normal, and the fact that they are smaller indicates that there isn't a normal distribution. Categorical measurements of the patients that participated in the study were summarized in numbers and percentages, and the numerical measurements are summarized as averages and standard deviation (median and minimum-maximum when necessary). The correlation between the continuous measurements is examined using spearman correlation coefficient.

RESULTS

Sixty patients were included in the study in total. 43.3% of the patients that participated in the study were female (n:26), 56.7% were male (n:34), and the mean age of the patients was 70.58±14.86 (min: 21, max: 93) years. When we examined diagnoses of the patients, we found that sepsis diagnosis was made at the most at the rate of 43.3%, and it is followed by acute renal failure at the rate of 13.3%, and gastrointestinal bleeding and aspiration pneumonia each at the rate of 8.3%. Mean CVP value of the patients that participated in the study

was $5.70 \pm (5.01)$ cmH₂O (min: 0, max: 15). Mean value of VCI-e diameter was $14.72 \pm (5.0)$ mm (min: 2, max: 24), mean value of VCI-i diameter was $9.1 \pm (5.9)$ mm (min: 2, max: 21), mean value of VCI-CI was $52.49\% \pm (26.53)$ (min: 13, max: 100), mean value of VCI -DI was $61.29\% \pm (\text{min}:14 \text{ max}:100)$ (**Table 1**).

Table 1. Mean values of CVP, VCI-e diameter, VCI-i diameter, VCI-CI and VCI-DI of the patients					
	CVP (cmH₂O)	VCI-e (mm)	VCI-i (mm)	VCI-CI (%)	VCI-DI (%)
N	60	60	60	36	24
Mean	5.7	14.72	9.1	52.49	61.29
Sd.	5.01	5.00	5.90	26.53	31.35
Min	.00	6.40	2.00	13.00	14
Max	15.00	24.40	21.00	100.00	100

The study found that there is a weak positive correlation between systolic blood pressure, diastolic blood pressure, mean arterial pressure and CVP, and there is a weak negative correlation between heart rate and CVP. No correlation was found between respiratory rate and CVP (Table 2). In addition, the study also found that there is a weak negative correlation between systolic blood pressure, diastolic blood pressure, average arterial pressure, respiratory rate and VCI-CI, and a weak positive correlation between heart rate and VCI-CI (**Table 2**).

Table 2. Correlation among Systolic and Diastolic Blood Pressure, Heart Rate, Respiratory Rate, Mean Arterial Pressure and CVP and VCI-CI			
		CVP	VCI-CI
Heart Rate	R	-.32	,39
	P	.01	,002
	N	60	60
Diastolic Blood Pressure	R	.29	-,26
	P	.02	,04
	N	60	60
Mean arterial pressure	R	.28	-,27
	P	.02	,03
	N	60	60
Systolic Blood Pressure	R	.24	-,26
	P	.05	,04
	N	60	60
Respiratory Rate	R	-.00	-,14
	P	.96	,26
	N	60	60

The study found a positive correlation between CVP and VCI-e diameter, VCI-i diameter, and a strong negative correlation between CVP and VCI-CI/VCI-DI (**Table 3**) (**Figure 2,3,4,5**).

Table 3. Correlation among CVP and VCI-e diameter, VCI-i diameter, VCI-CI, VCI-DI				
	CVP and VCI-e diameter	CVP and VCI-i diameter	CVP and VCI-CI	CVP and VCI-DI
R	.86	.88	-.81	-.85
P	.00	.00	.00	.00
N	60	60	36	24

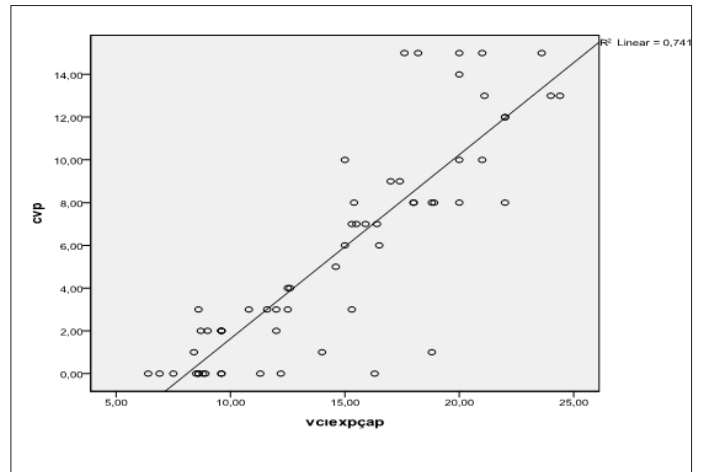


Figure 2. Chart of Correlation between CVP and VCI-e Diameter

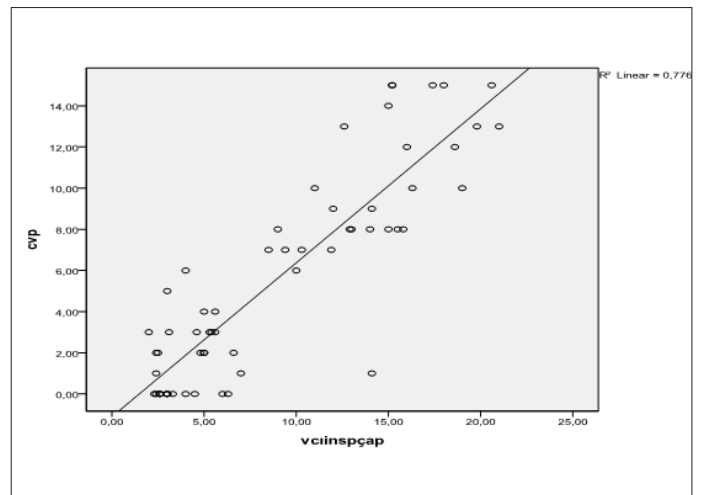


Figure 3. Chart of Correlation between CVP and VCI-i diameter

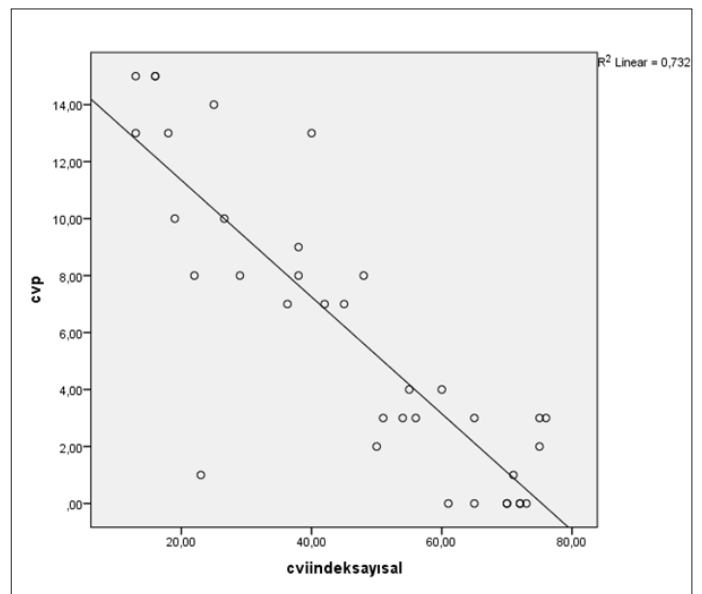


Figure 4. Chart of Correlation between CVP and VCI-CI

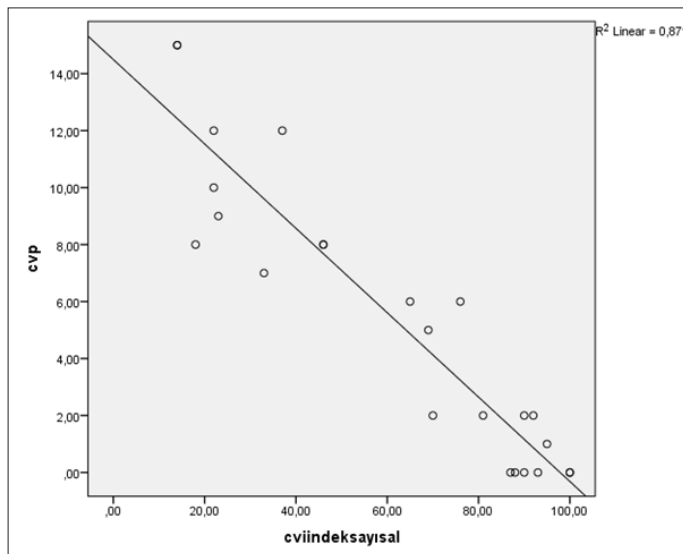


Figure 5. Chart of Correlation between CVP and VCI-DI

The study found that mean CVP value of the patients whose VCI-CI value is 20% and below (n:9) is 13.2 (2.58) cmH₂O (min: 8, max: 15); mean CVP value of those whose VCI-CI is below 50% (n:28) is 10.42 (2.989) cmH₂O (min: 7, max: 15); mean value of those whose VCI-CI is above 50% (n:32) is 1.78 (1.869) cmH₂O (min: 0, max: 6) (**Table 4**).

Table 4. Comparison of CVP values of the patients with VCI-CI above and below 50% and those below 20%

CVP	N	Mean	Sd.	Min.	Max
VCI-C < %50	28	10.42	2.98	7	15.00
VCI-CI > %50	32	1.78	1.86	.00	6.00
VCI-CI < %20	9	13.22	2.58	8.00	15.00

The study found that CVP measurement was made on 30 patients (50%) to whom a central venous catheter was inserted through internal jugular vein (IJV) and the other 30 patients (50%) to whom a central venous catheter was inserted through femoral vein (FV). The study also found that VCI-CI mean of the patients to whom central venous catheter was inserted through IJV is 54.86± (27.69) %; average VCI-CI value of the patients to whom the catheter was inserted through FV is 50.13± (25.579) %. No significant difference is found among CVP and VCI diameters of the patients to whom central venous catheter was inserted through IJV and FV (p>0.05) (**Table 5**).

Table 5. Comparison of CVP, VCI diameters values the patients with IJV and FV catheterization

		N	%	Mean	P
CVP	IJV	30	50	5.43±5.09	.72
	FV	30	50	5.96±4.96	
VCI-e	IJV	30	50	14.72±5.14	.95
	FV	30	50	14.72±4.9	
VCI-i	IJV	30	50	9.01±6.19	.96
	FV	30	50	9.19±5.69	

The study found that 60% of the patients that participated in the study have spontaneous respiration, and 40% receive mechanical ventilatory support. No significant difference was found in VCI-i, VCI-e diameter, VCI-CI, VCI-DI and CVP between the two groups (**Table 6**).

Table 6. Comparison of CVP, VCI-e, VCI-i diameter, VCI-CI values of the patients with mechanical ventilation support and spontaneous respiration

	N	%	CVP	VCI-e diameter	VCI-i diameter	VCI-DI/ VCI-CI
Mechanical ventilation	24	40	14.71±5.93	9.37±6.03	5.41±4.97	61.29±31.35
Spontaneous respiration	36	60	14.73±4.81	8.91±5.89	5.88±5.09	52.49±26.53
P			.62	.60	.98	.051

The study found that 38.3% of the patients that participated in the study were discharged with full recovery, and 61.7% died.

DISCUSSION

Volume treatment is crucial in treatment of the patients in critical condition and the patients with acute circulatory failure in particular. The purpose of volume resuscitation is to preserve sufficient tissue perfusion while avoiding any significant interstitial edema. Assessment of intravascular volume status in a timely, accurately and repeatable manner is important in adequate treatment of the critical patients.^[12,14] Hemodynamic dysfunction occurs in various clinical environments ranging from polyclinics to trauma-resuscitation areas. In addition, it can also occur in a wide range of clinical presentations ranging from mild systemic inflammatory response syndrome to advanced hemodynamic collapse and shock. For this reason, many researchers have carried out studies to develop reliable techniques or introduce biomarkers in order to foresee the volume response in the patients in critical condition.^[13] Therefore, it is crucial for clinicians to use POCUS which is quick, reliable and easy-to-use device in assessment of volume status of the patients in critical condition.

The reference point for assessing vascular volume status and heart preload in the patients in critical condition is monitorization of central venous pressure. CVP must certainly be monitored in cases that volume replacement must be made carefully such as shock and circulation failure, pediatric or cardiac diseases of the patients that apply to emergency service.^[15] However, CVP monitorization requires central venous catheterization by an invasive method which is often difficult in an urgent resuscitation and even impossible if the physician is inexperienced or in a countryside.^[16] POCUS is a technique that is often used in emergency services and intensive care units. It is even used in prehospital interventions and battlefields. POCUS is reliable, non-invasive and portable and images are easily interpreted by a broad range of specialists. Accurate measurements of internal organs and blood veins in particular can be made easily by POCUS.^[17]

Vena cava inferior is the largest vein with the lowest pressure in the venous system. Vasodilation reflects the changes in venous pressure to a certain extent. These changes also reflect volume excess. Therefore, VCI diameter can be used as an important means of diagnosis in assessment of hypervolemia and volume status.^[17,18]

Morishita et al.^[19] compared base deficit (Bee), lactate, circulating blood volume calculated by using pulse dye densitometry (PDD) which is a new method employed in detection of circulating blood volume to VCI-e, ANP and BNP in their study on healthy volunteers and hemodialysis patients; they stated that VCI-e is more important than both ANP and BNP in detecting circulating blood volume in hemodialysis patients. In 2010, Akilli et al.^[20] carried out a study on 50 volunteers and 28 sequential hemorrhagic shock patients and they that VCI diameter measured by POCUS is a stronger predictor value than other non-invasive indicators that are commonly used in estimation of shock index and acute hemorrhage such as blood pressure, pulse rate, serum lactate level and base deficit in hemorrhagic shock patients. In 2018, Sahalaby et al.^[21] carried out a study on 50 patients and they stated that there is a high correlation between VCI-CI, VCI diameter max. and CVP. In 2017, Vaish et al.^[22] carried out a study on 50 patients staying in pediatric intensive care unit and stated that there is a positive correlation between VCI diameter and CVP, and a negative correlation between VCI diameter and VCI-CI. Besides, they indicated in this study that effective volume treatment increases VCI diameter and decreases VCI-CI. A recent study was conducted with 76 patients who were followed up for septic shock and were breathing under mechanical ventilation support. As a result of this study, it was reported that VCI-DI value is a good predictor to evaluate fluid response in septic shock in mechanically ventilated patients.^[23] In literature, there are many similar studies indicating the correlation between ultrasonographic measurement of VCI and CVP.^[16,24,25] In this study, we found that there is a high positive correlation between CVP and VCI-e diameter, VCI-i diameter and high negative correlation between CVP with VCI-CI and VCI-DI (**Table 3**) (**Figure 2,3,4,5**). Findings of this study are consistent with the literature.

Zhang et al.^[1] investigated the publications until May 2013 and reviewed 8 studies in total. All studies included critical patients, one study dealt with pediatric patients. There were 235 patients in total. Five studies included the patients receiving mechanical ventilatory support; two studies included the patients with spontaneous respiration; the remaining study didn't state the respiratory pattern. All these studies indicated that vena cava inferior measurements by POCUS is of great importance in estimating the volume response on particularly the patients receiving mechanical ventilatory support and the patients that were resuscitated with colloids. In their study Joerg C.Schefold et al. assessed the correlation between the sonographic changes in VCI diameter and the invasive

hemodynamic changes in 30 patients who were diagnosed with sepsis and septic shock and monitored in mechanical ventilation support in intensive care unit and they indicated that sonographic measurement of VCI is correlated to CVP and other invasive parameters in assessment of volume status in the patients receiving mechanical ventilatory support.^[26] In this study, we found that 60% of the patients that participated in the study have spontaneous respiration, 40% of them receive mechanical ventilatory support and there is not any significant difference in VCI-i, VCI-e diameter, VCI-CI, VCI-DI and CVP values between the two groups (**Table 6**). In consequence of the literature and this study, one can say that mechanical ventilation doesn't pose an obstacle for measuring VCI and VCI-CI by POCUS in detecting volume deficit of patients and assessing the response to the volume treatment.

Pacheco Sda et al.^[27] indicated that there is a correlation between the CVP measured in 60 patients to whom central venous catheter was inserted through internal jugular vein or subclavian vein after heart surgery and the CVP measured at reference (zero) level on femoral central venous catheter of the same patients. Boone et al.^[28] indicated that measurement of CVP in 40 patients through femoral veins is sufficient as much as the measurement of CVP through IJV and subclavian vein in the postoperative patients. This study found that CVP measurement was made on 30 patients (50%) to whom a central venous catheter was inserted through IJV and the other 30 patients (50%) to whom a central venous catheter was inserted through FV. CVP mean of the patients to whom central venous catheter was inserted through IJV was $5.43 \pm (5.09)$; mean CVP value of the patients that were catheterized through FV was $5.96 \pm (4.96)$. No significant difference was found among CVP and VCI diameters in the patients to whom central venous catheter was inserted through IJV and FV ($p > 0.05$) (**Table 5**). This study reveals findings that are consistent with the literature.

Limitations

FV catheterization is not recommended for CVP measurement. CVP measurement was made from FV in 50% of the patients in our research. This is one of the limitations of our study. However, in our study, no statistical difference was detected between the CVP values measured from FV and IJV.

CONCLUSION

Consequently, measurement of respiratory variability in VCI diameter by POCUS which is a rapid, reliable, easily applicable, cost-efficient and non-invasive method for the critical patients that have spontaneous respiration or receive mechanical ventilatory support in emergency services and intensive care units can be used in order to assess volume status and estimate CVP. In addition, we consider that the use of POCUS should be extended in emergency services and intensive care units where critical patients are monitored.

ETHICAL DECLARATIONS

Ethics Committee Approval: The study was carried out with the permission of Ondokuz Mayıs University Medical Faculty Clinical Research Ethics Committee dated 30.05.2015 (OMU CREC decree no: 2014/675).

Informed Consent: All patients signed the free and informed consent form.

Referee Evaluation Process: Externally peer-reviewed.

Conflict of Interest Statement: The authors have no conflicts of interest to declare.

Financial Disclosure: The authors declared that this study has received no financial support.

Author Contributions: All of the authors declare that they have all participated in the design, execution, and analysis of the paper, and that they have approved the final version.

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