

Comparing Outcomes and Complication of Central Venous Cannulation Using Both Conventional and Ultrasound Guide

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Central venous cannulation (CVC) is required for management of critically ill and hemodialysis patients which has different complications in conventional procedure. The aim of this study was to compare the outcomes and complication of central venous cannulation using both conventional and ultrasound guide. A randomized controlled trial study of 336 recently hospitalized hemodialysis patients was conducted (168 in intervention group and 168 in control group). CVC was done by an ultrasound machine in intervention group that the needle was inserted perpendicular to the skin under visualization on the US screen while CVC was performed by the conventional landmark approach in the other group. The time for insertion, attempts required, and complications were measured in both groups. Data were analyzed with ANOVA Repeated Measure. In intervention group 22 patients (13.09%) required more than one attempt, while in the control group 75 patients (44.6%) required more than one attempt. Statistically this difference was significant ($P.V = 0.000$). In the control group, arterial puncture was happened in 10 patients (5.9%), and hematoma in 5 patient (2.9%), while were 2 patients (1.1%) and 1 patients (0.59%) in the ultrasound group respectively ($P.V = 0.04$ and $P.V = 0.05$). The results of our study showed that USG approach took lesser time, required lesser attempts, and had lower incidence of complications for cannulation of the internal jugular vein.

Key words: Central venous cannulation, Hemodialysis, Conventional, Ultrasound guide.

Central venous cannulation (CVC) is required for management of critically ill and hemodialysis patients. In patients with liver disease and renal failure, CVC is an important intervention for fluid, drugs and concentrated substances such as glucose solution administration and conducting some procedures such as biopsies and measuring venous pressure¹. Insertion of CVC has both vascular and non-vascular complications, which are decreased in

recent years with the introduction of ultrasound guidance (USG) cannulation²⁻⁵.

USG has different performance such as cause of abdominal distention, DVT and pulmonary status assessment, pericardial tamponade, etc. A common use of ultrasonography in hemodialysis patients is in central venous cannulation, which has been a put blindly using landmark. This conventional cannulation had different complications such as arterial puncture, pneumothorax, hemothorax and air embolism which have been reduced by appearing ultrasonography that the needle can be visualized entering the vein,⁶ while the conventional method is by direct

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palpation of the artery and puncture with a catheter and followed by threading of the cannula into the vessel which make different complications as mentioned before⁸.

Indications of USG cannulation is for hypotensive patients with difficult palpate of carotid artery, measuring the central venous pressure, administration of inotropes, parenteral nutrition, hemodialysis, etc. Moreover, some of these indications are of emergency that USG CVC can be performed faster than conventional procedure. In some countries, CVC under US guidance is likely to be made compulsory in the near future [9]. This study was designed to investigate the outcomes and complication of central venous cannulation using both conventional and ultrasound guide and compared them in hemodialysis patients.

MATERIALS AND METHODS

Procedures

This randomized controlled trial study compared the outcomes and complication of central venous cannulation using both conventional and ultrasound guide in patients needing hemodialysis undergone central venous cannulation with ultrasound guide (study group) to the conventional procedure (control group). An inclusion criterion was all patients who have had indications for central venous cannulation. An exclusion criterion was patients with previous CVC within 15 days, anatomical deformity (such as neck surgery, malignancy and burns on the site), having emergency conditions and bleeding disorders.

In the intervention group, the ultrasound machine SiteRite II (Bard Access, Inc., Salt Lake City, Utah) with 7.5 MHz probe was used. Special jelly as a matching layer for ultrasound was rubbed over the area. The probe was covered with sterile sheath and placed over the anterior neck triangle (between two heads of the sternocleidomastoid muscle (SCM)). The vessels were seen in the transverse section. Internal carotid artery was seen as pulsating, while the internal jugular vein in lateral was nonpulsating which got compressed by pressuring the probe. Then the needle was inserted directed vertically to the skin under visualization on the US screen. After successful aspiration of blood, a J-shaped guide wire was

inserted through the hollow needle, then dilator was inserted through the guide wire which was withdrawn and sutured.

In control CVC was performed by the conventional landmark. The patient was in supine position with slight head down with contralateral side position in order to palpate the SCMs and ICA. The ICA was pressed slightly with fingers so that it does not overly the IJV. Then the central line needle was inserted in the lateral of ICA pulsations site. After successful aspiration of blood, rest of the procedure was similar to the intervention group. After the procedure, a chest X-ray was performed for all the patients to rule out a pneumothorax.

Participants

Participants were recruited between August and December 2014 from Golestan hospitals in Ahvas in Iran. All patients who were hospitalized for central venous cannulation diagnosed by a nephrologist during this time were eligible for participation. From patients 340 cases were selected based on inclusion and exclusion criteria, 4 of them not participate to the study and other 336 cases allocated with a simple random sampling method in 2 groups of intervention and control (168 cases in each groups). The study received ethics approval from the relevant institutional review committees, and all participants gave written informed consent.

Data analysis

Analyses were performed with statistical package of SPSS (version 20) using descriptive statistics such as mean and standard deviation and analytical statistics such as ANOVA repeated measure.

RESULTS

The mean age of the patients in intervention group (90 males, 78 females) was 54.8 ± 7.6 years and in control group (87 males, 81 females) was 49.9 ± 10.6 years. Table 1 shows comparison of demographic findings of two groups.

The mean time to successful aspiration of venous blood after completion prep of skin in intervention group was 132.52 secs, while in control group was 169.2 secs which was statistically significant ($P.V=0.000$). In intervention group 22 patients (13.09%) required more than one attempt, while in the control group 75 patients (44.6%)

Table 1. Comparison of demographic variables in two groups

Groups Variables	Intervention	Control	P- value
Age: Mean(\pm SD)	54.8 (7.6)	49.9 (10.6)	0.073
Sex: Number (%)			
Male	90 (53.5)	87 (51.7)	0.74
Female	78 (46.4)	81 (48.2)	
Time of renal failure : Mean(\pm SD) (year)	9.23 (4.5)	10 (5.1)	0.701

Table 2. Comparison of outcomes and complications in both studied groups

Groups Variables Mean(\pm SD , n)	Intervention	Control	P- value
Time from skin prep to successful aspiration (in seconds)	132.52(\pm 15.1 , 168)	169.2(\pm 16.21 , 168)	0.000
Attempts for			
No. of patients required more than one attempt (%)	22 (13.09)	75 (44.6)	0.000
successful cannulation			
Mean No. of attempt	1.4 (\pm 0.42 , 168)	1.98 (\pm 0.61 , 168)	0.03
Complications			
Arterial puncture	2 (1.1)	10 (5.9)	0.04
Hemothorax	0	1 (0.59)	0.970
Pneumothorax	0	2 (1.1)	0.87
Hematoma	1 (0.59)	5 (2.9)	0.05
Sudden death	0	0	N.S

required more than one attempt. Statistically this difference was significant ($P.V = 0.000$). An average of 1.4 attempts per cannulation was required for intervention group, whereas for control group average 1.98 attempts were required (Table 2).

In the control group, arterial puncture was happened in 10 patients (5.9%), and hematoma in 5 patient (2.9%), while were 2 patients (1.1%) and 1 patients (0.59%) in the ultrasound group respectively ($P.V = 0.04$ and $P.V = 0.05$) (Table 2).

DISCUSSION

Side effects of CVC are not rare and on the other hand they can be serious problems with poor prognosis in some cases. Recently the use of ultrasound for CVC has been studied to reduce the complications which are dependent to operator and the availability of the equipment¹⁰. The use of ultrasonography with professional operator decrease the number of complications and attempts compared with the routine method (landmark method). In some studies, it was indicated that the complications of CVC insertion are increased in three or more attempts compared with a single attempt¹¹. In our study, it was found that in

intervention group 22 patients (13.09%) required more than one attempt, while in the control group 75 patients (44.6%) required more than one attempt which increase the complications as mentioned before. Ishii *et al.* performed a study looking at infants and small children requiring radial artery cannulation and They showed that success rates after a single attempt were significantly higher in the US group (76.3%) than those in the conventional group (35.6%) ($p < 0.001$)¹².

On the other hand Ganesh *et al.* found no statistically significant differences between the US-guided cannulation and palpation technique groups in the time to successful cannulation, total number of attempts, number of successful cannulations during the first attempt or number of cannulae used for catheterization¹³.

Moreover our study showed that the complications of conventional method was more in totally compared with ultrasound group but just two of complications show significant differences between two groups (arterial puncture and hematoma). Several randomized controlled trials have showed the value of US in arterial catheter insertion compared with conventional procedure. Shiver and colleagues demonstrated that a first-pass

success rate was 87% in the US group compared with 50% in the conventional group. Moreover they showed that US method was associated with a 43% reduction in the development of hematoma at the insertion site¹⁴. Levin *et al.* indicated a first-pass success rate was 62% in US group compared with 34% in control group¹⁵. In the study performed by Ankit Agarwal *et al.* demonstrated that arterial puncture happened in four patients (10%), and pneumothorax in one patient (2.5%) in conventional procedure, while there were no such complications in the ultrasound group¹⁶.

While Tada *et al.* showed that US guidance offered no additional benefit in cases where the radial arterial pulse was palpable¹⁷. This difference in results of mentioned study and our study was the sample size which was 336 in our study and 166 in Tada's study. Moreover, the procedures are different that we used central venous, whereas they study on the peripheral artery line insertion.

On the other hand in some literatures indicated that disadvantage associated with USG-guided CVC, is procedure-related increased incidence of infection, which can be reduced by the use of a two-operator technique with sterile self-adhesive plastic and povidone iodine solution¹⁸.

CONCLUSION

The results of our study showed that USG approach took lesser time, required lesser attempts, and had lower incidence of complications for cannulation of the internal jugular vein. Regular use of USG for CVC will benefit for patients required CVC for most of time especially hemodialysis patients. It would be a costly investment in a developing country like Iran, one must keep in mind that use of USG is a prudent approach as USG-guided CVC is easier, quicker, and safer than landmark approach.

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