The effect of changes in patients' body position on the back pain intensity and hemodynamic status during and after radiofrequency catheter ablation of cardiac dysrhythmias

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ABSTRACT

Background: After radiofrequency catheter ablation of arrhythmias, patients have to bed rest for 4-6 h to prevent bleeding and hematoma. However, such a rest may cause back pain in the patients. The aim of this study was to determine the effects of continuous change in body position during and after the radiofrequency ablation on the back pain.

Materials and Methods: In a quasi-experimental design 75 patients referring to university-affiliated hospitals were randomly assigned to a control group, receiving no change in body position, group A subjected to changes in body position during and after ablation, and group B subjected to changes in body position during ablation. The intensity of pain, blood pressure, heart rate, and extent of bleeding and hematoma were measured.

Results: The groups were not significantly different in terms of demographic characteristics, blood pressure, heart rate, overall bleeding, or hematoma at the entry into the coronary care unit. While not significantly different from each other, the intensity of back pain between group A and B were significantly lower than that of group C. Compared to group C, group A and B had a significantly lower pain score up to 6 and 4 h after the procedure, respectively. Group B had a significantly higher pain score at 2, 4, and 6 h post ablation than group A.

Conclusions: The findings show that changing the body position during and after the ablation procedure would reduce or prevent the back pain without increasing the chance of bleeding and hematoma.

Key words: Back pain, bleeding, hematoma, Iran, patient positioning, radiofrequency catheter ablation

INTRODUCTION

ysrhythmia caused or contributed to 479,000 of more than 2,400,000 deaths in the United States in 2003.^[1] Atrioventricular node reentry tachycardia is the most common supraventricular tachycardia.^[2] Atrial fibrillation is the most common sustained clinical arrhythmia.^[3]

In case the drugs are not effective in decreasing or elimination of dysrhythmia, non-medical treatments such as ablation

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would be applied.^[4] Radiofrequency catheter ablation is the standard therapy for treatment of different types of dysrhythmias^[5] such as supraventricular dysrhythmia in patients with Wolfss–Parkinson–White syndrome and atrioventricular node reentry tachycardia.^[6] The success of catheter ablation to eliminate accessory pathways is almost 95%.^[7] Catheter ablation destroys specific cells, which are the reason or the center of dysrhythmia.^[4] The duration of the procedure differs from 1 to 6 h based on the type of arrhythmia and other factors.^[8]

Back pain is quite common after cardiac catheterization, and is caused by the position of the body, which is surrounded by the bed.^[9-13] To avoid bleeding and hematoma, patients are instructed to rest in bed with their foot in a straight position for 4-6 h after the procedure.^[14] Such a position, while reducing the vascular complications of the procedure, often leads to back pain and other adverse effects such as hemodynamic status instability.^[1,15] Through the activation of autonomic nervous system and the release of epinephrine,^[16] pain leads to many harmful effects including increased heart rate,^[17,18] blood pressure,^[19,21]

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and myocardial work load^[22] and also dysrhythmia.^[4] Lying on the back for a long time imposes pressure, and causes cellular ischemia and pain in the lumbar and the back.^[23] Therefore, patients intend to change their position so as to reduce the pain and discomfort.^[9,11]

Considering the possible incongruence of the patients' views with the care providers perspectives^[24], the most important factor in comforting the patients is the relief of pain and a suitable position from the patient's attitude.^[25] Patients' positioning is based on experience and thought, and there is no scientific basis for that.^[2,26] Inner-muscle pressure in lumbar muscles has a direct relation with the patients' position and the imposed load on the muscles.^[27] Changes in patients' position following angiography could result in a stable hemodynamic status by reducing their pain without increasing vascular complications.^[15] The present study was designed to examine the effects of a combination of reducing or preventing techniques, such as adjusting the angle of the bed with the horizontal level, changing the body position, and supportive pillow under the body and knees, on back pain following radiofrequency catheter ablation in patients with cardiac dysrhythmia.

MATERIALS AND METHODS

This is a quasi-experimental study performed at Shahid Faghihi and Kowsar hospitals in Shiraz, Iran, in 2010-2011. The inclusion criteria were eligibility for treatment by radiofrequency catheter ablation, a systolic blood pressure less than 190 mmHg, a diastolic blood pressure less than 110 mmHg, and consent to participate in the study. The exclusion criteria were spinal cord disorders, inter-vertebral disk disorders, bleeding disorders, the use of anticoagulant and analgesic drugs 24 h prior to the procedure, advanced heart failure, a procedure duration less than $1\frac{1}{2}$ h and longer than $3\frac{1}{2}$ h, increased bleeding or any other disorders during the procedure, severe decrease of consciousness level, performing ablation procedure through artery, and tranquilizers' use during the procedure.

Using data from a previous study,^[15] an alpha value of 0.05, power of 0.80, and effect size of 2 for pain score, the sample size was calculated to be 16 patients in each group. The samples were increased to 25 in each group (total 75 patients) to account for missing data due to any reason. The positioning protocol of the study was designed based on Rezaei-Adaryani and colleagues'protocol.^[15] We also used pillow for knee positioning. The study protocol was approved by the research team and then Ethics Committee, Shiraz University of Medical Sciences. The patients (N = 75) were selected using purposive sampling. After obtaining written consent, the participants were randomly allocated to two case groups (A and B) and a

control group (25 each).

Patients in the control group received routine care (with no change in position) including lying on their back for 6-8 h, which comprised of duration of the procedure.

Patients in group A did lie on their back during the procedure with the head of their bed tilted up for 15°, using a small mattress $(3 \times 30 \times 80 \text{ cm})$ under their body and their knee bent for 30° using a small cylindrical pillow. In the first and second hours after the procedure, they continued lying on their back with the head of the bed tilted up for 30° and 45°, respectively. During the 2 hours post procedure, a small pillow was placed alternatively every 30 min under right and left sides of the body from shoulder to gluteal. Moreover, they continued to use the small cylindrical pillow under their knees. During the third hour, head of the bed was tilted up to 30° and the patients did lie on right and left sides for 30 min each. Within the fourth hour, the patients lied on their side (right or left) for 30 min with the head of the bed tilted up for 15°, followed by sitting for the next 30 min. In case of group B, the samples were positioned precisely like patients in group A during the procedure, and they were in the routine position after the procedure (without any alteration). The intensity of the back pain, blood pressure, heart rate, and the amount of bleeding and hematoma were measured right after the patients entered the coronary care unit (CCU) and 2, 4, 6, and 24 h after ablation in the three groups.

The data were collected using a questionnaire containing questions regarding patients' demographic characteristics including age, gender, body mass index, marital status, education level, type of arrhythmia, previous history of ablation, angiography, open-heart surgery and changing the heart valves, the duration of ablation procedure, blood pressure, heart rate, prothrombin time, and the international normalized ratio, as well as a numeric rating scale, a standard mercury sphygmomanometer (ALPK2, Japan), a two-dimensional ruler, a scale, and a goniometer.

The numerical scale used for the evaluation of pain intensity scored pain from 0 (least severe) to 10 (most severe), and its validity and reliability had been established. Different forms of pain evaluation numeric scale have provided acceptable data with coefficients more than 99%.^[28,29] Blood pressure was measured using a standard mercury sphygmomanometer. The amount of bleeding and hematoma was measured using two-dimensional ruler with the precision of 1 cm². The ruler was reported to be highly consistent with a correlation coefficient of 0.96.^[15] The goniometer was

used to measure the angle of head of the bed and the patients' knee angles.

The findings were analyzed using Chi-square and Kruskal-Wallis test. Where a significant difference was found with Kruskal-Wallis, pairwise comparisons were performed using Mann-Whitney U-test with Bonferroni adjustment. Statistical Package for Social Sciences (SPSS version 11.5) was used for data analysis at a $P \le 0.05$.

RESULTS

Seventy-five patients did participate in the study, and all of them did finish it. The patients' baseline characteristics are shown in Table 1. There was no significant difference among the three groups in terms of gender, marital status, education level, type of arrhythmia, age, and body mass index, duration of ablation procedure, heart rate, mean arterial pressure, prothrombin time, or international normalized ratio. Also, the groups were not significantly different in terms of previous use of radiofrequency catheter ablation or coronary angiography.

The pain intensity was significantly (P < 0.001) different among the three groups at immediately, 2, 4, or 6 h after ablation. However, it was not significantly different at 24 h after ablation among the three groups.

There was a significant difference (P < 0.001) between the control group and groups A and B in terms of pain intensity immediately after ablation. However, there was no significant difference between groups A and B. At 2 h after ablation, there was significant difference between the control group and groups A and B, as well as between groups A and B. At 4 h after ablation, there was a significant difference between groups A and B. At 4 h after ablation, there was a significant difference between groups A and B, as well as between groups A and B. At 4 h after ablation, there was a significant difference between groups A and B, group A and control group (P < 0.001), and group B and control group (P = 0.010) in terms of pain

intensity. At 6 h after ablation, there was a significant difference in pain intensity between groups A and B (P < 0.005) and group A and control group (P < 0.001). However, it was not significantly different between group B and control group (P = 0.144). There was no significant difference in the pain intensity among the three groups at 24 h after ablation. Table 2 shows the comparison of the pain intensity among the three groups at different times after ablation.

There was no significant difference in the arterial blood pressure or heart rate of the three groups at immediately, 2, 4, 6, or 24 h after ablation.

There was no significant difference among the three groups regarding the amounts of overall bleeding and overall hematoma formation.

DISCUSSION

The findings of this study indicate that patients in the control group experienced more pain compared to those in group A or B. It could be due to the prolonged rest without any change in the patients' position. This shows that the more the patients' rest in a flat position after ablation, the more intense back pain they would experience. No significant difference between groups A and B immediately after ablation could be due to similar body position in the two groups. Compared to patients in groups A, those in group B experienced more pain after ablation at 2, 4, and 6 h after ablation. Previous studies showed that changes in body position resulted in significant decrease of pain intensity at various times after angiography.^[9,15,30,31] Also, no significant difference was observed in the mean arterial pressure or heart rate among participants at various times after ablation. However, such a finding is in contrast to that of Adaryani et al. who showed that mean arterial

Table 1: Patients' baseline characteristics and group matching (the results of Kruskal-Wallis and Chi-square test)

Variable	Group A (<i>n</i> =25)	Group B (<i>n</i> =25)	Control group (<i>n</i> =25)	P value
Gender (female/male), %	72/28	64/36	80/20	0/425*
Marital status (single/married), %	12/88	8/92	12/88	0/869*
Education (illiterate and primary school/middle and high school/university education), %	48/24/28	48/28/24	32/40/28	0.693*
Type of arrhythmia (AVNRT/AVRT/other), %	68/28/4	84/12/4	76/8/16	0.140*
Age (years)	45.2±11.8	46.4±15.3	49.4±12.8	0.506**
Body mass index (BMI)	25.7±4.2	25.6±4.0	26.1±3.4	0.596**
Duration of the procedure (minutes)	122.8±22.4	118.6±22.0	121.8±27.2	0.747**
Heart rate (beats per minute)	81.8±12.3	83.2±14.2	77.0±11.8	0.154**
Mean arterial pressure (mmHg)	1.0±14.1	95.6±11.6	98.2±11.5	0.431**
Prothrombin time (seconds)	12.8±0.7	12.5±0.6	12.8±1.2	0.379**
International Normalized Ratio	1.0±0.1	1.0±0.0	1.1±0.1	0.622**

*Chi-square, AVNRT: Atrioventricular node re-entry tachycardia, AVRT: Atrioventricular re-entry tachycardia, **Kruskal Walis

Table 2: Comparison of the pain intensity (mean \pm SD) among the three groups at different times after cardiac radiofrequency ablation

Group time	Group A (<i>n</i> =25)	Group B (<i>n</i> =25)	Control (<i>n</i> =25)	P value
T1 (after ablation)	1.0±1.1*	1.0±1.3*	3.7±1.8	<0.001
T2 (2 h later)	1.0±1.0*†	3.3±2.2*	4.9±2.2	<0.001
T3 (4 h later)	0.4±0.6*†	4.4±2.7*	6.5±2.4	<0.001
T4 (6 h later)	0.1±0.3*†	1.0±1.6	1.8±2.3	<0.001
T5 (24 h later)	0.0±0.2	0.1±0.3	0.6±1.4	0.057

*Shows significant difference from the control group, [†]Shows significant difference from group B

pressure and heart rate at 6 and 8 h after catheterization were significantly higher in the control group than those in the intervention groups.^[15] Pain could lead to high blood pressure and heart rate.^[17,18,20,21] The lack of significant difference in mean arterial pressure or heart rates among the three groups could be due to small number of the patients enrolled. In other words, compared to pains related to back and muscle spasms, pains with higher intensities may result in hemodynamic changes. Our study did not show significant difference in the amounts of overall bleeding or hematoma formation at femoral puncture site of the three groups at the measurement time points.

The findings of this study should be interpreted in the light of a number of limitations including variable pain threshold among patients, and the observation and reporting of outcomes was not blinded. Other studies involving varied changes in body position would be helpful to come up with a better nursing guideline for managing patients' pain and discomfort after cardiac radiofrequency catheter ablation.

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