



# Effect of Localized Cold Compress on Pain Severity with the Subcutaneous Injection of Enoxaparin: A Randomized Controlled Trial

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Received 2020 November 30; Revised 2021 February 08; Accepted 2021 February 09.

## Abstract

**Background:** Subcutaneous injection of enoxaparin causes local reactions, such as bruising and pain at the injection site. The pain induced by subcutaneous injection may lead to anxiety, fear of needle, and distrust in healthcare providers in the long run.

**Objectives:** The present study aimed to evaluate the impact of topical cold compress on bruising and pain intensity at the subcutaneous injection site of enoxaparin.

**Methods:** This clinical trial was conducted on 100 patients in five groups. In the control group, subcutaneous injection was performed using the routine method. In the intervention groups, local cold compress was applied five minutes prior to the injection, five minutes before and following the injection, five minutes after the injection, 20 minutes following the injection, and five minutes before and 20 minutes following the injection within 10 seconds. Data were collected using a researcher-made checklist containing two sections of individual characteristics and pain severity. In addition, pain intensity was evaluated using the visual analog scale (VAS) immediately after each injection.

**Results:** Group five showed a significant difference with the control group only in the mean pain severity ( $P < 0.05$ ), while the mean pain intensity scores had no significant difference in the other groups compared to the control group.

**Conclusions:** According to the results, the application of local cold compress before and after the subcutaneous injection of enoxaparin sodium could effectively reduce pain intensity at the injection site.

**Keywords:** Cold Therapy, Enoxaparin, Injections, Pain, Subcutaneous

## 1. Background

The safe administration of drugs is a basic skill and a common intervention performed by nurses. Anticoagulants are frequently prescribed as prophylaxis or treatment agents for venous thromboembolism. The subcutaneous injection of anticoagulants often causes complications such as pain and hematoma at the injection site. Previous studies have indicated that these complications may increase the physical and psychological discomfort of the patients (1).

Enoxaparin is applied in the treatment of patients with deep vein thrombosis (DVT) and hospitalized patients with or without pulmonary embolism, as well as DVT prevention in abdominal surgeries and hip replacement associated with severe movement restrictions, prevention of the ischemic complications of unstable angina and myocar-

dial infarction, and treatment of myocardial infarction after percutaneous coronary interventions (2, 3). Furthermore, enoxaparin is an anticoagulant with a low molecular weight, immediate effect, and long half-life. It is obtained from the de-polymerization of heparin and injected subcutaneously (4); the subcutaneous injection of heparin causes bruising and tissue injury in hospitalized patients. In general, the pain induced by subcutaneous injection causes anxiety, fear of needles, and distrust in healthcare providers in the long run (5). The rate of injection, type of drugs, and volume of the injectable drug could also affect the severity of pain. Recent findings have indicated that bruising and pain at the injection site of the mentioned group of drugs is a grave concern as aggravation may result from two different phenomena. Firstly, the insertion of a needle into the skin stimulates and damages neurons, thereby leading to the sensation of pain (6). On the other

hand, the sulfate bonds in the drug exacerbate its acidic properties, increasing the sensation of pain at the injection site (7).

Safe and accurate injection is an important responsibility of nurses. Therefore, they should be familiar with appropriate and effective strategies based on scientific evidence in order to reduce potential damages and complications in hospitalized patients (7, 8). Some of these strategies include applying pressure before injection, needle cooing, injection at a specific time, and cooling the injection site beforehand or warming afterwards. In addition, nurses must have an adequate knowledge of the possible side-effects of injectable drugs.

Several studies have been focused on reducing the complications of subcutaneous injection and the pain caused by sodium enoxaparin (9, 10).

## 2. Objectives

Given the importance of anticoagulant use in different patients and the undeniable topical side-effects of these drugs on patient cooperation and treatment adherence, the present study aimed to investigate the impact of topical cold application on the pain intensity of patients at the subcutaneous injection site of enoxaparin sodium.

## 3. Methods

This randomized clinical trial was conducted to assess the effectiveness of cold compress on the severity of subcutaneous injection pain in the cardiac patients admitted to Ayatollah Rouhani hospital in Babol, Iran in 2015. The patients were divided into five groups, including a control group and four intervention groups.

The sampling process was performed at the cardiac section of the hospital; initially, the researchers selected the patients who were prescribed enoxaparin 6,000 units daily. Following that, the study groups were assigned a code each, and the codes were placed in a container. At the next stage, five envelopes labeled A, B, C, D, E (intervention groups), and F (control group) were selected and provided to the patients, who were blinded to the procedures. By drawing lots from the patient code container, a piece of paper was taken out, the desired code was selected from one of the envelopes, and the intervention groups were identified accordingly. At the next stage, the specified codes were returned to the container, so that the participants would have an equal chance to choose their intervention group. The process continued until reaching the desired sample size. Following that, each group received their specific treatment. The samples were selected from the patients admitted to the cardiology and CCU wards via convenience sampling. Notably, the wards were in the same

medical center and covered by a joint healthcare team. As such, the type of the provided care and environmental conditions of the wards were also the same.

Sample size was determined to be 100 individuals in five groups of 20 based on the previous studies in this regard (7). The inclusion criteria of the study were as follows: (1) age of 18-65 years; (2) not pregnancy/lactation during the study; (3) no previous injection at the site within 12 hours before the test; (4) full consciousness and ability to perceive and indicate pain and (5) no history of coagulation disorders (Figure 1). The patients received the same injectable dose of enoxaparin (6,000 units). The patients who were unwilling to continue participation or were discharged earlier than 72 hours were excluded from the study.

Initially, random sampling was performed by the researcher to enroll the patients, followed by examining the complications after injection. The control group received the injection without an intervention using a standard injection technique, and the injection was performed in the right or left abdomen within five centimeters around the umbilicus to the sides with no aspiration within 10 seconds without massage after the injection.

The patients also received training on avoiding the manipulation of the injection site. Cold compress was applied at different intervals. In the first intervention group, the cold compress (water and ice bags) was applied five minutes prior to the injection. In the second group, the cold compress was applied 20 minutes following the injection. In the third intervention group, the cold compress was applied five minutes before and 20 minutes after the injection, and in the fourth group, the cold compress was applied five minutes before and five minutes after the injection.

Data were collected by a trained researcher using a researcher-made checklist consisting of two sections, which were demographic characteristics (age, gender, disease type) and pain intensity assessment by VAS. This standard instrument is used to measure pain intensity within a scale of 0 - 10, and the patient is asked to rate their perceived pain. VAS is a 100-millimeter ruler, and the patient indicated the level of pain within the mentioned scale (no pain = 0, moderate pain = 1 - 5, severe pain = 6 - 10); the measured accuracy of pain severity in this scale is based on the visual aspect. Notably, the validity, reliability, and sensitivity of VAS have been confirmed in several studies for acute, chronic, and cancerous pain. Accordingly, the validity of the scale is 76 - 86%, and the reliability is reported to be 60 - 77% (10). In the present study, pain severity was measured immediately, 24, and 72 hours after the injection.

Data analysis was performed in SPSS version 16 (233, South Wacker Drive, 11th Floor, Chicago, USA) using descriptive statistics, independent *t*-test, and one-way analysis of

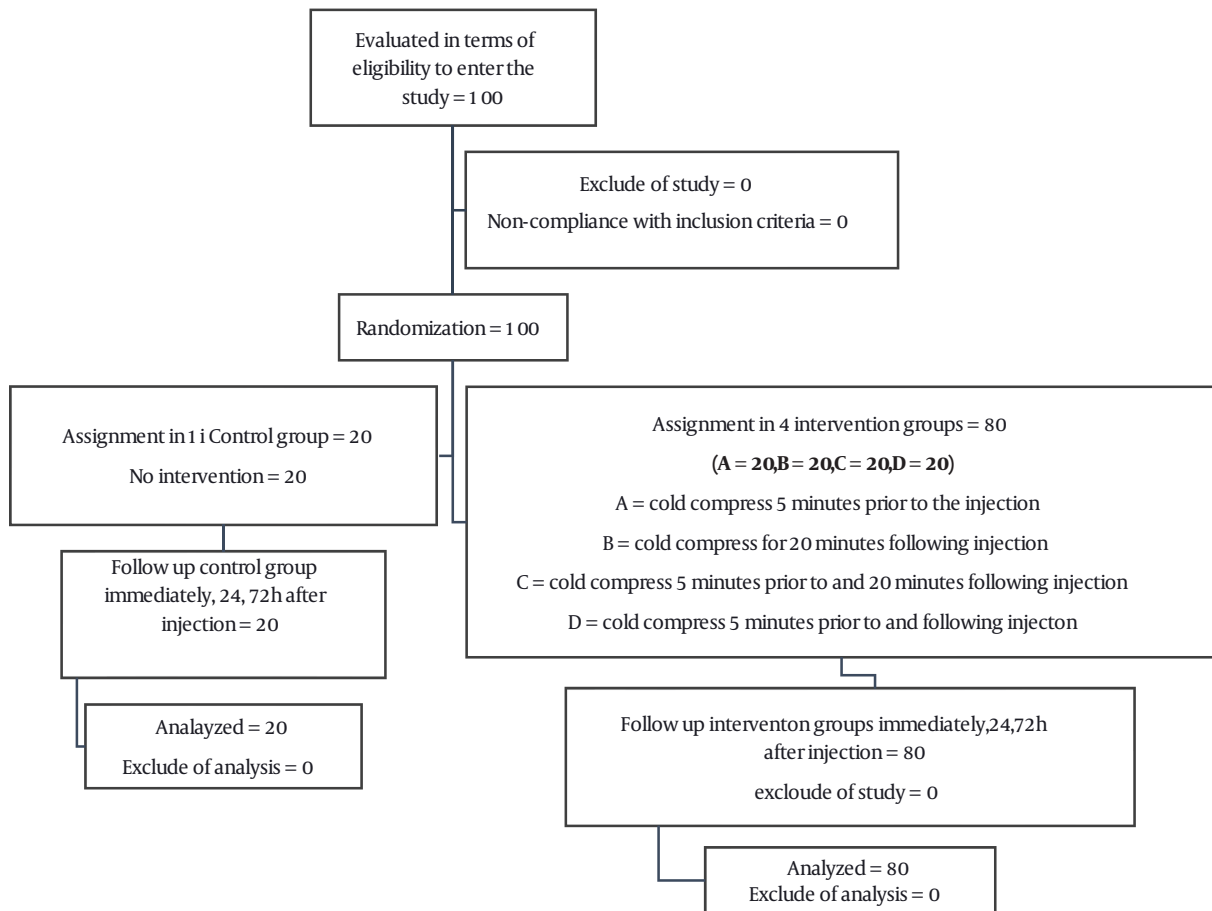


Figure 1. CONSORT diagram

variance (ANOVA). In all the statistical analyses, the significance level was set at 0.05.

#### 4. Results

According to the descriptive statistics, the mean age of the patients was  $54.66 \pm 8.603$  years, and gender distribution was not significant (Table 1). All the participants completed the research, and none were excluded. The results of ANOVA indicated a significant difference in the mean pain severity of the patients in the fourth group compared to the control group, while the mean pain intensity scores in the intervention groups had no significant difference with the control group (Figure 2).

#### 5. Discussion

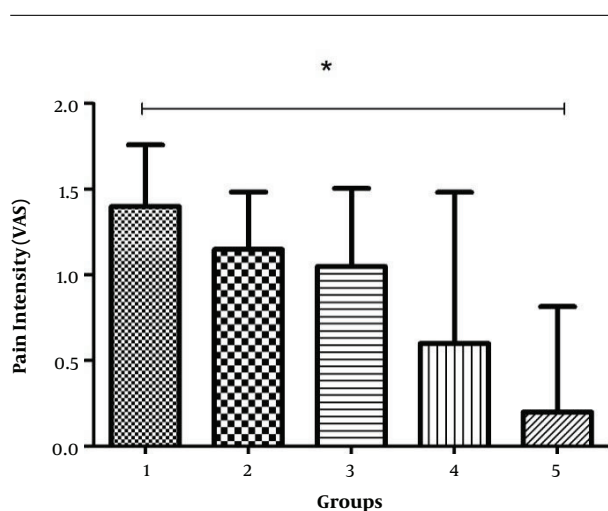
The present study aimed to compare four methods of cold compress therapy and their effect on the severity of

the pain induced by enoxaparin injection at the site. According to our findings, pain severity had a significant difference between the male and female patients and was higher in the women ( $t = 3.39$ ;  $P = 0.001$ ) (Table 2) compared to the men, which indicated the lower pain threshold and pressure tolerance in women ( $P < 0.001$ ). This is consistent with the results obtained by Soetanto (10).

The results of the present study indicated a significant difference in the mean pain intensity in the fourth group compared to the control group, while pain severity was significantly higher in the control group ( $P < 0.01$ ) (Table 3). These findings are in line with the results obtained by Farhang Ranjbar et al. in a study regarding the impact of local cold compress on the incidence of bruises and pain after enoxaparin injection. Their mentioned study indicated no significant difference in pain severity between the intervention group (local cold compress five minutes before injection) and the control group. However, the mean pain intensity in the intervention group was reported to be

**Table 1.** Comparison of Five Study Group Based on Age and Gender

Groups	Age (Mean ± SD)	Gender		Total
		Female	Male	
Control	10.68 ± 53.45	12	8	20
Cold compress 5 minutes before injection	6.18 ± 57.25	12	8	20
Cold compress 20 minutes after injection	8.30 ± 55.4	10	10	20
Cold compress 5 minutes before injection; cold compress 20 minutes after injection	10.39 ± 53.4	7	13	20
Cold compress 5 minutes before injection and 5 minutes after injection	6.66 ± 53.7	9	11	20
<b>Total</b>		50	50	100



**Figure 2.** Comparison of pain intensity in study groups (data expressed as mean ± SEM; \* P < 0.01)

**Table 2.** Gender Distribution in Five Study Groups<sup>a</sup>

Test	Value	Degree of Freedom (df)	Significance Level (Bilateral)
Pearson's chi-squared test	3.600	4	0.463

<sup>a</sup> Chi-square test (significance = 0.463;  $\chi^2 = 3.600$ ) showed no significant difference between five groups in terms of gender distribution.

lower compared to the control group. In the mentioned research, the highest recorded pain score in the control and intervention groups was 100 and 56, respectively. The difference in this regard was considered clinically significant, indicating that local cold compress could reduce the pain induced by subcutaneous injection (11).

In the present study, pain reduced in all the study groups after using the cold compress at all the intervals, while only a significant reduction was observed in pain intensity following the cold compress five minutes before

and after the injection in the control group. This is in line with the study by Sendir et al., which investigated the impact of cold compress before and after injection on the pain intensity at the subcutaneous injection site using three injection methods. The mentioned study indicated that using the cold compress prior to and following injection could significantly reduce the pain intensity at the injection site (12). The difference between the two studies could be attributed to the pretest-posttest design, leading to prolonged analgesia.

According to the findings of El-deen, two intervention groups receiving cryotherapy had significantly lower pain intensity scores compared to the control group, which could be due to the cooling and analgesic effects of ice on the skin and the subsequent numbing sensation and pain relief (1). On the other hand, the study by Farnia et al. demonstrated no significant difference in the effects of Emla cream and cold compress on the intensity of the pain induced by enoxaparin injection. Considering the cost-effectiveness of cold compress, the use of this method is suggested for pain relief (13).

### 5.1. Limitations of the Study

As pain is a mental concept and its expression varies in individuals based on physiological, individual, and cultural characteristics, optimal accuracy and reliability cannot be achieved in pain assessment.

### 5.2. Conclusion

According to the results, the local cold compress could reduce the pain intensity and injuries caused by the injection. In order to improve the quality of care and minimize the pain experienced by patients, our findings could be applied as a guide to decrease the adverse effects of subcutaneous enoxaparin injection and also provide training to caregivers and therapists. In addition, the incorporation of these results into clinical nursing may increase the quality of nursing practice and enhance patient satisfaction and cooperation.

**Table 3.** Comparison of Mean Pain Intensity in Five Study Groups <sup>a, b, c</sup>

Groups	N	Minimum	Maximum	Mean ± SD
Control	20	0	4	1.40 ± 1.60
Five minutes before injection	20	0	5	1.15 ± 1.49
Twenty minutes after injection	20	0	7	1.05 ± 2.03
Five minutes before injection and 20 minutes after injection	20	0	2	0.6 ± 0.88
Five minutes before injection and 5 minutes after injection	20	0	2	0.2 ± 0.61

<sup>a</sup> Significant difference in mean pain intensity in five study groups (significance = 0.044; F = 2.255).

<sup>b</sup> Null hypothesis representing equality of pain intensity in five groups rejected.

<sup>c</sup> Significant difference observed in mean pain intensity in study groups.

## Acknowledgments

Hereby, we extend our gratitude to the Deputy of Research and Technology of Islamic Azad University (Babol Branch, Iran) and all the patients for assisting us in this research project.

## Footnotes

**Authors' Contribution:** Study concept and design, Fatemeh Taghlili; Acquisition of data, Maryam Ebrahimi Zavardeh; Analysis and interpretation of data, Fatemeh Taghlili; Drafting of the manuscript, Fatemeh Taghlili; Statistical analysis, Azadeh Kiapour; Administrative, technical, and material support, Fatemeh Taghlili; Study supervision, Fatemeh Taghlili.

**Clinical Trial Registration Code:** This study has been registered in the Iranian Registry of Clinical Trials (code: IRCT20180827040884N2).

**Conflict of Interests:** There is no conflict of interest.

**Ethical Approval:** The study protocol was approved by the Research Deputy (design number: 1569505230001) and hospital security. Before the study, the objectives of the research were explained to the participants, and they were assured of the safety of the procedures. Moreover, written informed consent was obtained from the subjects.

**Funding/Support:** Islamic Azad University (Babol Branch, Iran)

**Informed Consent:** Written informed consent was obtained from the subjects.

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