Comparison of the effect of ginger capsule and diclofenac tablet on the relief of pain following inguinal hernia surgery

Gholamreza Shabanian¹*, Ali Satari²

¹Department of Anesthesia, Shahrekord University of Medical Sciences, Shahrekord, Iran
²Student Research Committee, Shahrekord University of Medical Sciences, Shahrekord, Iran

*Corresponding Author: Gholamreza Shabanian, Department of Anesthesia, Ayatollah Kashani Hospital, Shahrekord University of Medical Sciences, Iran, Tel: +989133827110, E-mail: gshabanian@yahoo.com

Abstract
Background and aims: Postoperative pain control decreases negative effects on the health domains of the patients and their relatives in addition to reducing the costs and the duration of hospitalization. The aim of the present study was to comparatively investigate the analgesic effects of ginger and diclofenac tablets.

Materials and Methods: In this clinical trial, 80 patients undergoing inguinal hernia surgery were randomly divided into two groups of 40. One group received 250 mg ginger capsule while the other one received 100 mg diclofenac tablet. Then, the patients were examined within 6 hours (every two hours) and their pain was assessed using the standard 10-point visual analogue scale. Finally, the data were analyzed using the SPSS, version 22.

Results: Based on the results, there was no significant difference in pain severity between the two groups at 0, 2, 4, and 6 hours after recovery (P > 0.05). In addition, no significant difference was observed in the duration of pain since the time of patient recovery between the two groups (P > 0.05). Further, as regards the number of patients receiving pethidine, no difference was found between the two groups (P > 0.05).

Conclusion: In general, both ginger capsule and diclofenac tablet were similarly effective in relieving the pain after inguinal hernia surgery.

Keywords: Inguinal hernia, Ginger, Diclofenac, Pain

Introduction
Pain is one of the major public health issues around the world that imposes heavy economic burden on the community (1-4). In addition to bothering the patient, pain can impair the quality of life and thus adversely affect the family and social aspects of the patients’ lives (5). Therefore, pain relief in patients reduces the adverse effects of pain on the patient’s physical and mental conditions and provides comfort for his/her family (6). Postoperative pain relief, especially through certain regimens, can decrease mortality during and after the operation (7,8). Pain control with the least possible complications, as one of the dimensions of patient satisfaction with health care services, is one of the goals of pain management (6). There are several ways to reduce pain in patients. For example, nonsteroidal anti-inflammatory drugs and opioids can be used to treat various pains, including surgical pain (9).

Meanwhile, herbal remedies have drawn attention because of their cost-effectiveness and fewer side effects in long-term use, including chronic pains (10-14). Plants can even be used as food supplements and be effective in reducing the pain by inducing anti-inflammatory properties (15-18). Zingiber officinale rhizome is one of the oldest medicinal and preventive remedies for many conditions, including pain (19-23). However, despite the anti-inflammatory effects of ginger, its effect on pain relief is controversial (24). Therefore, the present study was conducted to compare the effects of the ginger capsule and diclofenac tablet on the relief of pain following inguinal hernia surgery.

Materials and Methods
In this prospective double-blind clinical trial, a total of 80 patients submitted to inguinal hernia repair surgery in Kashani hospital of Shahrekord were selected by convenience sampling technique. The sample size was calculated at 40 for each group using a sample size statistics formula and according to the protocol of a similar study (25). The inclusion criteria were those patients aged over 14 years who were eligible for inguinal hernia repair surgery and provided consent to participate in the study. In addition, the exclusion criteria included over 1-hour duration of surgery, classified as higher than ASA (American Association of Analgesia) II, a history of cardiovascular, liver, kidney, pulmonary, as well as psychiatric diseases, epilepsy, and diseases affecting the function of the nervous
and muscular system, addicted or alcoholic or the habitual use of any related drug, emergency surgery, gastrointestinal problems, the unintentional division of ilioinguinal nerve and/or iliohypogastric nerve during the surgery, receiving analgesics 24 hours before surgery, asthma, eczema or hypersensitivity to ginger, recurrent hernia, neurological lesions such as stroke, nerve division due to trauma and congenital neurological diseases, and the lack of ability to communicate verbally and express the pain. The data collection instrument was a checklist, the first section of which consisted of demographic characteristics such as age, a history of underlying diseases, hospitalization and surgery records, associated symptoms including nausea and vomiting, as well as postoperative complications (up to 10 days) including infection and hematoma after taking the studied drugs. The second section addressed pain severity and the duration of postoperative pain that was measured by a standard 10-point visual analogue scale (26). Anesthesia was recorded on a 10-point (1-10) scale, with the same visual analogue scale form used at all intervals, and therefore the patient would select the choice that indicated the severity of his/her pain. Before entering the operating room, patients received no pretreatment and were divided into two groups.

In order to control postoperative pain, the patients in the ginger group received one ginger capsule (Zintoma) orally with 50 mL water one hour before surgery, while the diclofenac group received diclofenac 100 mg with 50 mL water. Further, both groups received 2 mg midazolam one hour before the surgery. In recovery, pulse oximetry was monitored and oxygen was given via the facial mask. The first examination of the pain was conducted within 6 hours (once every two hours) after complete recovery in the recovery room and after the patient was transferred to the department. Furthermore, the pain severity was assessed by the visual analogue scale criteria while the incidence of vomiting, respiratory depression, urinary retention, shivering, and hemodynamic changes, as well as the amount and type of injectable drug were monitored by a contributor to the research project who was blind to the type of the drug prescribed for the patient. The duration of the pain for all patients was considered to begin from the time the patient recovered to when the pain severity decreased to less than three.

In addition, 25 mg of pethidine was injected to the patient if requested or needed (pain score higher than 5) and the number of injections of pethidine was recorded. The obtained data were analyzed by SPSS, version 22. The frequency and relative frequency were used for qualitative variables and mean (standard deviation) was employed for quantitative variables. Moreover, qualitative variables were analyzed by chi-squared and Fisher exact tests and the mean values in the two groups were compared by using Student’s t test or its nonparametric equivalent (Mann-Whitney test) depending on data distribution.

$P<0.05$ was considered statistically significant.

**Results**

The mean age of the participants in the ginger and diclofenac groups was 41.47±14.44 and 44.22±13.22, respectively, with no statistically significant difference ($P>0.05$). Additionally, the age range in all participants was 16-71 years and 41.7750±14.3574 and 44.025±13.3002 years in the ginger and diclofenac groups, respectively. The results of this study showed that there was no significant difference regarding the diseases between the two groups ($P>0.05$). It was not possible to investigate this issue because no side effects such as infection, vomiting, and the like were observed in the patients.

Table 1 demonstrates the results related to pain severity among the groups receiving ginger and diclofenac. Based on the results, no significant difference was found regarding pain severity between the two groups at 0, 2, 4, and 6 hours after the recovery ($P>0.05$). In addition, the results revealed no significant difference in the duration between recovery and pain improvement between the two groups ($P>0.05$).

The duration of pain in the two groups is shown in Table 2. Although there was no significant difference between the two groups, most patients experienced pain for 3 hours.

Twenty-five milligrams of pethidine was injected to the patient with a score of higher than 5 or on his/her own request. Accordingly, about 0.12 and 0.1 of the patients in the ginger and diclofenac groups, respectively, received pethidine on their own request or because of the pain score above 5, with no statistically significant difference ($P>0.05$).

**Discussion**

This study aimed at comparing the effects of ginger capsule and diclofenac tablet on pain relief after inguinal hernia surgery. The severity of pain was not significantly different between the two groups at different intervals after

<table>
<thead>
<tr>
<th>Time of study</th>
<th>Ginger capsule</th>
<th>Diclofenac</th>
<th>$P$ value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Recovery</td>
<td>5.02±1.44</td>
<td>4.82±1.33</td>
<td>0.552</td>
</tr>
<tr>
<td>2 h after recovery</td>
<td>3.17±1.12</td>
<td>3.05±1.03</td>
<td>0.608</td>
</tr>
<tr>
<td>4 h after recovery</td>
<td>1.52±1.10</td>
<td>1.4±1.03</td>
<td>0.603</td>
</tr>
<tr>
<td>6 h after recovery</td>
<td>0.27±0.84</td>
<td>0.25±0.77</td>
<td>0.893</td>
</tr>
</tbody>
</table>

**Table 2.** Duration of the pain in the groups receiving ginger capsule and diclofenac tablet

<table>
<thead>
<tr>
<th>Group</th>
<th>Ginger capsule</th>
<th>Diclofenac</th>
<th>$P$ value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Duration of pain (h)</td>
<td>5.75±1.12</td>
<td>5.70±1.15</td>
<td>0.845</td>
</tr>
</tbody>
</table>
the recovery. Further, the pain between the recovery and pain improvement failed to significantly differ between the two groups. Consistent with these results, the positive effects of ginger or its derivatives on various types of pains were confirmed by other studies, including those of Yip et al on knee pain (27), Chen et al on dysmenorrhea (28), Black et al on muscle pain (29), Al-Nahain et al on pain caused by rheumatoid arthritis (30), and Leach et al on osteoarthritis (31).

However, a review article reported that despite the anti-inflammatory effects of ginger, its effect on pain relief remains to be elucidated (24). This can be due to the differences in the methodology of other studies. Additionally, the type of the mental experience of pain, the culture, and the type of pain measurement scale could lead to inconsistency in the findings.

Ginger rhizome has been used for many human diseases such as gastrointestinal in various parts of the world since the ancient times, and its medicinal properties have been often attributed to the presence of phenolic compounds, along with its antioxidant and anti-inflammatory properties (32,33). By containing active ingredients such as gingerol, ginger can reduce inflammation and thus relieve the pain (34). Anthocyanins are glycoside polyhydroxyl derivatives and methoxy, 2-phenylbenzopyrylium salts, as well as non-toxic and water-soluble pigments that are widely found in nature. The red, blue, purple, amethystine, and black colors in many fruits, vegetables, and flowers are due to the presence of anthocyanins. These compounds are highly potent antioxidants that can suppress the pathways related to the pain by inhibiting cyclooxygenase and lipoxygenase. There are phenolic compounds and anthocyanins such as gingerdion, shaogol, and gingerol in ginger (35). These three compounds are strong prostaglandin inhibitors that prevent the metabolism of arachidonic acid by inhibiting cyclooxygenase and lipoxygenase. In addition, these compounds can impede inflammatory pathways by inhibiting nitric oxide synthase and thus exert their analgesic effects (36,37).

**Conclusion**

Overall, no significant difference was observed between ginger and diclofenac in postoperative pain relief, which indicates the acceptable analgesic effect of ginger. Besides, considering the dose of ginger in this study, it can be concluded that higher doses of ginger can even exhibit a greater analgesic effect compared to those of diclofenac if they cause no side effects. In this study, there was no specific side effect in the group receiving ginger and its analgesic effect was similar to that of diclofenac. Therefore, it seems that ginger supplementation can be used to relieve pain with comparatively fewer side effects.

**Conflict of Interests**

The authors have no conflict of interests.

**Ethical considerations**

This article was derived from a research project approved by the Research and Technology Deputy of the Shahrekord University of Medical Sciences (approval no.: 1964). Hereby, the authors gratefully thank the women who participated in this study.

**References**

17. Maroon JC, Bost JW, Maroon A. Natural anti-inflammatory
agents for pain relief. Surg Neurol Int. 2010;1:80. doi:
10.4103/2152-7806.73804.
18. Harrison AM, Heritier F, Childs BG, Bostwick JM, Dzidzako
MA. Systematic review of the use of phytochemicals for
22. Rahmani AH, Shabrami FM, Aly SM. Active ingredients of
ginger as potential candidates in the prevention and treatment of
diseases via modulation of biological activities. Int J Physiol
23. Rahnama P, Montazeri A, Huseini HF, Kianbakht S, Naseri M.
Effect of Zingiber officinale R. rhizomes (ginger) on pain relief
in primary dysmenorrhea: a placebo randomized trial. BMC
6882-12-92.
(Zingiber officinale) for the treatment of pain: a systematic
25. Noroozinia H, Mahoori A, Hassani E, Akhbari P. Diclofenac
suppository versus intramuscular pethidine in post
herniorrhaphy pain relief. Tehran University Medical Journal.
2011;69(3):198-203.
26. Kersten P, White PJ, Tennant A. Is the pain visual analogue
scale linear and responsive to change? An exploration using
journal.pone.0099485.
27. Yip YB, Tam AC. An experimental study on the effectiveness
of massage with aromatic ginger and orange essential oil
for moderate-to-severe knee pain among the elderly in
28. Chen CX, Barrett B, Kwekkeboom KL. Efficacy of oral ginger
(zingiber officinale) for dysmenorrhea: a systematic review
and meta-analysis. Evid Based Complement Altern Med.
29. Black CD, Herring MF, Hurley DJ, O'Connor PJ. Ginger
(Zingiber officinale) reduces muscle pain caused by eccentric
30. Al-Nahain A, Jahan R, Rahmatullah M. Zingiber officinale:
A potential plant against rheumatoid arthritis. Arthritis.
31. Leach MJ, Kumar S. The clinical effectiveness of Ginger
(Zingiber officinale) in adults with osteoarthritis. Int J Evid
Based Healthc. 2008;6(3):311-20. doi: 10.1111/j.1744-
1609.2008.00106.x.
32. Shukla V, Singh M. Cancer preventive properties of ginger:
a brief review. Food Chem Toxicol. 2007;45(5):683-90. doi:
33. Rafieian-Kopaei M, Nasri H. The Ameliorative Effect of Zingiber
officinale in Diabetic Nephropathy. Iran Red Crescent Med J.
34. Wilson PB. Ginger (Zingiber officinale) as an analgesic and
35. Ghaseemzadeh A, Jaafar HZ, Rahmat A. Identification and
concentration of some flavonoid components in Malaysian
young ginger (Zingiber officinale Roscoe) varieties by a high
performance liquid chromatography method. Molecules.
36. Aktan F, Henness S, Tran VH, Duke CC, Routogalis BD, Ammit
AJ. Gingerol metabolite and a synthetic analogue Capsarol
inhibit macrophage NF-kappaB-mediated iNOS gene
expression and enzyme activity. Planta Med. 2006;72(8):727-
al. In vitro antioxidant and anti-inflammatory activities of
1-dehydro-[6]-gingerdione, 6-shogaol, 6-dehydroshogaol and