Effect of Intraperitoneal Curcumin Instillation on Postoperative Peritoneal Adhesions

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Abstract
Objective: The aim of this study was to determine the effect of curcumin on adhesion formation in a rat cecum abrasion model.

Materials and Methods: Thirty Wistar rats were randomized into three groups; the control group received saline, the curcumin group received 10 mg/kg of curcumin after cecal abrasion, and in the sham group the abdominal wall was closed without any abrasion to the cecum. On day 15, adhesions were assessed blindly using a standardized scale, and histopathological samples were taken and examined.

Results: There were no incisional hernias or wound dehiscences in any animals of the three groups. A comparison of adhesion scores showed a significant difference between the curcumin (median = 1) and the control group (median = 2; p < 0.05). The grade of inflammation of the curcumin (median = 1) and the sham (median = 0) group was significantly lower than that of the control group (median = 3; p < 0.01 and p < 0.001, respectively). Hydroxyproline levels were significantly lower in the sham (48.3 ± 11.8 μg/mg) and the curcumin (63.8 ± 13.9 μg/mg) group compared to the control group (85.7 ± 22.1 μg/mg; p < 0.05).

Conclusion: These data suggest that curcumin, administered intraperitoneally, was effective in the prevention of peritoneal adhesion formation.

Introduction

Intraperitoneal adhesions are a common worldwide problem that causes chronic abdominal pain, dyspareunia, infertility, postoperative ileus and medico-legal issues [1]. The risk for the development of intraperitoneal adhesions after abdominal surgery is reported to be over 90% [1, 2]. Postoperative adhesions are the most common cause of small bowel obstruction, responsible for approximately 70–80% of cases, and are associated with high morbidity and mortality [1, 2]. While conservative treatment is the primary approach for postoperative ileus, about half of the patients require surgery [2]. Surgical intervention is not without its own complications; after adhesiolysis has been performed, adhesions tend to reform and are associated with an increased morbidity rate and a high risk of relapse of bowel obstruction [1, 2].
The pathogenesis of intraperitoneal adhesion formation is still not well understood, and there is no specific treatment. However, the suggestion has been made that inflammatory cells, fibroblasts, the endothelium and the mesothelium of the peritoneal serosa, as well as other factors, are in a complex interplay in the pathogenesis of adhesion formation [3]. Insufficient fibrinolysis is a widely accepted major factor in peritoneal adhesion formation [4], which inevitably occurs if fibrinolysis is insufficient. The two main factors that convert plasminogen into active plasmin, that is tissue plasminogen and urokinase-like plasminogen, are major activators in the fibrinolytic system [4].

Inflammatory mediators also play an important role in adhesion formation. Factors such as transforming growth factor and interleukins decrease the fibrinolytic capacity of the tissue and increase the formation of adhesions [5, 6]. Vascular endothelial growth factor (VEGF), recently found to be expressed by mast cells, has multiple effects on several crucial mechanisms in adhesion formation [7]. VEGF is a potent angiogenic cytokine and has been demonstrated to play a role in early inflammatory responses, wound repair and deposition of fibrinogen [7]. Some medications such as bevacizumab that inhibit the biological activity of human VEGF have been shown to reduce abdominal adhesions [8].

Various material-based strategies are clinically used to cover damaged peritoneal surfaces [9]. Besides, many drug-like agents, including antioxidants, anti-inflammatory agents, antibiotics, anticoagulants and fibrinolytics, have been shown to prevent the formation of adhesions [7, 10]. Curcumin, a yellow pigment extracted from the Indian spice tumeric, has been reported to possess pleiotropic activities, antibiotics, anticoagulants and fibrinolytics, have been shown to prevent the formation of adhesions [7, 10]. Curcumin, a yellow pigment extracted from the Indian spice tumeric, has been reported to possess pleiotropic effects as an antioxidant as well as to have anti-inflammatory, anticarcinogenic, antibacterial, antiviral, antifungal, antiproliferative and proapoptotic effects [11]. Curcumin has also been shown to increase fibrinolytic activity and cell migration towards the wound area by modulating urokinase-like plasminogen expression [11]. Moreover, curcumin has been shown to inhibit both VEGF secretion and VEGF-mediated angiogenesis [12]. Due to these aforementioned effects, we hypothesize that curcumin could play a role in the prevention of postoperative adhesion formation.

Materials and Methods

The study was conducted after having obtained approval from the Ethics Committee at Dicle University School of Medicine. The animal care was in accordance with institutional guidelines.

Materials

The mixture of curcumin was purchased from Sigma Aldrich (C7727; St. Louis, Mo., USA) and was dissolved in dimethyl sulfoxide (1 mg/ml) in brown glass vials for storage at 4°C.

Animals

Thirty male Wistar albino rats weighing 200–250 g were used. The animals were kept in cages under standard conditions with a balanced pelleted diet and water. The cages were changed twice a week and the water was changed once a day by experienced staff. The animals were acclimatized for 1 week before the experiments and were kept on a 12-hour light/dark cycle at a constant room temperature and humidity. After adaptation, they were randomly assigned to three different groups: the control, sham and curcumin groups, each including 10 rats.

Anesthesia and Surgical Procedure

The animals were fasted the night before surgery and then anesthetized with an intramuscular injection of 70 mg/kg of ketamine (Ketalar; Eczacıbaşı, Istanbul, Turkey). When anesthesia was achieved, the surgical procedures were performed under sterile conditions. Next, a minilaparotomy was performed to gain access to the abdominal cavity. To create a cecal adhesion model, a sterile surgical brush was used to obtain a bleeding surface on the antimesenteric side of the cecum and parietal peritoneum. The peritoneum, fascia and skin were then closed with 3/0 absorbable sutures.

In the sham operation group, the abdominal wall was closed without any abrasion to the cecum. In the control group, the cecal and peritoneal abrasion was conducted and 0.2 ml of saline was administered intraperitoneally before closing the abdomen. In the curcumin group, after cecal and peritoneal abrasion, 10 mg/kg of curcumin was administrated intraperitoneally before closing the abdomen.

Adhesion Assessment

On the 15th day of the study, all the animals were anesthetized with an injection of ketamine, and they were euthanized by intracardiac blood collection. A repeat laparotomy was performed to evaluate adhesion formation in the three groups. The cecum and the abdominal sidewall were evaluated for adhesion formation according to the classification of Mazuji et al. [13]. The cecum and adherent abdominal wall were then sent for histopathological evaluation.

Histopathological Evaluation

Peritoneal healing and fibrosis were quantitated by the measurement of hydroxyproline tissue levels. After fixation of the adhesion, specimens were buffered in formalin (10%) and were dehydrated and embedded in a paraffin wax. Serial sections of 5 μm were stained with hematoxylin and eosin (HE).

The samples were histopathologically examined under light microscopy by a pathologist blinded to the study. A semiquantitative scoring system of the histopathological grading scale of Zühlke was used for the evaluation of the inflammation. Grade I was defined as weak connective tissue, rich cells, old and new fibrin and thin reticulin fibers; grade II was defined as connective tissue with cells and capillaries, and few collagen fibers; grade III was characterized by firmer connective tissue, fewer cells, more vessels and few elastic and smooth-muscle fibers, and grade IV was determined by old, firm granulation tissue which was cell poor and se Rosal layers which were hardly distinguishable [14].
Effect of Curcumin on Postoperative Adhesions

Determination of Tissue Levels of Hydroxyproline

Because collagen plays an important role in the development of postoperative adhesions and hydroxyproline is a major component of the protein collagen, the extent of collagen deposition in the adhesion tissues was determined using the hydroxyproline analysis. From an adhesion, 1 g of tissue was hydrolyzed in 1 ml of acidic buffer at 121 °C for 5 h. Then, the tissue was centrifuged at 5,000 rpm for 20 min. The absorbance of this material was evaluated colorimetrically (spectrometrically), with a wavelength maximum of 560 nm and at 121 °C. Hydroxyproline levels were calculated in micrograms per milligram of tissue.

Statistical Analyses

Statistical evaluation was made using MedCalc software version 12.4.0. The Mann-Whitney U test was used to evaluate continuous variables. Two-sided p values are given and p < 0.05 was considered to be the limit of significance.

Results

All animals completed the study. No congenital adhesions were noted at the initial laparotomy. There were no incisional hernias or wound dehiscences present.

Macroscopic Adhesion Scores

The adhesion scores for each group are shown in table 1. On comparing adhesion scores, a significant difference was found between the curcumin and the control groups (p < 0.05). However, the curcumin and sham groups (p > 0.05) did not show any significant differences (p > 0.05). An example of abdominal adhesion tissue in the control group is shown in figure 1.

Histopathological Results

The Zühlke scores were significantly lower in the curcumin (median = 1) and sham (median = 0) groups compared to the control group (median = 3; p < 0.01 and p < 0.001, respectively; fig. 2, 3). There were 6 animals in the control group with a Zühlke grade 3 score whilst the other groups had no animals with a grade 3 score. Six animals in the curcumin group had grade 1 and the others had grade 2 scores. Six animals in the sham group had grade 0 and the others had grade 1 scores. The sham group (me-

Table 1. Distribution of adhesion scores and tissue hydroxyproline levels

<table>
<thead>
<tr>
<th>Adhesion scores</th>
<th>Sham group</th>
<th>Control group</th>
<th>Curcumin group</th>
</tr>
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<tbody>
<tr>
<td>Grade 0</td>
<td>7</td>
<td>0</td>
<td>3</td>
</tr>
<tr>
<td>Grade 1</td>
<td>3</td>
<td>4</td>
<td>5</td>
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<tr>
<td>Grade 2</td>
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<td>4</td>
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<td>Grade 3</td>
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<td>Grade 5</td>
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Hydroxyproline levels 48.3 ± 11.8 85.7 ± 22.1 63.8 ± 13.9

Data are either number (n = 10 in each group) or mean ± SD (μg/mg). Significant differences (p < 0.05) were found for the adhesion scores between the curcumin and control groups and for the hydroxyproline levels of the sham and curcumin groups compared to the control group.

Fig. 1. A rat from the control group with peritoneal adhesion formation.

Fig. 2. The histopathological scores in each group: those of the curcumin and sham groups were significantly lower than those of the control group (p = 0.0016 and p = 0.0001, respectively).
dian = 0) had significantly lower Zühlke scores compared to the curcumin group (median = 1; p < 0.01). Microscopic findings were similar to macroscopic findings.

**Quantitative Analysis of Hydroxyproline**

Because collagen plays an important role in the development of postoperative adhesions and hydroxyproline is a major component of the protein collagen, the extent of collagen deposition in the adhesion tissues was determined using hydroxyproline analysis. Hydroxyproline levels were significantly lower in the sham (48.3 ± 11.8 μg/mg) and curcumin groups (63.8 ± 13.9 μg/mg) compared to the control group (85.7 ± 22.1 μg/mg; p < 0.05; table 1).

**Discussion**

Both surgeons’ adhesion scorings and the histopathological grading showed significantly low adhesion scores in the curcumin group in comparison to the control group. Hence, the present study showed that curcumin effectively decreased the experimental postsurgical abdominal adhesion rate. Hydroxyproline levels were significantly lower in the curcumin group compared to the control group, thereby indicating that intraperitoneally administered curcumin decreased collagen synthesis. In some patients peritoneal tissues have a high tendency to develop adhesions, probably due to a reduced ability of the peritoneum to degrade fibrin deposits. It is widely accepted that the early fibrinolytic capacity plays an important role in the formation of adhesions [15]. In the first few days after peritoneal injury, fibrinolysis prevents adhesion formation during healing of the peritoneum. However, if the fibrinolysis process is ineffective, adhesion formation is inevitable. Fibrinolysis is carried out by the transition of plasminogen to plasmin, an enzyme with a broad proteolytic activity. In humans, it has been shown that peritoneal plasminogen activation is decreased during surgery, leading to subsequent adhesion formation [15]. Madhyastha et al. [11] demonstrated that curcumin treatment resulted in an increase in fibrinolytic activity around the wound area via an upregulation of urokinase-like plasminogen activator mRNA and protein.

Since VEGF has been shown to play a central role in the formation of postoperative intra-abdominal adhesions,
anti-VEGF monoclonal antibodies have been suggested for antiadhesion treatment. Ignjatovic et al. [8] stated that a single intraperitoneal dose of bevacizumab, a recombinant humanized monoclonal antibody that binds to and inhibits the biological activity of human VEGF, diminished both the grade and severity of peritoneal adhesions in a rat adhesion model. Fortunately, this therapy did not disrupt wound healing in a clinically important manner. Binion et al. [16] demonstrated that COX-2 plays an important role in VEGF-induced angiogenesis and curcumin blocks both COX-2 expression and angiogenesis induced by VEGF. They suggested that these findings may provide an understanding of the beneficial effects of curcumin in conditions such as chronic inflammation and cancer. Gururaj et al. [17] demonstrated a time-dependent response of curcumin in VEGF inhibition in endothelial cells. Similarly, curcumin has been shown to inhibit the transcript levels of VEGF in breast cancer cells [18]. Moreover, Bae et al. [19] found that curcumin suppressed transcriptional activity of HIF-1α, leading to a decrease in the expression of VEGF.

An important question that remains to be answered about intraperitoneal curcumin administration is whether or not it impairs wound healing. In our study, no wound dehiscences or abdominal wall hernias were seen in the groups, showing that administration of curcumin intraperitoneally does not disrupt wound healing in a clinically important fashion. Several studies have shown the nontoxic nature and beneficial effects of curcumin as a wound-healing agent [20, 21].

Jomezadeh et al. [22] claimed that curcumin was not effective in reducing postoperative peritoneal adhesion in rats. However, they performed relaparotomy on the 5th postoperative day to evaluate adhesion formation. Within the first 5 days after surgery, early fibrinolysis encourages peritoneal healing. If enough fibrinolysis does not occur within 5–7 days of the injury, the temporary fibrin matrix gradually becomes more organized and leads to adhesion formation. Therefore, adhesion formation is best evaluated after 7–10 days of the initial surgery [4, 8]. In our opinion, the 5th postoperative day is a very early point at which to evaluate adhesion formation. Therefore, we performed a second laparotomy 15 days after the first operation.

**Conclusion**

The present study shows that curcumin appeared to be safe and effective in the prevention of postoperative intra-abdominal adhesion formation in the rat. Due to the nontoxic nature of curcumin, we suggest that it could be used for the prevention of peritoneal adhesion formation. However, further studies are needed to determine the most suitable intraperitoneal dose of curcumin for peritoneal adhesions.

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**Disclosure Statement**

The authors have no conflicts of interest to declare.

**References**


