Lipoic acid decreases peritoneal adhesion formation in a rat uterine scar model

Abstract

Objective: To investigate the effects of lipoic acid in the prevention of postoperative pelvic adhesions by a visual scoring system and immunohistochemistry in a rat uterine horn model with full thickness injury.

Material and Methods: Twenty-eight female Wistar albino rats were randomised into four groups: uterine trauma control, 15 days and 30 days, and uterine trauma + lipoic acid, 15 days and 30 days. A full thickness defect was established by incising a segment of approximately 1.0 cm in length from each uterine horn, leaving the mesometrium intact. Extension and severity of the adhesions in each group were scored by a visual scoring system and evaluated immunohistochemically.

Results: Adhesion scores were 2.00±0.81, 2.14±0.69, 0.71±0.75, and 0.85±0.69 for extent and 2.28±0.48, 2.14±0.69, 0.85±0.69, and 1.14±0.69 for severity in Groups 1, 2, 3 and 4, respectively. Adhesion extent and severity were significantly less for groups treated by lipoic acid but no difference was observed between long and short interventions and the risk of intestinal injury, haemorrhage and inadequate site exposure is increased. Minimalising tissue trauma, the avoidance of foreign materials and prophylaxis for infection are common measures against adhesion formation. There are numerous studies on developing agents such as barrier materials, hormones and their agonist/antagonists, hyaluronic acid, fibrinolytic agents, non-steroidal anti-inflammatory drugs and antioxidants to prevent postoperative adhesion formation (3-7).

Conclusion: Lipoic acid was found to be effective in reducing postoperative adhesion formation in a rat model.

Key words: Lipoic acid, adhesion, rat, uterine scar, full thickness uterine injury

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Introduction

The process of incision, cauterisation and suturing during surgery inevitably results in tissue healing and postoperative peritoneal adhesions, which may lead to female infertility, chronic abdominal pain or bowel obstruction (1, 2). Safe surgical entry into the abdomen is difficult in subsequent interventions and the risk of intestinal injury, haemorrhage and inadequate site exposure is increased. Minimalising tissue trauma, the avoidance of foreign materials and prophylaxis for infection are common measures against adhesion formation. There are numerous studies on developing agents such as
peritoneal adhesions. Adhesion formation following uterine scarring in animal experiments has not been reported to date. In this study, we used a rat model resembling myomectomy or caesarean section with uterine scarring and suturing and tested the effect of Lipoic Acid (LA) on postoperative peritoneal adhesion formation.

Free radicals, namely superoxides, peroxides and hydroxyl radicals, are mediators of inflammation inducing adhesions by cellular membrane damage. Antioxidants such as methylene blue, vitamin E and N-acetyl cysteine have been reported to decrease development of peritoneal adhesions (7, 11, 12). There are few studies to our knowledge investigating the effect of LA in the prevention of adhesion formation. In this study, we aim to evaluate the effects of LA in the prevention of postoperative pelvic adhesions. To rule out subjective evaluation we have used both a visual scoring system and immunohistochemically we have used the wound healing markers, urokinase plasminogen activator (u-PAR) and vitronectin in a rat uterine horn model with full thickness injury of the myometrium.

Material and Methods

Twenty-eight female, non-pregnant Wistar albino rats were used; all experiments were performed in accordance with the guidelines provided by the Experimental Animal Laboratory and approved by the Animal Care and Use Committee of Dokuz Eylul University Faculty of Medicine. The rats weighed approximately 200-250 g, were housed three animals to a cage under standard laboratory conditions with a day cycle of 14 hours light and had free access to food and water. The rats were randomly assigned to one of four study groups: Group 1, uterine scar group (15 days) (n=7); Group 2, uterine scar group (30 days) (n=7); Group 3, uterine scar+LA therapy (15 days) (n=7); and Group 4, uterine scar+LA (30 days) (n=7).

To standardise the hormonal status of the rats, menstrual cycle was determined by vaginal smear and the experiment was done on day of the dioestrus phase. Experimental design was modified the full-thickness injury model described in 2012 by Lin et al. (13) All rats were anaesthetised by intraperitoneal injection of ketamine and xylazine (35 mg/kg). The first incision was made in the abdominal wall of each rat under sterilised conditions. Then a full-thickness defect was created by incising a segment of approximately 1.0 cm in length from each uterine horn, leaving the mesometrium intact (Figure 1). The margins of the uterine defect were marked with a 4-0 nylon line. The abdominal incision was closed in two layers with a monofilament 3/0 polyglactin suture for the peritoneum and 2/0 polyglactin suture for the skin. The operation time did not exceed 15 minutes and all animals recovered without any complications or infections. All animals were treated with an intramuscular injection of penicillin (80,000 units/100 mg) for 3 days after the surgery.

Alpha lipoic acid (Sigma, St Louis, MO, USA) was prepared by mixing 100 mg/kg with sterile saline in a dark bottle and adding 1 M NaOH until the suspension dissolved. The pH was adjusted to 7.4 by adding 1 M HCl. Fresh LA solution was administered by oral gavage for either 15 or 30 days after uterine scarring. After 15 or 30 days according to the study groups, the animals were anaesthetised, relaparatomy was performed, the extent and severity of intraabdominal adhesions were recorded and animals were sacrificed. All uterine horns of each rat in all study groups were evaluated separately (total 14 horns).

An author blinded to the medication status of the rats performed the visual assessment of adhesions. A published scoring system was used (14). The extent was evaluated as 0 for no adhesions, 1 for 25% of adhesions of the traumatised area, 2 for 50% of adhesions of the traumatised area and 3 for total involvement. The severity scores were 0 for no resistance to separation, 1 when minimal dissection was required and 3 for sharp dissection.

Uterine horns were fixed in neutral formalin fluid, dehydrated in graded series with ethanol and water, and embedded in paraffin. Serial sections 5 microns thick were collected on slides. For light microscopy evaluation, haematoxylin eosin staining was performed. Streptavidin-biotin technique was used for immunostaining with u-PAR (rabbit anti-rat urokinase receptor IgG, 3920, American Diagnostica; 10 mg/mL concentration) and Vitronectin (ab45139, Abcam; 1/100 dilution). Following overnight incubation at 60°C, sections were dewaxed in xylene for 20 minutes. A decreasing series of ethanol was used for rehydration and then sections were washed in distilled water followed by phosphate-buffered saline (PBS) for 10 min each. Then, they were treated with trypsin (Cat No: 00-3008 Digest All 2A, Zymed, San Francisco, CA, USA) at 37°C for 15 min. To inhibit endogenous peroxidase activity, sections were delineated with a Dako pen (Dako, Glostrup, Denmark) and incubated in a solution of 3% H2O2 for 15 min. Sections were incubated with a blocking solution (Invitrogen, Histostain-Plus Broad Spectrum Cat No: 94526-20840-10068) for 30 minutes. After washing in PBS, the sections were incubated in a solution containing 1% H2O2 for 5 minutes to inhibit endogenous peroxidase activity. Sections were then incubated in a solution containing the primary antibody diluted 1/50 in PBS for 24 hours. Following washing in PBS, sections were incubated in a solution of biotinylated secondary antibody diluted 1/200 in PBS for 20 minutes. After washing in PBS, sections were incubated in a solution containing streptavidin-horseradish peroxidase for 20 minutes. After washing in PBS, sections were incubated in a solution containing diaminobenzidine (DAB) for 5 minutes. Sections were then counterstained in haematoxylin for 20 minutes. After washing in distilled water, sections were dehydrated in graded series of ethanol and xylene, and coverslipped.

Figure 1. a, b. Macroscopic appearance of adhesion areas  c. Haematoxylin-eosin evaluation of adhesion areas
positive stained cells among 100 cells in 5 fields randomly chosen in each horn were counted. A total of 500 cells were counted for each horn. Statistical analysis was done using SPSS 15.0 (SPSS Inc., Chicago, USA) and Kruskal-Wallis test was applied. Results were given as mean +/- standard deviation.

**Results**

The extent of adhesions was 2.00±0.81, 2.14±0.69, 0.71±0.75, and 0.85±0.69 for Groups 1, 2, 3 and 4, respectively. Adhesion extent was significantly less for lipoic acid groups. When comparing 15 days and 30 days of LA administration, there was no difference. The severity of adhesions was 2.28±0.48, 2.14±0.69, 0.85±0.69 and 1.14±0.69 for Groups 1, 2, 3 and 4, respectively. Severity of adhesions was significantly less in the LA groups, but no difference was observed between long and short administration (Table 1).

Both Vitronectin and u-PAR staining were significantly increased in LA groups when compared to the scar group. There was no significant difference between short and long LA application groups (Table 2, Figure 2).

**Discussion**

Postoperative pelvic adhesions may lead to complications such as extended operation time, additional blood loss and visceral damage (7) in cases of relaparatomy. Caesarean sections are the most frequently performed obstetrical operations worldwide and recurrent caesareans are difficult due to intraabdominal adhesions. Similarly, after a myomectomy there is a risk of peritoneal adhesion formation. In this study, we demonstrated that when applied orally, both short and long duration treatments with lipoic acid were effective in preventing pelvic adhesions following surgical trauma in rats. Tissue remodelling markers were used to verify the results. The incision site of the myometrium and locations of subsequent suturing are areas where wound healing takes place. Healing is a result of proliferation and regeneration of the mesothelial cell layer and fibrinolysis producing a peritoneal scar (15). Adhesions are primarily the result of this scar. An increase in oxidative stress and the formation of reactive oxygen species (ROS) play an important role in the pathophysiology of adhesion formation. Antioxidants such as methylene blue, vitamin E and N-acetyl cysteine have been reported to decrease development of peritoneal adhesions (7, 11, 12). Alpha LA and its metabolites are antioxidant and, when in contact with free radicals, oxidation takes place (16).

There are several experimental models for producing peritoneal adhesions in laboratory animals: the damaged uterine horn model by electrocautery or scraping, caecal abrasion, peritonitis model and the bowel anastomosis model (4, 17-20). The traumatisation of the uterine horn is widely used to mimic abdominal surgery; however, in the most frequent gynaecology

<table>
<thead>
<tr>
<th>Table 1. Extent and severity of adhesion</th>
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<tr>
<td><strong>Group</strong></td>
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<tr>
<td>Uterine Scar Group (15d)</td>
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<td>Uterine Scar Group (30d)</td>
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<tr>
<td>Uterine Scar Group+LA therapy (15d)</td>
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<td>Uterine Scar Group+LA therapy (30d)</td>
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Uterine Scar 15 d vs. Uterine Scar 30 d (p=0.72, p=0.70) for extent and severity; (*) Uterine Scar 15 d vs. Uterine Scar 15 d+LA (p=0.01, p=0.003) for extent and severity; (**) Uterine Scar 30 d vs. Uterine Scar 30 d+LA (p=0.009, p=0.024) for extent and severity, Uterine Scar 15 d+LA vs. Uterine Scar 30 d+LA (p=0.674, p=0.431) for extent and severity

**Table 2. Immunohistochemical staining of Vitronectin and u-PAR of adhesion tissue**

<table>
<thead>
<tr>
<th><strong>Group</strong></th>
<th><strong>Vitronectin</strong></th>
<th><strong>u-PAR</strong></th>
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<tbody>
<tr>
<td>Uterine Scar Group (15d)</td>
<td>11.42±2.50*</td>
<td>13.7±2.05*</td>
</tr>
<tr>
<td>Uterine Scar Group (30d)</td>
<td>21.43±6.57</td>
<td>19.4±3.86</td>
</tr>
<tr>
<td>Uterine Scar Group+LA therapy (15d)</td>
<td>39.14±9.51**</td>
<td>40.7±8.42**</td>
</tr>
<tr>
<td>Uterine Scar Group+LA therapy (30d)</td>
<td>42.14±8.98***</td>
<td>43.5±6.72***</td>
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(*) Uterine Scar 15 d vs. Uterine Scar 30 d (p=0.001, p=0.011) for Vitronectin and u-PAR immunostaining; (**) Uterine Scar 15 d vs. Uterine Scar 15 d+LA (p=0.001, p=0.001) Vitronectin and u-PAR immunostaining; (***) Uterine Scar 30 d vs. Uterine Scar 30 d+LA (p=0.004, p=0.001) for Vitronectin and u-PAR immunostaining, Uterine Scar 15 d+LA vs. Uterine Scar 30 d+LA (p=0.902, p=0.436) for Vitronectin and u-PAR immunostaining.
and obstetrics operations there is a full thickness cut through
the uterine wall. Therefore, we have chosen a new model to
form adhesions. We cut the uterine horns using full layer thick-
ness and then sutured the incision resembling a caesarean
section or myomectomy where the uterine cavity is exposed.
Tissue remodelling and a normal healing process require plas-
minogen activators to form plasmin which will play a role in
fibrinolysis. Insufficiency in fibrinolysis after surgery may lead to
fibrin deposition causing adhesions. Urokinase plasmin activa-
tor binds to its receptor u-PAR which was used in the present
study. Vitronectin on the other hand activates plasminogen
activators and integrins and the balance of these molecules is
important in regeneration (21, 22).
ROS are shown to be involved in adhesion formation after sur-
gery. There is an increase of free radical activity of superoxide
anions, xanthine oxidase and MDA (23, 24). The surgical area is
a local hypoxic environment leading to an ischemia/reperfusion
process resulting in a decrease of free radical scavenger levels.
Restoration of these free radical scavengers have been shown
to prevent adhesion formation in animal studies with induced
intestinal ischemia (25). During peritoneal healing, oxidative
stress increases and a positive correlation between the level
of oxidative stress and the severity of adhesions has been
demonstrated (26, 27). Postsurgical adhesion formation was
reduced by the administration of antioxidants such as vitamin
E, selenium or resveratrol in previous studies (5, 12, 28). Ozler
et al. (29) have shown the existence of oxidative stress in a rat
model with caecal trauma and a decrease after the application
of lipoic acid. Even though the method of adhesion formation
induction differs from our study, it supports our hypothesis that
LA is effective in preventing postsurgical adhesions.
Lipoic acid is used in the therapy of diabetes, atherosclerosis,
neurodegenerative processes, joint diseases or acquired immune
deficiency syndrome (16, 30). It has a low redox potential and
thus participates in reactions neutralising ROS, and reduces the
oxidised forms of other antioxidants. Another advantage of LA is
the fact that it is soluble in water and in fats (16). This is the first
experimental study that combines a new method for adhesion
formation, an antioxidant molecule (lipoic acid) and immunohis-
tochemical methods to evaluate the results objectively.
Most studies about postoperative adhesion formation and
prevention use visual evaluation methods and remain as sub-
jective results. The authors have contributed to the literature
on adhesion evaluation via an immunohistochemical method
using u-PAR and vitronectin markers for objective evaluation
(4). In the present study, this method is used again to confirm
the anti-adhesion effect of lipoic acid administration after surgi-
cal intervention mimicking caesarean section or myomectomy.
A distinctive increase in both tissue regeneration markers has
been observed in animals treated with LA. The present results
suggest a possible preventive effect of LA on postoperative
adhesions after full thickness uterine trauma with minimal side
effects and minimal cost.

**Ethics Committee Approval:** Ethics committee approval was
received for this study from the Animal Care and Use Committee
of Dokuz Eylul University Faculty of Medicine.

**Informed Consent:** N/A.

**Peer-review:** Externally peer-reviewed.

**Author contributions:** Concept - S.C.M., A.G.; Design - S.C.M,
Materials - S.C.M., O.S., P.A.; Data Collection&/or Processing -
Critical Reviews - O.S., B.U.E.

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**Figure 1. Staining of vitronectin and u-PAR in uterine scar groups of control and LA after 15 and 30 days (arrows)**
Conflict of Interest: No conflict of interest was declared by the authors.

Financial Disclosure: No financial disclosure was declared by the authors.

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