Effect of Curcumin on the Formation of Epidural Fibrosis in an Experimental Laminectomy Model in Rats

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ABSTRACT

AIM: To clarify the effects of topical application of curcumin on the prevention of epidural fibrosis.

MATERIAL and METHODS: Twenty-one rats were randomly divided into three equal groups (control, spongostan, local curcumin) and a laminectomy procedure was performed between T11 and L1 in all rats. Subsequently, spongostan soaked with curcumin (100 mg/kg) was applied topically. After four weeks, the vertebral column from T9 to L3, which included the paraspinal muscles and epidural scar tissue, was removed as a single piece and the epidural fibrosis and arachnoidal scarring were graded and histopathological analysis carried out accordingly. Kruskal-Wallis and Pearson Chi-Square tests were used for statistical analysis. A p-value of less than 0.05 was considered to be significant.

RESULTS: The grading of epidural fibrosis was far lower in the experimental group with curcumin compared to the control and spongostan groups, but the difference was not statistically significant.

CONCLUSION: The findings of this study show that local curcumin decreases the formation of epidural fibrosis and this effect of curcumin is thought to be mediated by reducing the functions of inflammatory cells such as macrophages, neutrophils and fibroblasts, and the anti-inflammatory and antioxidant effects.

KEYWORDS: Laminectomy, Epidural fibrosis, Failed back surgery, Curcumin, Rat
MATERIAL and METHODS

Adult female Wistar Albino rats that weighed 220-230 grams were included in the experiment that was in line with the national institutes of health guidelines for care and use of laboratory animals. Approval was obtained from the local ethics committee on animal care (Aydın Adnan Menderes University, approval no: 6458310 /2017/123, date 28 November 2017).

Experimental Groups

Group 1: Control (C); T12 total laminectomy was carried out; nothing extra was applied, n=7.

Group 2: Spongostan (S); T12 total laminectomy; distilled water and spongostan were applied on the laminectomy area, n=7.

Group 3: Local curcumin (LC); T12 total laminectomy; 100 mg/kg curcumin (purity_97) (Sigma Aldrich, Chemical Co., USA) with spongostan, n=7.

Surgical Procedure and Preparation of Samples

The temperature of the medium of the animals was adjusted as 22-25° with average humidity and the light-dark cycle was adjusted as 12/12 hour. Food and water were given ad libitum. 50 mg/kg Ketamine hydrochloride (HCl) (Ketalar, Parke-Davis, Turkey) and 10 mg/kg Xylazine HCl (Rompun, Bayer, Turkey) were administered intraperitoneally (IP) for anesthesia while spontaneously breathing. The temperature of the body was 37°C in the prone position. The rats' surgical areas were shaved and disinfected with povidone-iodine by the same surgeon. A midline skin incision was made on T10-L2. Another incision was made in the thoracolumbar fascia. The paravertebral tissues were dissected and the T10-L1 laminae were accessed. Total laminectomy was carried out at the T12 level for the rats in all groups while dura mater was visible and intact. Cotton pads were employed for hemostasis. The wounds were sutured with 5-0 polypropylene suture after application of topical curcumin. Under deep anesthesia, mixed xylazine HCl (10 mg/kg IP) and ketamine HCl (50 mg/kg IP) were administered on the 30th day for cervical dislocation. T9 and L3 vertebral columns were removed en bloc.

Histopathological Evaluation

Vertebral columns T9-L3, paraspinal tissues and epidural scar tissue were excised en bloc, put into a container which was filled with 10% neutral buffered formalin solution and fixed. Decalcification was carried out in 7 days with 10% formic acid. The specimens were then taken from the laminectomy area. Tap water was used to wash them, and paraffin was used to embed the samples after routine tissue processing; 4-micron axial sections were taken. Hematoxylin-eosin and Masson's trichrome stains were used for staining. A blinded pathologist examined the histopathological sections and analyzed arachnoid involvement and EF grade. EF in the laminectomy area was assessed with the Olympus BX52 microscope after staining and was photographed by employing an Olympus DP 25 camera. The EF grading was analyzed in line with the scale of by He et al. (8). In addition, arachnoidal involvement was also examined.

RESULTS

Complications Associated with Wound Recovery and Medical Practice

No adverse effects of curcumin were seen in the wound area and the peripheral tissues.

Histopathological Results

The groups were graded in a histopathological manner in line with the He Scheme. Grade 3 (Figure 1A, B) and 2 EF were seen in six and one rats respectively in group C. Grade 3 and 2 were seen in four and three rats respectively in group S. Grade 3 and 2 (Figure 2A, B) EF were seen in two and five rats respectively in group LC. These results showed that group LC was superior to group C and group S in reducing EF. However, there were no statistically significant differences (p<0.097) (Figure 3).

The groups were also examined for arachnoid involvement. No statistically significant differences were detected (p<0.734), (Figure 4).

DISCUSSION

Inflammation is a complex defense reaction of endogenous or exogenous signals on vascularized tissues. Neutrophils, monocytes, and macrophages have a role in the inflammation process. Proinflammatory cytokines such as interleukin (IL)-1, IL-6, IL-8 and tumor necrosis factor alpha (TNF-α) and the growth factors of platelet-derived growth factor (PDGF), transforming growth factor-β1 (TGF-β1), TGF-α, insulin-like growth factor (IGF–1) and fibroblast growth factor (FGF) are produced in the injured area by the defense cells. Activated fibroblasts and epithelial cells function in the next step of healing by cleaning the waste material (5). Fibroblasts reproduce rapidly in response to activation of inflammatory cytokines and growth factors (TGF-β and TGF-α) in order to repair the local defective vertebral area where laminectomy is performed and reach the epidural space during spinal surgery (13). Fibroblasts transform into fibrocytes as a result of the production of collagen fibers. Scar tissue emerges from...
Figure 1: A) Hematoxylin-eosin (x40 magnification) staining for epidural fibrosis in the laminectomy sites. Grade 3 fibrosis in the control group: Scar tissue completely covered the laminectomy defects and adhered to the underlying dura mater (Arrows). B) Masson trichrome (x40 magnifications) staining for epidural fibrosis in the laminectomy sites. Grade 3 fibrosis in the control group: Scar tissue completely covered the laminectomy defects and adhered to the underlying dura mater (Arrows).

Figure 2: A) Hematoxylin-eosin (x40 magnification) staining for epidural fibrosis in the laminectomy sites. Grade 2 fibrosis in the curcumin group: Scar tissue adhered to the underlying dura mater and covered less than two-thirds of the laminectomy sites (Arrows). B) Masson trichrome (x40 magnifications) staining for epidural fibrosis in the laminectomy sites. Grade 2 fibrosis in the control group: Scar tissue adhered to the underlying dura mater and covered less than two-thirds of the laminectomy sites (Arrows).

Figure 3: Histopathological assessment of epidural fibrosis grades of groups. There was no statistically significant difference between the groups (p>0.05). However, grade 2 epidural fibrosis was common in the curcumin group.

Figure 4: Frequency of arachnoidal involvement of the groups. There was no statistically significant difference between the groups (p>0.05).
fibrous connective tissue. After this process, epidural fibrosis develops as a result of transformation of adipose tissue around dura mater into fibrous tissue (21).

Several clinical and laboratory studies have been conducted for avoidance of EF but satisfying results have not been reported yet (9,14). Whatever the underlying reason, inflammatory responses and fibroblasts seem to be the most important factors during fibrosis. Unfortunately, there is no management modality other than anti-inflammatory drugs and immune suppressants to solve these problems. The aim of this study was to evaluate the effects of curcumin, which has anti-inflammatory properties, on EF.

Curcumin has antioxidant, anticancer, anti-apoptotic and anti-neurodegenerative properties (16). The impact of curcumin on EF after lumbar operations has not yet been investigated and this is the first study in the literature. As a result of this laboratory study, the Grade II EF rate in the curcumin group was found to be higher than Grade III in control group. Several effects of curcumin may have caused this result. Firstly, it might be the anti-inflammatory property of curcumin because curcumin blocks tissue proliferation and post traumatic tissue renewal by acting on macrophage cells with its anti-inflammatory property (1). It is known that macrophages show these effects by increasing the function of factors such as TGF-β, TGF-α, basic FGF, PDGF and vascular endothelial growth factor that increase proliferation and cell synthesis. Several studies suggest that curcumin decreases the amount of these factors (23). On the other hand, IL-6 and TGF-β1 has been shown to be effective on EF (3). TNF-α has also been found to be another factor in EF (15). TNF-α especially increases the proliferation and differentiation of fibroblasts and the formation of α-smooth muscle actin and the extracellular matrix by transforming TGF-β; also, TNF-α and IL-6 are strongly implicated as promoters of fibrosis (3,15). Therefore, decreasing the effect of TNF-α and IL-6 by using curcumin can result in a failure of fibroblast cell function. Curcumin has been found to have a suppressant effect on pro-inflammatory cytokine such as TNF-α which is a significant mediator in the inflammatory response (1,7). Prior studies have also suggested that this mediator induces the amount of IL-6, prostanandin E2 (PGE2), cyclooxygenase (COX-2) that have an important role in inflammation. Curcumin suppresses TNF-α-induced neuroinflammation IL-6, PGE2, COX-2 at a significant level (23); Moreover; curcumin has been shown to decrease post traumatic inflammation due to its anti-inflammatory properties in experimental spinal cord trauma studies (2). Spinal cord repair and neural function recovery were reported to be accelerated by curcumin via inhibition of glial scar formation and decreased inflammation (25). On the other hand, in the experimental study by Daishun Liu et al., it was suggested that curcumin may decrease pulmonary fibrosis via inhibiting TGF-β2 which drives the differentiation of lung fibroblasts to myofibroblasts (12). These studies strengthen the hypothesis that fibroblasts may be inhibited by curcumin and that it may therefore play a role in EF via decreasing inflammation. Our results therefore indicate that decrease of EF is due to the effect of curcumin on several steps of inflammation via its anti-inflammatory properties.

Moreover; some studies suggest that curcumin, which has antioxidant effects, may inhibit nitric oxide and reactive oxygen species in macrophages. This suggestion supports an EF-decreasing effect of curcumin (10,20).

The study also has some limitations. These are the inadequate number of rats, not comparing different doses of curcumin, and no comparison of the effect of the same dose with oral usage.

■ CONCLUSION

As a result of this study, the observation that the majority of the observed epidural fibrosis was grade II in the curcumin group and grade III in the control group indicates that curcumin has the potential to be used effectively in EF. Curcumin may decrease EF via anti-inflammatory mechanisms. However, further studies are needed to define which anti-inflammatory mechanisms play a role in this process.

■ REFERENCES