Effects of Intraoperative Local Mitomycin and 5-Fluorouracil Application on Intraocular Pressure and Wound Healing Process in the Experimental Glaucoma Filtering Surgery in Rabbits


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Summary

Purpose: The purpose of this study is to investigate the effects of intraoperative local mitomycin and 5-fluorouracil application on intraocular pressure and the wound healing process in the experimental glaucoma filtering surgery in rabbits.

Methods: Taking 42 eyes of 21 rabbits as a sample, the effects of intraoperative local mitomycin and 5-FU application on the intraocular pressure and wound healing process in glaucoma filtering surgery were studied. Mitomycin was applied on 11 and 5-FU on the remaining 10 right eyes. All 21 left eyes served as controls. On the 32nd day after the operation, the eyes of three rabbits from both mitomycin and 5-FU groups; and on the 60th day, all remaining eyes were enucleated for histopathological examination. Ultrastructure evaluation was based on transmission electron microscopy.

Results: Intraocular pressure was found to be significantly lower in all glaucoma filtering surgery groups (p<0.001). Mitomycin and 5-FU application further lowered the intraocular pressure as compared to control group (p<0.01). While intraocular pressures were lower in the first 5 days in the mitomycin group, the difference between mitomycin and 5-FU groups was not significant after 5 days (p>0.05). Bleb formation was significantly higher in the mitomycin and 5-FU groups than the control group (p<0.01). There was no difference among groups with respect to their anterior chamber depth, corneal epithelial defects and other complications (p>0.05). Histopathologic examination by light and electron microscopy revealed that mitomycin and 5-FU application delayed the wound healing process; and this effect was found to be more marked with the use of mitomycin.

Conclusions: Intraoperative local mitomycin and 5-FU application further lowered the intraocular pressure and delayed the wound healing process.

Key Words: Glaucoma filtering surgery, Mitomycin, 5-fluorouracil, Wound healing


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Filtration insufficiency is one of the most important postoperative complication of trabeculectomy procedures. The scarring of conjunctiva and Tenon's capsule is the most common cause. (1)
use of antimetabolites to prevent subconjunctival scarring in glaucoma surgery has recently been proposed. It has been shown in clinical trials that postoperative subconjunctival application of 5-fluorouracil (5-FU) and intraoperative application of mitomycin locally on the scleral flap increased the surgical success in high risk patients who went through glaucoma filtering surgery.(2-12)

Use of 5-FU and mitomycin in glaucoma filtering surgery have caused some complications. Intraoperative local application of mitomycin has been reported to cause conjunctivitis, local corneal oedema and corneal epithelial defect.(9-15) More serious complications such as corneal epithelial defects and weakness at the wound site has been associated with the postoperative subconjunctival use of 5-FU.(16-18) Therefore; recently, new trials, like mitomycin, are underway to apply 5-FU locally during trabeculectomy procedure.(19-21)

The aim of this study is to investigate the effects of intraoperative local mitomycin and 5-FU on intraoperative pressure and wound healing in glaucoma surgical filtering surgery in rabbits.

**Material and Methods**

This study was carried out in 42 eyes of 21 albino rabbits in both sexes weighing between 2 and 3 kg. They were divided randomly into two groups: Group I was the mitomycin group and Group II was the 5-FU group. The left eyes of rabbits in both groups formed the control.

**Surgical procedure**

Surgical procedures were carried out with the help of a binocular loop. General anaesthesia was induced with intraperitoneal injections of Xylazine hydrochloride (Rompun®, Bayer), 10 mg/kg and ketamine hydrochloride (Ketalar®, Eczacibasi), 50 mg/kg. A superiorly placed limbal-based conjunctival flap was then raised. Sponges soaked in mitomycin (Mitomycin C®, Kyowa) in a concentration of 0.5 mg/ml and in 5-FU (Fluorouracil®, Roche) in a concentration of 50 mg/ml were placed between the conjunctiva and the sclera in the right eyes of the rabbits in Group I and II, respectively. On the other hand balanced salt solution (BSS Plus®, Alcon) soaked sponges were placed in all left eyes. After waiting for 5 minutes, application sites were irrigated with BSS Plus solution. Sclerostomy was done with the help of 1.5 mm Elliot’s trephine and a peripheral iridectomy was performed through the sclerostomy. The conjunctival incision was closed with 8/0 polygalactine (Vicryl®, Ethicon).

**Intraocular pressure measurements**

Intraocular pressures were measured using pneumotonometer (Biorad Digilab Modular One Pneumo-Tonometer) after local anaesthesia was applied with Oxibuprocain hydrochloride (Benoxinate® 0.4%, Thilo) preoperatively, on postoperative days 1, 3, 5 and 7, twice weekly for one month and once every week until 2 months.

**Biomicroscopical examinations**

On days 1, 7, 15, 32 and 60, biomicroscopical examinations were carried out. Bleb appearances, depths of anterior chambers, hyphaemas, epithelial defects with fluorescein, leaks from wounds, corneal pigmentation, corneal haze, pannus, rubeosis and cataract were all noted. Bleb appearances were graded according to the criteria of Kay (22) et al, and anterior chamber depths were evaluated according to the criteria of Stewart (23) et al.

**Histological examination**

On day 32, three rabbits randomly chosen from both mitomycin and 5-FU groups and killed by intracardiac injection of high dose sodium thiopentone BP (Pentotal® sodium, Abbott). After immediate enucleation of the eyes, corneal and scleral pieces of tissue including the bleb regions were taken vertically for histological examination. For light microscopy (Olympus BH-2), fixation with neutral formaline 10% and paraffin blocking were followed by Haematoxylin and Eosin staining. After pre-fixation with phosphate-buffered glutaraldehyde 2.5%, irrigation with phosphate buffer 0.1 N and post-fixation with phosphate-buffered osmium tetroxide 1% were carried out for electron microscopy (Zeiss EM-9). Blocks from the filtration area were prepared with araldite semi thin sections of 1 micron and stained with Toluidin blue 1% and thin sections were stained with uranyl acetate and lead citrate.

The remaining rabbits were followed up once weekly until the sixtieth day, when their eyes were enucleated and the above mentioned procedures
were applied for examinations under light and electron microscopy.

In the statistical analysis of the data, two-sided analysis of variance, t-test, X²-test and Kruskal Wallis tests were performed.

Results

This study was conducted on three groups of eyes: Mitomycin group, 11 eyes; 5-FU group, 10 eyes; control group, 21 eyes. After the measurements on day 32, the eyes of three rabbits from each of the mitomycin and 5-FU groups were enucleated for histological examination. Because of the decrease in the number of subjects (8 eyes in the mitomycin; 7 eyes in the 5-FU and 15 eyes in the controls), each group was evaluated for their preoperative values, and follow-up measurements for postoperative values until day 32.

Mean intraocular pressure (IOP) values for three groups are shown in Figure 1.

In the comparison of the IOP values of the three groups, there was no difference in the preoperative values (p>0.05). For the values of the first 5 days postoperatively, although there was no difference between the 5-FU and control groups and 5-FU and mitomycin (p>0.05), values for the mitomycin group were found to be significantly lower than the control group (p<0.01). In the comparison of the IOP values for postoperative days 5-32, values for the mitomycin and 5-FU groups were found to be significantly lower than the control group (p<0.001). There was no statistical difference between the mitomycin and 5-FU groups (p>0.05).

The IOP values in all three groups measured on postoperative day 32 and were found to be significantly lower than the preoperative values (p<0.001). There were no differences among groups themselves (p>0.05).

Figure 2 shows the mean IOP values in all three groups measured preoperatively and postoperative day 32 and after.

There was no statistical difference in the preoperative IOP values of mitomycin, 5-FU and control groups. While there was no difference between the mitomycin and 5-FU groups in the measurements postoperative day 32 and after, IOP values of both groups were significantly lower than the control group (p<0.001).

The appearances of blebs in three groups on postoperative day 60 were as follows: 7/8 (87.50%) in the mitomycin, 5/7 (71.43%) in the 5-FU and 4/15 (26.67%) in the controls. When the mitomycin and the 5-FU groups were compared in this respect to the controls, the difference was statistically sig-

![Figure 1. Comparison of intraocular pressure values from preoperative to postoperative day 32 in mitomycin, 5-FU and control groups.](image)
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Figure 2. Comparison of the mean IOP values in mitomycin, 5-FU and control groups measured preoperatively, postoperative day 32 and after.

Significant (p<0.001). In the comparison of the mitomycin and the 5-FU groups the difference was not significant (p>0.05).

There were no differences among all three groups in anterior chamber formation, epithelial defects and hyphaema (p>0.05).

In light and electron microscopic examination of sections from the controls on postoperative day 32, the sclerostomy fields were almost filled with fibrotic tissue and were about to appear histologically like the normal sclera. It was late phase scarring according to the wound healing pathogenesis. Mononuclear cells, proliferated fibroblast in active synthesis, increased number of collagen fibers, macrophages and plasmocytes could be seen (Figure 3). In relation to the increased synthesis of collagen fibers, their phagocytosis could be observed in the cytoplasms of some mononuclear cells and Langhans type giant cells (Figure 4a and b). The findings for the control group on postoperative day 60 were similar to those on postoperative day 32. There was proliferation in the fibroblasts in the site of scarring which was proliferative late phase granulation tissue. Almost all of the sclerostomy site was filled with regular collagen bands which gave the area an appearance like that of the normal sclera. In some eyes, the iridocorneal angle

Figure 3. Active fibroblasts, increased number of collagen fibers and mononuclear cells in the control group. Electronmicrograph (EM), X4500.

was observed to be filled with fibrotic tissue which made the angle prolapse towards the anterior chamber and extend into the Descement's membrane and obstruct the filtration on a large scale.

In the histological sections of mitomycin and 5-FU groups on postoperative day 32, sclerostomy site was seen to be filled with fibrovascular tissue which was not yet formed completely between the
scleral and collagen bundles. This fibrovascular tissue was found to have formed collagen bundles in a much thinner area as compared to the controls. From the findings of postoperative day 60, it was interesting to note that both the fibrovascular tissue which filled the sclerostomy site was loose, thin and in a structure with less fibroblast activation and that the wound healing lagged much behind the duration of the experiment (Figure 5). Cicatrisation was even less in some eyes than that seen in control eyes on postoperative day 32. In the comparison of

mitomycin and 5-FU groups with respect to wound healing, the examination of light microscopic serial sections revealed that the tissue in the filtration site was looser and thinner in the mitomycin group as compared to the 5-FU group. This difference was quite minor in the comparison to the control. Electron microscopic investigation, on the other hand, revealed more prominent difference between the mitomycin and 5-FU groups than that seen through the light microscopy. These two groups (Figure 6a and b) went through a wound healing
process quite different from the controls. (Figure 7). The healing process was also in a positive way from the prognosis of the filtrating surgery point of view. A marked delay in scarring was seen in the drug treated groups. In contrast to active fibroblasts seen in the control group (Figure 8), fibroblast were mostly inactive and showed no proliferation in drug treatment groups (Figure 9). Cells characteristic of early wound healing phase were observed in drug treatment groups. There were neutrophils in the mitomycin and eosinophils in the 5-FU groups (Figure 6a and b). The most striking ultrastructural difference was that myofibroblasts were seen only in the mitomycin group. These cells were in clumps in cicatrisation sites which were looser and thinner as compared to the 5-FU group (Figure 10a and b).

**Discussion**

Scar formation in the subconjunctival tissue and the tenon capsule causes filtration failure which by itself is the most important factor affecting the success of the trabeculectomy operations. (1) There are different studies dealing with the
question of how to decrease post trabeculectomy scar formation. The post operative subconjunctival 5-FU injection (first trial done by Gressel (30) et al and first true pilot studies done by Heuer (2) et al) has showed an increase in the success rate of the surgery. In the latter studies, the application of 5-FU in complicated and uncomplicated glaucoma patients showed improvement in the prognosis of the operations. (2-8) Because the postoperative subconjunctival application of 5-FU has many serious complications; many other metabolites have recently been studied. According to these studies, intraoperative local applications of mitomycin into the flab improves the prognosis of not only uncomplicated glaucoma but also in completed ones. (9-12) From the complications of 5-FU injections and the good results of the local mitomycin injections we deduced the use of local 5-FU. (19-21)

In this study we were interested in the results of peroperative local use of mitomycin and 5-FU in filtering glaucoma operations. The results of the postoperative follow up of IOP of the three groups showed a significant fall in all the groups. In Figure 1 we notice a fall during the first postoperative week followed by a rise later on, in the IOP, in all three groups. Therefore it has been thought that the post trabeculectomy success is associated with the healing of the wound during first postoperative month. Khaw (21) and Pasquale (31) found that the operations in which mitomycin had been applied became more successful. In our study we have not seen statistical differences between mitomycin and 5-FU groups, but mitomycin group showed lower IOP than those of control from the first postoperative day on, while the 5-FU group showed lower IOP than that of control group only after the 5th postoperative day. We deduced that the surgical procedure is important in lowering the IOP and the application of mitomycin or 5-FU will enhance that effect.

Presence of the functional filtering bleb during trabeculectomy operation is one of the factors affecting the success of the operation. The filtering flabs have an important role in the passage of the aqueous humours into the subepithelial field, and in this study, the postoperative blebs in mitomycin, 5-FU and control groups are assessed according to the Kay (22) et al criteria and we noticed that there are statistically important differences in the postoperative blebs in mitomycin and 5-FU groups than that of control groups and that this difference stays for 60 days postoperatively. However we did not find any difference between mitomycin and 5-FU groups, Khaw (21) et al stated that in mitomycin group the difference stays for 30 days.

The anterior chamber depth is assessed according to the Stewards (23) et al criteria, and in all three groups we found shallowness during the first few days which improved after the first week, and there was no difference between the groups.
There are many studies about the erosive effect of mitomycin and 5-FU on the cornea. (13-18) In our study we have not found any difference between the groups and we noticed that a single application of 5-FU decreased the occurrence rate of corneal defects.

In trabeculectomy hyphemia formation effects the prognosis. The clot in the site of the sclerotomy is regarded to be an important factor in the activation of the fibroblasts. (26) In our study, we did not find statistically important differences between the groups.

Contrary to all other surgical procedures, in trabeculectomy, the healing and the closure of the wound is not the aim. Miller (32), in his study, applied filtering glaucoma surgery in rabbit eyes and studied the healing time of the wounds. Zalish (6) et al applied 5-FU after filtering glaucoma surgery and found that, although the opening in the flab is partially closed with fibrovascular tissue, it still remains intact. Our study showed similar results.

In Mandelbaum (33) endophthalmitis the micro-organisms are thought to be able to find access to the eye; though, the flab's field indicate that the weakness at the site of the flab may increase the risk of endophthalmitis. However, endophthalmitis is a serious complication of infiltration surgery. (34-35) In our study, we did not face any endophthalmitis during sixty-day follow up. Heuer (2) et al found that in case of post-operative subconjunctival injection of 5-FU, 41% shows weakness at the site of the conjunctival wound. Seidel test in none of our cases was positive. This indicate that the peroperative application of 5-FU prevents one of its most important complications. However, such an important complication obligates us to be more careful with the use of mitomycin. The absence of myofibroblasts in 5-FU may indicate that they have a function in preventing weakness at the site of the wound.

The success of filtran surgery is barely evaluated on the basis of the delay in the healing process of the wound in sclerostomy area. It is seen in our study, that this expectation has been fulfilled. However, the delay in the healing process would cause weakness at the site and related complications. The ideal is to have a filtration area that releases aqueous humour but forms a barrier for extracellular macromolecules and microorganism. For this ideal to be realised, the myofibroblasts are lacking.

Recent studies show that myofibroblasts having intrasitoplasmic contractile elements make it easier for the aqueous humour to be externally released. (36) It has been found that proteins such as vimentin, spectrin, and F-actin play an important role in releasing aqueous humour, and these proteins be marked in Schlemm's channel endotel immunohistochemically. It has also been found that myofibroblasts in trabecular meshwork supports the Schlemm's channel endotel. (37)

On the other hand, it is thought that myofibroblasts cause contraction in the wound area and make it easier for the aqueous humour to be released in early postoperative phase (between the 7th-14th day). (38) In light of the fact that myofibroblasts were not seen in 5-FU and control group may be the cause for the failure of late filtration. The high filtration caused by the myofibroblasts in the mitomycin group; in addition may lead to bleb leakage and late endophthalmitis. (39-40) Regarding the continuation of filtration, mitomycin is as effective as 5-FU with respect to structural predisposing factors for the complication and the myofibroblasts. We stated that the myofibroblast like cells have an important role in the angiogenesis and the control of the blood flow, and that the increase in the number of the aqueous veins is one of the ways of the postoperative reduction of the IOP. (41-42) There fore the reduction in the IOP in mitomycin group could be related to the effect of the mitomycin in transforming fibroblasts to myofibroblasts, even though the superiority of mitomycin in the postoperative reduction of IOP is statistically not important.

Mitomycin and 5-FU increases the efficiency and success of glaucoma filtering surgery. Further histopathological studies would explain the success of this filtration in terms of its causes in relation to the transformation evident in myofibroblasts so that the importance and role of these cells is identified.

REFERENCES


