ABSTRACT

Background: The transient increase of intraocular pressure (IOP) following neodymium YAG laser capsulotomy can occur in a significant number of patients, which requires prophylactic treatment with IOP reducing drugs, and in some patients, postoperative IOP monitoring. This study was performed to compare the efficacy of brimonidine 0.2% versus latanoprost 0.005% (Xalatan) in preventing the IOP elevation after YAG laser posterior capsulotomy in patients visiting ophthalmology clinic in Alavi Hospital.

Methods: This study was a randomized, double-blind clinical trial that included 100 patients who had developed posterior capsule opacification (PCO) as a result of previous cataract surgery and were candidate for undergoing YAG laser posterior capsulotomy. The patients were randomly divided into two groups of 50 patients. One group received brimonidine 0.2% one hour before surgery, and the other group received Xalatan 0.005% in the night before laser surgery. In both groups the patients’ IOP was measured in baseline, 1, 2, 3, 24 hours, 3 days and one week after surgery. The gathered data were analyzed using statistical methods in SPSS.16.

Results: The mean IOP, 1, 2, 3, and 24 hours, 3 days, and one week after surgery didn't show any significant difference between two groups. IOP one hour before surgery changed significantly compared to one hour after surgery and a statistically significant relationship was found between the two groups, though at other times of measurement, the differences were not significant.

Conclusions: Results showed that using Brimonidine 0.2% or Latanoprost 0.005% as prophylactic before YAG laser posterior capsulotomy could be effective in preventing IOP after treatment.

Keywords: Brimonidine, Latanoprost, Xalatan, YAG laser capsulotomy

INTRODUCTION

Cataract surgery is one of the most common ocular surgeries. The studies done in different regions have reported prevalence of the Cataract 50% and 75% in the age groups 65-74 and at the age above 75 years respectively.1,2 The most common complication of cataract surgery is posterior capsule opacity (PCO), which results from normal posterior capsule (28 percent over five years). This capsule opacification is caused by the lens epithelial cells that have survived after the loss of the nucleus and cortex, and have proliferated.1,3 Fortunately, PCO responds to neodymium YAG laser posterior capsulotomy. Transient elevated IOP can occur in a significant number of patients following neodymium YAG laser posterior capsulotomy that requires prophylactic treatment with intraocular pressure (IOP) reducing drugs, and in some patients, postoperative IOP monitoring.4

Currently, for this complication prevention, various drugs, including acetazolamide and Timololare has been used that these drugs have less side effects and need to be consumed many times by patients. Both brimonidine and Xalatan are of new drugs used for treating glaucoma, that due to their high efficacy and lower side effects are
highly prescribed by physicians and many studies in many countries have examined the effectiveness of these drugs and have found satisfying results. And now, according to these studies, brimonidine is administered before YAG laser surgery in most countries. The aim of this study was to compare the effectiveness of Brimonidine 0.2% or Xalatan 0.005% (Latanoprost) in preventing the IOP spikes after YAG laser capsulotomy.

METHODS

This is a randomized, double-blind clinical trial study that has been done on 100 patients who were candidate for undergoing YAG laser capsulotomy in ophthalmology clinic of Alavi Hospital. The patients were randomly divided into two groups of 50 patients. One group received brimonidine 0.2% one hour before surgery, and the other group received Xalatan 0.005% in the night before laser surgery.

Inclusion criteria

Inclusion criteria were having the history of cataract surgery with posterior capsular opacification and being a candidate for undergoing YAG laser capsulotomy.

Exclusion criteria

Exclusion criteria were as follows:

1. The history of previous glaucoma,
2. The history of consumption of anti-glaucoma medications,
3. Severe adhesion of lens to surrounding tissues,
4. The history of uveitis,
5. The age under 21 years,
6. Unstable cardiovascular disease,
7. Being pregnant or lactating.

The same ophthalmologist (the researcher) examined all the patients, measured their IOP (before and after surgery), and performed YAG laser posterior capsulotomy on them.

The YAG laser capsulotomy was performed at 9 am, and then in the specified intervals (Seven times) their intraocular pressure was measured and recorded.

1. Just before surgery,
2. One hour after surgery,
3. Two hours after surgery,
4. Three hours after surgery,
5. 24 hours after surgery,
6. Three days after surgery,
7. A week after surgery.

All patients received Betamethasone eye drops every 6 hours for a week after surgery. The patients whose IOP after surgery was above 40 mm Hg received other medicines in addition to these drugs, in order for their IOP to reduce. To collect data, a checklist was completed for each patient. The used drugs did not have major side effects, and the patients after signing the informed consent were included in the study. This study was approved by the ethics committee of the Ardabil Medical University. In both groups the patients’ IOP was measured in baseline, 1, 2, 3, 24 hours, 3 days and one week after surgery. The gathered data were analyzed using statistical methods in SPSS.

RESULTS

The average age of patients was 67.3 years in the brimonidine group and 61.4 years in the Xalatan group. The average of used laser energy was 217 mJ in the brimonidine group and 268.2 mJ in Xalatan group. The mean IOP before surgery was 15.30±3 in the brimonidine which was similar to Xalatan Gorup with 14.34±3.7. The mean IOP was 13.8±4.6 one hour after surgery in the brimonidine group, and 15.44±6.8 in the Xalatan group, and the difference between two groups was not statistically significant (Figure 1).

![Figure 1: Mean of IOP in two groups in various times.](image)

There was a significant change in IOP between two groups one hour after surgery, but the changes in IOP rate in other times not statistically significant between two groups (Table 1).

The type of cataract surgery in both groups in tobrimonidine and Xalatan with 70% and 60% was phacosection, respectively (Table 2).

The greatest degree of visual improvement in both groups of brimonidine and Xalatan, with 46% and 38%, respectively, was “2 lines or more in Snellen chart”, and there was no statistically significant difference between the two groups in terms of visual improvement after surgery (Table 3).

DISCUSSION

Studies on the use of drugs brimonidine 0.2% or latanoprost (Xalatan) 0.005% as anti-glaucoma eye drops for preventing IOP spikes following YAG laser
capsulotomy showed that the impact of these drops in preventing IOP elevation was similar to other drugs.\textsuperscript{3-15}

<table>
<thead>
<tr>
<th>Times IOP changes</th>
<th>group</th>
<th>Decrease from baseline</th>
<th>No change</th>
<th>Increase less than 5 mmhg</th>
<th>Increase more than 5 mmhg</th>
<th>Increase more than 10 mmhg</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 hour after surgery</td>
<td>Brimonidine</td>
<td>33(66)</td>
<td>3(6)</td>
<td>11(22)</td>
<td>3(6)</td>
<td>0</td>
<td>0.038</td>
</tr>
<tr>
<td></td>
<td>Xalatan</td>
<td>20(40)</td>
<td>8(16)</td>
<td>13(26)</td>
<td>5(10)</td>
<td>4(8)</td>
<td></td>
</tr>
<tr>
<td>2hour after surgery</td>
<td>Brimonidine</td>
<td>27(54)</td>
<td>4(8)</td>
<td>11(22)</td>
<td>4(8)</td>
<td>4(8)</td>
<td>0.7</td>
</tr>
<tr>
<td></td>
<td>Xalatan</td>
<td>21(42)</td>
<td>7(14)</td>
<td>14(28)</td>
<td>3(6)</td>
<td>5(10)</td>
<td></td>
</tr>
<tr>
<td>3 hour after surgery</td>
<td>Brimonidine</td>
<td>25(50)</td>
<td>1(2)</td>
<td>11(22)</td>
<td>7(14)</td>
<td>6(12)</td>
<td>0.3</td>
</tr>
<tr>
<td></td>
<td>Xalatan</td>
<td>25(50)</td>
<td>4(8)</td>
<td>13(26)</td>
<td>2(4)</td>
<td>6(12)</td>
<td></td>
</tr>
<tr>
<td>24 hour after surgery</td>
<td>Brimonidine</td>
<td>23(46)</td>
<td>5(10)</td>
<td>13(26)</td>
<td>3(6)</td>
<td>6(12)</td>
<td>0.6</td>
</tr>
<tr>
<td></td>
<td>Xalatan</td>
<td>24(48)</td>
<td>6(12)</td>
<td>13(26)</td>
<td>5(10)</td>
<td>2(4)</td>
<td></td>
</tr>
<tr>
<td>3 days after surgery</td>
<td>Brimonidine</td>
<td>24(48)</td>
<td>3(6)</td>
<td>14(28)</td>
<td>4(8)</td>
<td>5(10)</td>
<td>0.43</td>
</tr>
<tr>
<td></td>
<td>Xalatan</td>
<td>19(38)</td>
<td>6(12)</td>
<td>16(32)</td>
<td>7(14)</td>
<td>2(4)</td>
<td></td>
</tr>
<tr>
<td>one week after surgery</td>
<td>Brimonidine</td>
<td>19(38)</td>
<td>3(6)</td>
<td>22(44)</td>
<td>6(12)</td>
<td>0</td>
<td>0.66</td>
</tr>
<tr>
<td></td>
<td>Xalatan</td>
<td>16(32)</td>
<td>6(12)</td>
<td>20(40)</td>
<td>7(14)</td>
<td>1(2)</td>
<td></td>
</tr>
</tbody>
</table>

Table 1: Compare the IOP changes between two groups in various times.

<table>
<thead>
<tr>
<th>Type of surgery</th>
<th>Phacoemulsification</th>
<th>Phacosection</th>
<th>Extra-capsular cataract extraction</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>groups</td>
<td>N %</td>
<td>N %</td>
<td>N %</td>
<td>N %</td>
</tr>
<tr>
<td>Brimonidine</td>
<td>10 20</td>
<td>35 70</td>
<td>5 10</td>
<td>50 100</td>
</tr>
<tr>
<td>Xalatan</td>
<td>16 32</td>
<td>30 60</td>
<td>4 8</td>
<td>50 100</td>
</tr>
</tbody>
</table>

Table 2: Type of cataract surgery in two groups.

<table>
<thead>
<tr>
<th>Degree improvement</th>
<th>No change</th>
<th>From H.M to F.C</th>
<th>From F.C to CLEAR F.C</th>
<th>From F.C to Snellen chart</th>
<th>One degree improvement in Snellen chart</th>
<th>two degree or more improvement in Snellen chart</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>groups</td>
<td>N %</td>
<td>N %</td>
<td>N %</td>
<td>N %</td>
<td>N %</td>
<td>N %</td>
<td></td>
</tr>
<tr>
<td>Brimonidine</td>
<td>8 16</td>
<td>0 0</td>
<td>5 10</td>
<td>9 18</td>
<td>5 10</td>
<td>23 46</td>
<td>0.44</td>
</tr>
<tr>
<td>Xalatan</td>
<td>6 12</td>
<td>4 8</td>
<td>5 10</td>
<td>10 20</td>
<td>6 12</td>
<td>19 38</td>
<td></td>
</tr>
</tbody>
</table>

Table 3: The degree of improvement in vision after YAG in two groups.

Teresa, Nancy and Unal in their study, showed that the use of brimonidine 0.15% before surgery is comparable to the efficacy and safeness of apraclonidine 0.5% and 0.1% in preventing IOP elevation after laser anterior segment capsulotomy.\textsuperscript{5,6,16}

Seong et al in a study identified the positive influence of brimonidine 0.2% as an eye drop in preventing IOP elevation after YAG laser capsulotomy.\textsuperscript{7}

Yeom et al in a study used two different concentrations of brimonidine (0.2% and 0.15%) and showed their high effectiveness in preventing IOP rise after YAG laser capsulotomy and also they suggest that both drugs had a similar impact on preventing IOP increase.\textsuperscript{3}

Negar and et al in a study showed that in patients with high IOP and wide-angle glaucoma, latanoprost is more effective in IOP control than laser trabeculoplasty.\textsuperscript{11}

Robert and et al in a study showed that the effect of latanoprost alone is equal to the combined effect of Dorzolamide / Timolol in lowering IOP.\textsuperscript{3}

Catherine and et al in a study demonstrated that Latanoprost decreases the IOP elevation in patients after YAG laser iridotomy (LI).\textsuperscript{12} In the present study, like other studies, both medications were proved to be effective in controlling IOP after surgery.

Considering the results obtained from two groups, no significant difference was seen between groups regarding the mean pressure in the intervals of measurement. But in terms of peak rise in IOP, the greatest increase in the brominidine group (16 patients) was observed 24 hours after surgery, and in the Xalatan group (12 patients) three days after surgery. However, this difference was no statistically significant. The groups varied from each other significantly concerning IOP change one hour after
surgery, while it was not significant in other measurements carried out at other times.

As the patients in this study underwent YAG laser capsulotomy when the administered drugs were in their peak effect, that is, Bromidine 0.2% one hour before surgery (with a peak effect 2 hours after dosing), and Xalatan 0.005% the night before surgery at 10 p.m. (with a peak effect 10-14 hours after dosing), and since the previously done studies haven't taken Xalatan as such, they obtained results and statistics in this study may yield more accurate outcomes.

Moreover, in as much as the risk of IOP elevation after surgery is until 3 days after the surgery, in this study, IOP was measured 1 hour, 2 hours, 3 hours, 24 hours, 3 days, and one week after surgery. By so doing, it can be claimed that the present study can be more accurate in IOP monitoring after surgery than other studies undertaken in this area to date.

Haydar et al., examined the effect of Latanoprost 0.005% and 0.2% brimonidine in preventing IOP elevation after phacoemulsification and intraocular lens implantation surgery, and found the efficacy of two eye drops was similar.8

Novak-Laus et al., in their study found that latanoprost or bromine are more effective than pilocarpine in reducing IOP following YAG laser iridotomy.9

William et al. also compared the effect of brimonidine 0.2% and Unoprostone 0.15% (Another prostaglandin analog) on IOP reduction and found that consuming brimonidine twice a day has greater effect on reducing peak IOP than Unoprostone.10

CONCLUSION

Based on the results of this study, it can be said that the prophylactic use of a single dose of latanoprost 0.005% before YAG posterior capsulotomy is similar to use of bromine 0.2%, and is effective in preventing IOP rise. Because elevated IOP after YAG posterior laser capsulotomy occurs in most patients, and requires drug treatment after surgery it is suggested that a single dose of brimonidine or latanoprost before YAG laser capsulotomy be used as a prophylactic medicine, so that there wouldn't be need for consuming multiple drugs after surgery.

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Conflict of interest: None declared
Ethical approval: The study was approved by the Institutional Ethics Committee

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