Safety & efficacy of single subconjunctival triamcinolone 5 mg depot vs topical loteprednol post cataract surgery: less drop cataract surgery

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Abstract
● AIM: To do a randomized prospective interventional study for comparing the effects of a single subconjunctival triamcinolone acetonide (SCTA) injection to tapering topical loteprednol in patients undergoing phacoemulsification surgery under topical anesthesia.
● METHODS: A total of 400 patients were randomized into 2 groups; Group A (200 patients) received 5 mg SCTA at the end of surgery and topical ketorolac tromethamine (0.5%) with ofloxacin (0.3%) combination for 3wk. Group B (200 patients) received tapering topical loteprednol etabonate (0.5%) along with ofloxacin (0.3%) and ketorolac tromethamine (0.5%) for 3wk. Outcomes evaluated were intraocular pressure (IOP), anterior chamber cells/flare and macular oedema postoperatively at 1, 6 and 12wk.
● RESULTS: Baseline parameters were almost similar in both the groups. No statistical difference was seen between the preoperative and postoperative IOP values for Group A (P=0.82) and Group B (P=0.61) and postoperative IOP values in between both groups (P=0.14) at 1wk. Incidence of cells/flare postoperative was statistically not significant (P=0.82) in both groups at all follow up visits. Postoperative macular oedema was not observed at any follow up visit.
● CONCLUSION: SCTA appears to be an effective alternative to prolong postoperative topical steroid use.
● KEYWORDS: intraocular pressure; subconjunctival triamcinolone acetonide; phacoemulsification; loteprednol; cells/flare; less drop cataract surgery

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INTRODUCTION
Cataract surgery is one of the most common and most successful surgical interventions today. Yet, from intra and postoperative complications to patient compliance; it is not without issues. Like other types of surgeries, cataract surgery also induces an inflammatory response, commonly anterior segment inflammation and macular oedema. Postoperative regimens to minimise inflammation vary widely among clinicians. Studies[1-2] have evaluated the efficacy of alternative administrative routes of corticosteroids apart from topical to deliver higher and more stable intraocular concentrations and improve patient compliance. Injection of depot corticosteroids into sub-Tenon’s capsule is an established approach of treating various ocular inflammatory diseases[3-4]. Postoperative topical drops such as NSAIDs (e.g. ketorolac and nepafenac)[5-6] and steroids (e.g. dexamethasone and prednisolone)[7] have been the mainstay for prophylaxis and treatment of inflammation. Unfortunately, adherence to topical medication regimens is less than ideal[8].

The current study is designed to evaluate the effects of a single subconjunctival triamcinolone acetonide (SCTA) injection to that of topical loteprednol on intraocular pressure (IOP), anterior chamber inflammation and macular oedema post cataract surgery.

SUBJECTS AND METHODS
Ethical Approval This study was conducted at a tertiary eye care centre in Southern India. Institution Ethics committee approval was obtained before commencement of the study and informed consent taken from each participant. The study was performed in accordance with the tenets of the Declaration of Helsinki.

It was designed as a randomised prospective interventional study and comprised of patients who underwent phacoemulsification surgery for cataract over duration of 6mo.
from July 1st 2016 to Dec 31st 2016. A consort flow diagram for recruitment and analysis of patients in this study is shown below (Figure 1). Patients were randomised 1:1 on the basis of simple coin flipping technique. Group A patients were to receive 5 mg SCTA at the end of surgery with topical ketorolac tromethamine (0.5%) and ofloxacin (0.3%) combination 4 times per day for 7d followed by 2 times per day for 2wk. Group B patients were administered tapering topical loteprednol (0.5%) 6-4-3-2-1 times per day for 7d each with ofloxacin (0.3%) and ketorolac tromethamine (0.5%) 4 times per day for 1wk followed by 2 times per day for 2wk. Patients were followed up at 1wk (visit 1), 6wk (visit 2), and 12wk (visit 3) for evaluation of IOP, anterior chamber cells/flare and macular oedema and other adverse events if any.

**Inclusion/Exclusion Criteria**
Patients eligible for inclusion were all those who had a cataractous lens according to Lens Opacities Grading System III (LOCS III)\[^9\]. Patients excluded were those who had cataract with grade 5, 6 nuclear colour/opalescence, a prior history of uveitis or evident intraocular inflammation, suffering from glaucoma, high myopes (axial length >25 mm), known steroid responders, previous ocular surgeries, corneal diseases. Also excluded were patients with diabetes, hypertension and asthma or on any systemic anti-inflammatory therapy. In addition, those who developed any intra-operative complications such as posterior capsule rupture, zonular dialysis or those who could not complete the required follow up visits were also to be excluded.

**Baseline Evaluation**
Patients were advised not to take any systemic anti-inflammatory drugs during the study period. Preoperative parameters measured were IOP in mm Hg by Non-contact tonometry (NCT; Topcon CT-80, Topcon Corp, Japan). Slit lamp biomicroscopy examination (Topcon SL 1E, Topcon Corp, Japan) was used to rule out the presence of cells/flare in the anterior chamber and macular edema. Best corrected visual acuity (BCVA) using Snellen visual acuity was noted for every patient.

**Study Medication**
A vial of triamcinolone acetonide injection (Tricort 10, Cadila Pharmaceuticals) of 1 mL containing 10 mg/mL was used. A 1-mL syringe with 26 gauge needle was utilised to withdraw 0.5 mL (5 mg) which was subsequently injected subconjunctivally at the end of surgery.

**Surgical Technique**
Routine phacoemulsification under topical anaesthesia and implantation of a foldable hydrophobic acrylic monofocal 1 piece posterior chamber intraocular lens (Technis 1, Santa Ana, California, AMO Inc., USA) was performed by a single surgeon. After the conclusion of surgery, 5 mg SCTA was injected under the inferior bulbar conjunctiva in patients of Group A (Figure 2).

**Outcome Measures**
Follow up visits were advised at 1, 6, and 12wk and BCVA was recorded. Primary outcomes evaluated were IOP, anterior chamber inflammation and cystoid macular oedema. IOP was measured with the help of NCT with the mean of two measurements used for analysis. Anterior chamber cells/flare were evaluated on slit lamp and graded according to Standardization of Uveitis Nomenclature Working Group criteria (SUN)\[^10\]. Macular oedema was assessed clinically and using Spectralis optical coherence tomography (Cirrus HD OCT 4000).

Secondary outcomes evaluated were the size of SCTA depot in millimetres (mm) at every visit and also local adverse effects like subconjunctival haemorrhage (SCH), chemosis, congestion or necrosis, if any.

**Statistical Analysis**
Data was analysed using commercial software (SPSS version 14.0 Inc., Chicago, Illinois, USA). Paired t-test (within group) and unpaired t-test (between groups) were used to compare the IOP values. Pearson Chi-square test was used to analyse patient demographics and the presence of anterior chamber cells/flare and development of macular oedema. \(P\) value less than 0.05 was considered statistically significant (95% confidence interval) for all variables. Secondary outcomes except size of SCTA depot were analysed qualitatively.

**RESULTS**
This study enrolled 400 patients. Each treatment group...
comprised of 200 patients. There was no loss to follow up for any patient in either group. The demographics of the study population have been presented in Table 1.

Figure 3 depicts the mean IOP and their standard deviations (SD) at baseline, postoperative 1, 6, 12wk in Group A receiving SCTA and Group B receiving topical loteprednol. The mean IOP values at each postoperative visit were not statistically significant between Group A and Group B at visit 1 \( (P=0.14) \), visit 2 \( (P=0.06) \) and visit 3 \( (P=0.12) \). Nine patients in Group A had an IOP>21 mm Hg on visit 1, which reduced to 3 patients on visit 2 and to 2 patients on visit 3. There was also no statistical difference in the mean preoperative and postoperative IOP values within Group A \( (P=0.82) \) and within Group B \( (P=0.61) \) at visit 1. Similar findings were noted at visit 2 where the difference in IOP values were not significant for both Group A \( (P=0.57) \) and Group B \( (P=0.33) \) compared to baseline values. Visit 3 also revealed similar results for Group A \( (P=0.65) \) and Group B \( (P=0.39) \). Two patients who had consistent IOP values of greater than 25 mm Hg at all visits underwent excision of SCTA after 12wk follow up.

Five percent patients among Group A developed anterior chamber cells/flare by visit 1 compared to 6% among Group B. The incidence of cells/flare reduced to 1% and 1.5% of total patients in Groups A and B respectively by visit 2. There was complete resolution of cells/flare in both groups by visit 3 as seen in the data given in Table 2. The difference in anterior chamber cells/flare scores between both the groups were not statistically significant at both visit 1 and visit 2 \( (P=0.82) \).

Among both the groups receiving treatment, there was no evidence of postoperative macular oedema on slit lamp examination which was then confirmed on optical coherence tomography (OCT). There weren’t any other abnormal clinically significant ophthalmic findings.

Mean size of the SCTA depot at the time of injection was 6.5 mm. On follow up the mean sizes were 3.5 mm at visit 1, 2.25 mm at visit 2 and 1.15 mm at visit 3. Minimal SCH was noted in 5 patients on visit 1 which completely resolved by visit 2. No other local adverse effects like chemosis, congestion, necrosis or ulceration were seen in any of the follow up visits.

**DISCUSSION**

With improvement in cataract surgical techniques, the incidence of postoperative inflammation has declined but still persists. To counter it, various modalities are being tested one of which is the subconjunctival injection of steroids immediately after cataract surgery. With increasing incidence of cataracts in recent times, lack of compliance to postoperative topical medications from patients, increasing cost, and patient inconvenience etc. a single subconjunctival injection of steroid could be a viable alternative to postoperative topical steroid use. Our study shows that a single injection of SCTA at the end of an uneventful cataract surgery is equally effective if not more to prolonged topical steroid use.

Weijtens et al \(^{[11]} \) suggested that periocular injections of corticosteroids induce a higher concentration of the steroidal agent than topical administration. Of particular concern, is the increase in IOP after administration of systemic corticosteroids \(^{[12]} \). But changes in IOP, our main outcome measure were not significant in the group receiving SCTA in comparison to the baseline values. Nine patients had an IOP>21 mm Hg on visit 1 (among those 4 had an IOP>30 mm Hg). Follow up visits showed improvement in IOP values and by visit 3, IOP values of almost all patients returned to their baseline levels, except only 2 patients among Group A who had an IOP>25 mm Hg. Surgical excision of depot injection has been suggested and may be required in patients who have received sub-Tenon’s capsule injections of corticosteroids and are not responding to maximal anti-glaucoma therapy \(^{[13]} \). Considering the above, those 2 patients underwent excision of SCTA under topical anaesthesia, the indication being uncontrolled IOP. Post excision, the IOP in these patients promptly reduced to the normal range without requiring any further topical anti-glaucoma therapy.
phacoemulsification surgery can be a useful alternative to prolonged tapering topical steroid use in preventing intraocular inflammation and macular edema. Since the number, frequency and duration of instilling eye drops is significantly reduced in the SCTA group it definitely contributes to faster rehabilitation coupled with greater patient comfort. Further studies are needed for detailed analysis and to cover a broader spectrum of patients.

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