Improvement of corneal fluorescein staining in postcataract surgery of diabetic patients by an oral aldose reductase inhibitor, ONO-2235

H Fujishima, K Tsubota

**Aim:** While the mechanism in the pathogenesis of diabetic corneal disease is unclear, aldose reductase has been implicated in corneal disease. The effects of an oral aldose reductase inhibitor (ARI) on the ocular surface of diabetic patients after cataract surgery were studied.

**Methods:** This clinical trial was designed to be randomised, double blinded, and placebo controlled. Pseudophakic patients with diabetes were assigned to receive either oral ARI (ONO-2235) or placebo for 12 weeks. Vital staining of the ocular surface, tear production and clearance, break up time in tears (BUT), corneal and conjunctival sensation, and symptom scores were measured before and after treatments.

**Results:** After 12 weeks of oral ARI administration, fluoroscein staining scores (from 2.04 (SD 1.12) to 1.46 (1.18); p=0.016), conjunctival sensation (from 1.15 (0.37) to 1.36 (0.31); p=0.0006), and symptom scores (from 5.38 (1.93) to 4.00 (2.07); p=0.0002) recovered significantly. Fluoroscein staining of oral ARI administration also decreased compared with placebo (p=0.017). Rose bengal staining, tear clearance, and corneal sensation were improved although this increase was minor. Tear production, BUT, and specular microscopic evaluation of the corneal epithelium and endothelium did not demonstrate a significant change.

**Conclusion:** Oral ARI opposes the ocular surface changes caused by diabetes, by recovery of ocular surface sensitivity as demonstrated through an improvement in vital staining.

Corneal epithelial abnormality in patients with diabetes mellitus has been a lively topic from the end of 1970s when ocular surgeries such as vitreous surgery gained popularity. Since that time, it is well understood that diabetes mellitus results in the ocular complications of cataract, retinopathy, and iritis. Patients with diabetes mellitus exhibit abnormalities of the corneal epithelium and endothelium, first observed in the 1980s using specular microscopy. Ocular surface disorders, such as corneal epithelities following ocular surgery, are thought to result from the disease progression of diabetes mellitus itself. Irrespective of ocular surgery, patients with diabetes experience a variety of corneal complications, including superficial punctate keratopathy, recurrent corneal erosion, persistent epithelial defect, or trophic ulceration.

Although the molecular mechanism generating diabetic ocular surface abnormalities is unclear, the effects of aldose reductase, the first enzyme of the sorbitol pathway, in the complications in other tissues of diabetic patients suggests its involvement in the pathogenesis of this condition. The observation that diabetics demonstrate a decrease in corneal sensitivity and the complications of sterile ulcers in diabetic corneas suggests a neurotrophic involvement. Experimental studies described the effectiveness of aldose reductase inhibitors (ARI) in promoting corneal re-epithelialisation. The topical ARI, CT-112, was reported to improve corneal sensitivity, with some clinical benefits. The improvement of corneal sensation and tear dynamics in patients with diabetes mellitus also resulted from oral administration of ARI, ONO-2235 ((E)-3-carboxymethyl-5-[(E)-2-methyl-3-phenylpropenyliden] rhodanine). These data, although suggestive, are not conclusive. We therefore established a randomised, double blinded, and placebo controlled clinical trial, administering ARI orally for 12 weeks to patients with diabetes mellitus after cataract surgery to determine the effect of this treatment on the ocular surface condition of diabetic patients.

**MATERIALS AND METHODS**

**Patients**
A randomised, double blind, and placebo controlled study was performed to determine the efficacy of oral ARI in the treatment of diabetes mellitus epitheliopathy. This study was conducted in Ichikawa General Hospital, Tokyo Dental College, Chiba, Japan following approval by the institutional review board. Following their informed consent, we enrolled 27 patients with diabetes mellitus in this study who had undergone bilateral cataract extraction and implantation of posterior chamber intraocular lenses. There were no surgical complications, such as vitreous loss into the anterior chamber or haemorrhage. Diabetes mellitus was diagnosed by internal medicine doctors. Four patients discontinued their involvement in this study because of a lack of compliance or because of side effects of the medication and two patients did not perform all examinations; 21 patients (nine patients from the placebo controlled group) were followed for the duration of the study (Table 1). Although severe clinical abnormalities such as corneal ulcers or epithelial defect were not observed, all patients had had surgery performed from 3 to 9 months before the beginning of the study. All had symptoms such as foreign body sensation. Four patients had pre-proliferative retinopathy, seven had proliferative retinopathy, 15 had neuropathy, and one had renal failure. Ten patients were treated with photocoagulation. Two of the placebo controlled patients had a history of vitrectomy surgery as a result of the vitreous haemorrhage.

**Drug administration**
After discussing the nature of the study with the patients, a total of 150 mg of either placebo or ONO-2235 per day were administered three times daily for 12 weeks, 30 minutes before each meal. No other drug with known or suspected interactions with ONO-2235, or any systemic and topical
drugs which have effects on tear production or the ocular surface, was prescribed during the study.

Clinical examination
Each measurement was made by the same investigator under the same conditions. Corneal sensation, rose bengal stain and fluorescein stain, tear break up time (BUT), Schirmer’s test, and the cotton thread test, were used to evaluate the ocular surface. Central corneal sensation was measured with a Cochet and Bonnet aesthesiometer (Luneau Ophthalmologie, France). Rose bengal staining was graded from 0 to 3+ at the nasal conjunctiva, the temporal conjunctiva, and the cornea (maximal grade of 9+). Fluorescein staining was graded 0 to 9+ at the cornea. Fluorescein staining was graded 0 to 9+ at the cornea. BUT is defined as the number of seconds between the last complete blink and the first disturbance of the precorneal tear film. Schirmer’s test was performed 5 minutes after instillation of a drop of solution containing 0.5% fluorescein and 0.4% oxybuprocaine hydrochloride solution into the cul de sac.

Symptoms were recorded at the end of 4, 8, and 24 week treatment period by a single investigator under blinded conditions. Before treatment and 4, 8, and 12 weeks after treatment, subjective comfort was evaluated using a questionnaire giving a face score defined by nine faces, each showing a different expression. Specular microscopy was used to evaluate the corneal epithelium and endothelial changes.

Statistical analysis
Data were reported as real numbers. Statistical analysis utilised the paired t test and Student’s t test. Probability values smaller than 0.05 are considered to be statistically significant.

RESULTS
Before treatment, mean corneal fluorescein staining score in these patients was 2.04 (SD 1.12). After a 12 week period of oral administration of ONO-2235, this value improved to 1.46 (1.18) (p=0.016). The score of the placebo group, however, did not improve significantly (from 1.33 (0.77) to 1.61 (0.98) (p=0.31) (Fig 1). Student’s t test demonstrated significant improvement within both groups (p=0.017).

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Background of patients</th>
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<td>Patients</td>
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</tr>
<tr>
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DM = diabetes mellitus

Figure 1 Corneal fluorescein staining. Vertical bar is SD; After treatment means after 12 weeks of treatment. After treatment of ONO-2235, fluorescein staining was significantly improved (from 2.04 (1.12) to 1.46 (1.18); p=0.016), but not of placebo (from 1.33 (0.77) to 1.61 (0.98); p=0.31). Student’s t test demonstrated significant improvement within both groups (p=0.017).

Figure 2 Conjunctival sensation. Vertical bar is SD; After treatment means after 12 weeks of treatment. After treatment of ONO-2235 (from 1.15 (0.37) to 1.35 (0.31); p=0.0006) and placebo (from 1.04 (0.40) to 1.20 (0.26); p=0.045), conjunctival sensation was significantly improved.
In the cornea.

May be involved in the accumulation of pyridine nucleotides limiting enzyme in this metabolic pathway, aldose reductase, attention. Corneal autofluorescence reported that the rate activation of the polyol pathway, has received worldwide abnormalities associated with hyperglycaemia, such as the utilization to retinal hypoxia; recently, however, the role of metabolic complications may be prevented by ARI.

Theoretically, inhibition of this polyol metabolism enzyme reduces the impaired functions and histological changes in nerves. The onset and progression of diabetic complications may be prevented by ARI.

Oral ARI improves corneal epithelial changes caused by diabetes, through the recovery of corneal sensation and increased tear production. The study, an open label trial, allowed both the patients and examiners to be aware when active medication was being given, possibly biasing the results. To explore the clinical safety and benefits of oral ARI, we conducted a randomised and controlled clinical study of oral ARI. Corneal fluorescein staining was significantly improved compared with the placebo, with parallel improvements in conjunctival sensation and symptom score. The distribution of fluorescein scores also revealed that women and older patients had increased staining over placebo groups (data not shown). Improvement in the dynamics of tear production may result effective suppression of aldose reductase activity in the Schwann cells in a similar way to “dry eye” patients. Improvement of tear dynamics may improve the conjunctival and corneal abnormalities seen in “dry eye.” Improvements in rose bengal and fluorescein staining scores concurrently with a reduction of the debris in tears is attributable to improved tear dynamics.

Before treatment, the corneal fluorescein staining score of the ARI treating group was higher than the placebo group. We have no idea why it happened, but we thought it happened by chance because patients were enrolled after the double blind study.

Topical oxybuprocaine (proparacaine) increases tear film osmolality in rabbit eyes, suggesting that the increase results from decreases in tear secretion secondary to decreased ocular surface sensitivity. In this study, although cornea and conjunctival sensation demonstrated an increasing trend, significant increase were observed only for conjunctiva. The improvement of symptom score may also be due to these slight improvements of tear production dynamics and conjunctival sensitivity resulting from drug treatment. All these significant improvements may be due to this drug.

Although previously surgery was an extensive procedure, recent advances in cataract operation procedures resulting in reduced inflammation and operation time has led to decreased side effects. In this study, the ocular surface demonstrated less optical damage (our former report included two patients with corneal epithelial defect and two with corneal ulcer); and were better controlled, although only fluorescein staining and not other tear factors improved. Rose bengal staining, tear clearance, and corneal sensation were improved, but tear production or BUT, and specular microscopic evaluation for corneal epithelium and endothelium did not improve significantly. These factors may result from improvements in surgical techniques. In addition, the two case of vitrectomy belonged in the placebo control group, possibly affecting the results. These results suggest that fluorescein staining is the most sensitive measurement; this method should be used to evaluate ocular surface conditions in patients with diabetes mellitus.

In the present study, after treatment with oral ONO-2235, at 150 mg/day for 12 weeks, improved the abnormalities associated with diabetes, including ocular surface disorders and symptoms. In an animal model, the serum density of oral ONO-2235 amounted to a maximum density of 15 minutes after oral administration in male rats. Following a single oral administration, choroidal density peaked 1–3 hours with a serum density approximately 20% that of male rat. We have not clarified how ARI reaches the cornea. However it has been reported that positive radioactive findings were observed by radioimmunological assay in ocular tissues, Harder’s gland (rat specific accessory lacrimal gland) and the lens in rats after oral administration of “C labelled ARI. ARI probably reaches the cornea via tears or aequous humour.” Four patients discontinued their involvement in the study because of colour change in urine or other problems, not because of systemic side effects of this drug. ONO-2235, 150 mg/day, is safe, making it a promising candidate for effective treatment in human eyes. Administration of ONO-2235 for 12 weeks to patients with diabetes and a diabetic ocular surface condition led to improvements in tear dynamics and the symptoms of postcataract extraction keratopathy.

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REFERENCES

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