Sodium hyaluronate eye drops of different osmolarity for the treatment of dry eye in Sjögren’s syndrome patients

P Aragona, G Di Stefano, F Ferreri, R Spinella, A Stilo

Aim: To study the effect of the treatment of dry eye in Sjögren’s syndrome patients with hypotonic or isotonic hyaluronate eye drops.

Methods: 40 Sjögren’s syndrome patients were divided in two groups and treated as follows: group 1 with hypotonic (150 mOsm/l) 0.4% hyaluronate eye drops; group 2 with isotonic 0.4% hyaluronate eye drops. The eye drops were instilled six times a day for 90 days. Grading of subjective symptoms, break up time (BUT), corneal fluorescein staining, conjunctival rose bengal staining, Schirmer’s I test, and conjunctival impression cytology were carried out at 0, 15, 30, 90 days from the beginning of the study. Patients were examined in a blind fashion. For the statistical analysis the Student’s t test, Mann-Whitney U test, and χ² test were performed.

Results: Symptoms were statistically significantly improved at day 15 in both groups but group 1 patients had a global score statistically significantly better group 2 (p=0.02). At day 15 group 1 patients had an improvement from baseline values of BUT (p=0.0003), fluorescein, and rose bengal score (p=0.000001 and p=0.0004 respectively). Group 2 patients had, at day 15, an improvement of BUT and fluorescein score compared to baseline values (p=0.05 and p=0.0001 respectively). A comparison between the two groups showed better results for group 1 patients at day 15 for rose bengal stain (p=0.01) and for BUT (p=0.05) and fluorescein score (p=0.0003) at day 90. The conjunctival impression cytology showed that group 1 had a statistically significant better total score than group 2 starting from day 15 and lasting throughout the study (p<0.02). Also group 2 patients showed an improvement from baseline values starting from day 30 (p=0.000005).

Conclusion: Hyaluronate eye drops are useful for treating severe dry eye in Sjögren’s syndrome patients. The use of a formulation with pronounced hypotonicity showed better effects on corneconjunctival epithelium than the isotonic solution.
divided into two groups and assigned to a treatment by personnel not involved with the patients’ examination.

The visits and cytological examinations were carried out in a blind fashion by investigators who were not aware to which group the patients belonged. The study drugs were commercially available eye drops and the two groups were treated as follows: group 1 (17 female and three male, age 44.3 (9.8)) with unpreserved hypotonic (150 mOsm/l) 0.4% sodium hyaluronate eye drops (lalurex, Fidia Oftal, Catania, Italy); group 2 (18 female and two male, age 44.7 (12)) with 0.4% unpreserved isotonic sodium hyaluronate eye drops (Dropstar TG, Farmigea, Pisa, Italy). In both eye drops the sodium hyaluronate had a molecular weight of 750 000 D. The eye drops were administered six times a day for 90 days and the subjects enrolled in the study underwent grading of subjective symptoms and clinical examination at time 0 and after 15, 30, and 90 days.

The patients were allowed to know the brand name of the eye drops they were using. They were asked to bring, at all visits, the used boxes of the eye drops in order to control the compliance with the study protocol.

Tests performed
The following symptoms were evaluated according to a scoring system from 0 (absent) to 3 (severe): burning, foreign body sensation, dryness, mucous discharge, itching, photophobia; a global score, obtained adding the score of each symptom, was considered for the evaluation of ocular discomfort.

The following tests were performed: break up time (BUT) (seconds), corneal fluorescein and conjunctival rose bengal staining,

\[ \text{Corneal fluorescein} \]

\[ \text{Conjunctival rose bengal} \]

Tests performed
The following symptoms were evaluated according to a scoring system from 0 (absent) to 3 (severe): burning, foreign body sensation, dryness, mucous discharge, itching, photophobia; a global score, obtained adding the score of each symptom, was considered for the evaluation of ocular discomfort.

The following tests were performed: break up time (BUT) (seconds), corneal fluorescein and conjunctival rose bengal vital staining, Schirmer's I test (mm/5 minutes), and persisting up to the end of the study.

The ocular tests revealed a statistically significant improvement in BUT, fluorescein, and rose bengal score (p = 0.003, p = 0.000001, and p = 0.0004 respectively) starting from day 15 and persisting up to the end of the study.

Statistical analysis
Primary efficacy variables were considered—the global symptoms score and the impression cytology scores. The statistical analysis of the results was carried out in a blind method by a person not involved in the study, using the software sas (version 8.1). Only the results of right eyes were considered for statistical analysis. Student’s t test, Mann-Whitney U test, and \( \chi^2 \) test were used as appropriate. A value of p <0.05 was considered statistically significant.

RESULTS
Symptoms
The global discomfort index for ocular symptoms showed in both groups a statistically significant improvement, starting from day 15 and lasting throughout the study. The improvement was more pronounced in group 1 at day 15 (p = 0.02 versus group 2). At days 30 and 90 there were no statistically significant differences between the two groups for the global discomfort index score. In both groups all symptoms, with the exception of photophobia, were statistically significantly reduced starting from day 15 and persisting up to the end of the study. Photophobia was statistically significantly reduced in both groups starting from day 30 (Table 2).

Tests
Group 1
The ocular tests revealed a statistically significant improvement in BUT, fluorescein, and rose bengal score (p = 0.003, p = 0.000001, and p = 0.0004 respectively) starting from day 15 and persisting up to the end of the study.

Table 2
<table>
<thead>
<tr>
<th>Time (days)</th>
<th>Burning</th>
<th>Foreign body sensation</th>
<th>Dryness</th>
<th>Mucous discharge</th>
<th>Itching</th>
<th>Photophobia</th>
<th>Discomfort index</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group 1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>2.4 (0.8)</td>
<td>2.7 (0.5)</td>
<td>2.4 (0.5)</td>
<td>1.9 (0.5)</td>
<td>1.3 (1.1)</td>
<td>1 (0.8)</td>
<td>11.7 (3.1)</td>
</tr>
<tr>
<td>15</td>
<td>1.4 (0.5)*</td>
<td>1.2 (0.8)*</td>
<td>1.4 (0.9)*</td>
<td>1.0 (0.8)*</td>
<td>0.4 (0.5)</td>
<td>0.6 (0.5)</td>
<td>6.2 (2.2)*</td>
</tr>
<tr>
<td>30</td>
<td>0.5 (0.5)</td>
<td>0.6 (0.5)</td>
<td>0.3 (0.5)</td>
<td>0.4 (0.5)</td>
<td>0.2 (0.2)</td>
<td>0.2 (0.4)†</td>
<td>2.3 (1.4)</td>
</tr>
<tr>
<td>90</td>
<td>0.5 (0.5)</td>
<td>0.5 (0.5)</td>
<td>0.2 (0.2)</td>
<td>0.3 (0.5)</td>
<td>0.1 (0.4)</td>
<td>0.0 (0.2)</td>
<td>1.6 (1.2)</td>
</tr>
<tr>
<td>Group 2</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>2.2 (0.6)</td>
<td>2.6 (0.5)</td>
<td>2.4 (0.7)</td>
<td>1.8 (0.4)</td>
<td>1.3 (0.9)</td>
<td>1 (0.6)</td>
<td>11.4 (3.8)</td>
</tr>
<tr>
<td>15</td>
<td>1.4 (0.8)*</td>
<td>1.5 (0.7)*</td>
<td>1.6 (0.8)*</td>
<td>1.2 (0.6)*</td>
<td>0.6 (0.5)*</td>
<td>0.7 (0.6)</td>
<td>7.5 (1.7)*</td>
</tr>
<tr>
<td>30</td>
<td>0.7 (0.5)</td>
<td>0.7 (0.5)</td>
<td>0.4 (0.5)</td>
<td>0.5 (0.5)</td>
<td>0.3 (0.5)</td>
<td>0.3 (0.5)†</td>
<td>3 (1)</td>
</tr>
<tr>
<td>90</td>
<td>0.5 (0.5)</td>
<td>0.5 (0.5)</td>
<td>0.3 (0.5)</td>
<td>0.4 (0.5)</td>
<td>0.2 (0.2)</td>
<td>0.2 (0.4)</td>
<td>2.1 (1)</td>
</tr>
</tbody>
</table>

*p <0.001 v time 0 starting from day 15, †p = 0.0002 v time 0 starting from day 30; §p = 0.02 v group 2 patients at the same period of observation.
An improvement of BUT and fluorescein corneal staining was present starting from day 15 (p = 0.05 and p = 0.0001 respectively); also the rose bengal score became statistically significantly improved, starting from day 30, when compared to the baseline value (p=0.000001).

Comparison between the groups
It was possible to observe that, in group 1 patients, statistically significantly better results were achieved for rose bengal conjunctival stain starting from day 15 (p = 0.01) and for BUT and fluorescein corneal stain at day 90 (p = 0.05 and p = 0.0003 respectively) (Figs 1–3).

No differences were shown, at any time, for Schirmer’s I test either comparing the two groups or from time 0 in the same group (Fig 4).

Conjunctival impression cytology
Group 1
The conjunctival impression cytology showed a statistically significant improvement of the mean total score at day 15 (p = 0.02 versus baseline) and a further improvement at day 30 (p = 0.000005 versus day 15); thereafter the amelioration was stable (p = 0.1 day 30 versus day 90) (Fig 5).

When the single parameters were taken into account, it was possible to observe an improvement of the cell to cell junctions (p = 0.0002) and of nuclear chromatin (p = 0.04). At day 30 all the other parameters showed a statistically significant improvement versus baseline values (Fig 6).

When considering the class of cytology it was possible to observe a statistically significant improvement starting from day 15 (p = 0.002), improving at day 30 (p = 0.05 versus day 15), and remaining unchanged at day 90 (Table 3).

Group 2
An improvement of BUT and fluorescein corneal staining was present starting from day 15 (p = 0.05 and p = 0.0001 respectively); also the rose bengal score became statistically significantly improved, starting from day 30, when compared to the baseline value (p=0.000001).

Comparison between the groups
It was possible to observe that, in group 1 patients, statistically significantly better results were achieved for rose bengal conjunctival stain starting from day 15 (p = 0.01) and for BUT and fluorescein corneal stain at day 90 (p = 0.05 and p = 0.0003 respectively) (Figs 1–3).

No differences were shown, at any time, for Schirmer’s I test either comparing the two groups or from time 0 in the same group (Fig 4).

Conjunctival impression cytology
Group 1
The conjunctival impression cytology showed a statistically significant improvement of the mean total score at day 15 (p = 0.02 versus baseline) and a further improvement at day 30 (p = 0.008 versus day 15); thereafter the amelioration was stable (p = 0.1 day 30 versus day 90) (Fig 5).

When the single parameters were taken into account, it was possible to observe an improvement of the cell to cell junctions (p = 0.0002) and of nuclear chromatin (p = 0.04). At day 30 all the other parameters showed a statistically significant improvement versus baseline values (Fig 6).

When considering the class of cytology it was possible to observe a statistically significant improvement starting from day 15 (p = 0.002), improving at day 30 (p = 0.05 versus day 15), and remaining unchanged at day 90 (Table 3).
Comparison between the groups

It was possible to observe a statistically significant difference for the total score starting from day 15 and lasting throughout the study. The analysis of the single parameters showed no differences at day 15, while at day 30 group 1 values were statistically significantly better for nucleus/cytoplasm ratio (p = 0.02), chromatin pattern (p = 0.009), and goblet cells distribution (p = 0.0001). Comparing the cytological classes of the two groups, group 1 showed statistically significant better results starting from day 15 (p = 0.002) and persisting up to day 90 (p = 0.02).

Pictures representing the conjunctival impression cytology specimens in the two groups are shown in Figure 7.

DISCUSSION

Tear hyperosmolarity was indicated as the primary causative mechanism leading to discomfort, ocular surface damage, and inflammation. The eye drops used in the present study are commercially available, unpreserved eye drops, currently used for the treatment of dry eye and characterised by a same molecular weight of the hyaluronate molecule but a different osmolarity.

Sodium hyaluronate is a natural polymer and it was shown that its concentration increases in response to ocular damage and during corneal wound healing.

It was recently demonstrated that a hyaluronate receptor, CD44, is expressed in corneal and conjunctival cells and that its activation promotes the interaction with cytoskeletal proteins, suggesting a role for hyaluronate in cell adhesion and motility.

Hyaluronate promotes, in vitro, cell migration and can stabilise the ocular surface epithelial barrier suggesting that it may be directly involved in the process of epithelial repair by means of the activation of the CD44.

It was also proposed that hyaluronate may have a role in controlling the localised inflammation often present in patients with keratoconjunctivitis sicca. Interestingly, it has been recently reported that the expression of CD44 is increased in patients with moderate dry eye and superficial keratitis and that sodium hyaluronate given for 2 months period is associated with a decreased expression of this adhesion molecule. So sodium hyaluronate might have a direct role in the control of ocular surface inflammation in dry eye patients.

A previous study carried out on patients with a mild dry eye, did not show any difference in dry eye symptom relief between a hypotonic sodium hyaluronate solution of 215 mOsm/l compared with an isotonic sodium hyaluronate solution (305 mOsm/l). Furthermore, both treatments determined an improvement of signs of corneoconjunctival epithelial suffering. In the present study, carried out on Sjögren’s syndrome patients with severe dry eye, it was possible to demonstrate that a hypotonic solution of sodium hyaluronate with an osmolarity of 150 mOsm/l, lower than that previously tested, is able to induce a statistically significant improvement of ocular surface conditions, demonstrated either by impression cytology or by vital staining such as fluorescein and rose bengal. This improvement was better than that induced by an iso-osmotic solution of sodium hyaluronate of similar molecular weight. Therefore, it appears that a lower osmolarity may be important to obtain a better therapeutic result in a more severe dry eye condition.
patients with highly compromised lacrimal gland secretion like Sjögren’s syndrome patients with severe dry eye.

The efficacy of sodium hyaluronate in reducing the epithelial changes in the ocular surface, in patients with dry eye, was evaluated by Nelson following a treatment period of 8 weeks. That study showed that although sodium hyaluronate was able to improve rose bengal staining, it did not change the degree of squamous metaplasia. It was concluded that significant changes of conjunctival epithelium could take longer than 8 weeks of treatment.

In the present study impression cytology, studied with a scoring system which included seven different cytological parameters, was used as an indicator of conjunctival epithelial damage. We found an improvement of ocular surface features after treatment with both hypotonic and isotonic sodium hyaluronate but the hypotonic solution showed a quicker and better improvement in the conjunctival conditions. It is noticeable that the method used to score the impression cytology specimens allowed us to study analytically seven parameters so it was also possible to have, together with a global assessment of the specimens, an evaluation of the single aspects related to the epithelial conditions and to analytically differentiate the epithelial features following the different treatments.

Our results seem to give further support to the findings of Gilbard et al., which showed a significant positive correlation between tear film osmolarity and ocular surface damage demonstrated by rose bengal staining.

The impression cytology demonstrated that total score and cytological classes were always better in patients treated with the hypotonic solution, although starting from day 30 an improvement was also present in subjects treated with the isotonic solution.

The seven parameters used for the evaluation of conjunctival impression cytology can be divided in two categories: the former is related to the cellular features and includes cell to cell junction, nucleus/cytoplasm ratio, nuclear chromatin, and keratinisation; the latter is related to tissue parameters and includes specimen cellularity, goblet cells distribution, and presence of inflammatory cells.

In group 1 subjects an improvement in cellular parameters such as cell to cell junctions and nuclear chromatin was observed at day 15. The former indicates an amelioration of the junctional complexes between the epithelial cells, so that the cells which initially appeared isolated were clumped in sheets where it was not possible to distinguish the cellular borders; the latter showed, before treatment, signs of cellular activation, such as presence of nucleoli, or damage like unevenly dispersed chromatin with presence of nuclear “snakes”; already after 15 days of treatment the chromatin pattern was nearly normal, being characterised by the presence of evenly dispersed chromatin. From day 30...
onwards, all the other parameters showed a statistically significant improvement from baseline values.

In group 2 subjects, a statistically significant improvement of cell to cell junctions and specimen cellularity, goblet cell distribution, and presence of inflammatory cells was observed only from day 30. The amelioration of these four parameters was enough to maintain a statistically significant improvement of the mean cytological score if compared with baseline features.

The goblet cell distribution is modified in response to irritating stimuli that, in more advanced stages of damage, bring a dramatic reduction or the disappearance of goblet cells from the conjunctival surface, this being a sign of squamous metaplasia like that typical of the advanced dry eye. When the irritating stimuli on the conjunctiva are relieved, goblet cells tend to appear again on the conjunctival surface.29

It is noteworthy that both isotonic and hypotonic sodium hyaluronate were able to induce an amelioration of the cell to cell junctions, since this effect could be an indirect demonstration of the interaction of hyaluronate with its receptor CD44, whose activation plays a part in cell adhesion and motility.29

From the results of this study it is possible to assess that sodium hyaluronate represents a useful treatment for severe dry eye in Sjögren’s syndrome patients. In particular, high concentrations of sodium hyaluronate were able to provide a quick reduction in ocular symptoms in these patients. The treatment with hypotonic sodium hyaluronate eye drops gives a statistically significantly better improvement in clinical and cytological signs of the disease, compared with isotonic sodium hyaluronate at the same concentration.

The use of hypotonic hyaluronate eye drops should be encouraged for the treatment of patients with severe dry eye, where an increased tear osmolarity could play a part in the pathogenesis of the ocular surface damage.

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