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Long-term outcome after topical ciclosporin in severe dry eye disease with a 10-year follow-up

Morgane Straub,1 Alain M Bron,2,3 Aurore Muselier-Mathieu,2 Catherine Creuzot-Garcher2,3

ABSTRACT

Aim To report a 10-year follow-up of patients suffering from severe dry eye syndrome (DES) initially treated with topical ciclosporin A (tCSA) for 6 months.

Methods The charts of 26 patients with severe DES related to keratoconjunctivitis sicca (KCS) and followed for a minimum 10-year follow-up were retrospectively reviewed. All of them were treated initially with tCSA for 6 months. The Schirmer I test, green staining scores and tear Im break-up time (TIBUT) were recorded to assess clinical symptoms before, during and after treatment. The subjective signs were evaluated with the ocular surface disease index (OSDI) questionnaire. Prolongation and reintroduction of tCSA after the initial treatment and combined treatments were also noted.

Results Overall the median (IQR) duration of tCSA treatment was 23 (7–51) months after a prolonged induction treatment lasting 20 (8–41) months during the 10-year follow-up. For symptoms, a statistically significant difference in the OSDI between baseline and the end of the 10-year follow-up was not found (p=0.67). We noted a statistically significant improvement in all clinical signs after the initial treatment period, still present at the end of follow-up. Only 6.5% of the patients needed reintroduction of tCSA after their prolonged induction treatment.

Conclusions The improvement observed after an initial tCSA treatment was sustained after a long-term follow-up with few cases requiring additional tCSA treatment. A prolonged induction treatment to decrease initial in ammatory local signs is a promising option in KCS.

INTRODUCTION

Dry eye disease (DED) is a frequent disease with a prevalence reaching 15% after 65 years of age. Women are particularly impacted after menopause because of hormonal status modifications.1 DED is associated with symptoms of ocular discomfort such as burning, foreign body sensation, dryness, grittiness, ocular pain, photophobia, blurred vision and visual fatigue.1 Keratoconjunctivitis sicca (KCS) is the most severe presentation of DED with subjective symptoms and objective signs related to ocular surface impairment. Several factors have been identified in DED: in animation plays a major role with an increased expression of inflammatory markers, namely HLA-DR, intracellular adhesion molecules, in ammatory cytokines (interleukin (IL) 1, IL-6, IL-8), tumour necrosis factor and increased activity of metalloproteinases in the tear fluid.6 First-line treatments for KCS are palliative and do not treat the underlying cause of the disease. Unlike arti cial tears, topical ciclosporin A (tCSA) works to restore the ocular surface, allowing increased production of tears by inhibiting the T-cell activation pathway. As an immunomodulatory agent, ciclosporin A, acting through the cell-mediated in ammatory cascade, can reduce dry eye syndrome (DES).3 In two multicentre, randomised, prospective, phase III trials, tCSA 0.05% was shown to increase Schirmer values, reduce corneal staining, improve subjective symptoms and reduce the dependence on arti cial tears.7 Therefore, ciclosporin has been shown to be an effective therape u tic agent for moderate-to-severe dry eye and was approved for this indication in December 2002 in the USA. The purpose of this retrospective study was to report the outcome of patients treated with an initial 6-month induction phase with tCSA 0.05% (Restasis, Allergan, Buckinghamshire, UK) followed by an ‘as-needed’ tCSA treatment period during a 10-year follow-up.

PATIENTS AND METHODS

Patients Patients suffering from moderate-to-severe KCS were enrolled in a clinical trial conducted by Allergan (Study 192371-501) in May 2002 to assess the ef cacy and the safety of tCSA in patients with moderate-to-severe dry eye. The study followed the Declaration of Helsinki guidelines for research involving human subjects and was approved by the local institutional review board. Once included, all the patients were treated for at least 6 months according to the design of the trial (period A). At the end of this trial, patients presenting persistent clinical signs and symptoms of dryness, tCSA was maintained through a temporary use authorisation according to the examiner’s decision (period B). In some cases, tCSA was temporarily stopped after period B and then reintroduced (period C). Gender, age and duration of ocular dryness at inclusion as well as associated systemic diseases were collected. For this retrospective study, only patients with a minimum follow-up lasting 10 years were studied (see figure 1). All the patients were followed during this 10-year period in the Department of Ophthalmology, Dijon University Hospital, Dijon, France. The inclusion criteria were the Schirmer I test ≤5 mm/5 min in both eyes at baseline, tear break-up time (TIBUT) ≤10 s (average of three tests) and fluorescein score >2 (Oxford Scheme) in both eyes. To be included, patients had to have felt uncomfortable for at least 3 months with a minimum of two symptoms among the following: feeling of dryness, foreign body sensation, burning or stinging. Exclusion criteria were the presence of uncontrolled systemic disease, active


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ocular infection or non-KCS ocular in ammation, known allergy or sensitivity to the study medications or its components, pregnancy, previous photorefractive surgery, abnormal lid position and contact lens wear.

Clinical examination
A complete ophthalmic examination with visual acuity, ocular pressure measurement and biomicroscopy was performed by a single examiner (CC-G). Patients presenting meibomitis (mild, moderate or severe) were identi ed. Corneal and interpalpebral conjunctival staining, TBUT and Schirmer test results were successively collected. The subjective symptoms were evaluated using the ocular surface disease index (OSDI) score and the number of arti cial tear eye-drops or ointment was recorded. For corneal t incic staining, the entire cornea was examined using the slit lamp with blue cobalt illumination and a yellow barrier lter. Staining was graded using the Oxford Scheme 6-point scale, from 0 to 5, using the same set of photographs as a guide.5 Lissamine green was then instilled, and interpalpebral conjunctival staining was evaluated, between 30 s and 2 min after instillation. Staining was graded using white light of moderate intensity, on interpalpebral regions of temporal and nasal conjunctiva, using the same Oxford Scheme. The sum of corneal and interpalpebral staining was therefore on a 0-point to 15-point scale. A negative change from baseline indicated an improvement. TBUT was performed with a slit lamp at ×10 magni cation using blue cobalt illumination. The Schirmer test I (without anaesthesia) was graded in millimetres after 5 min. A positive change from baseline indicated improvement. These clinical tests were performed before, during and after tCSA treatment and at each visit during the follow-up. Subjective symptoms were graded with the OSDI questionnaire.6 A negative change from baseline indicated improvement.

Treatment
During the 6-month initial period, patients were asked to use tCSA 0.5% twice a day. They were allowed to use concomitant arti cial tear eye-drops if needed. At the end of this initial period (period A), if relieved, patients could stop tCSA and use only arti cial tears as needed. In other cases, tCSA was prolonged and prescribed twice daily as long as necessary (period B) to obtain a signi cant improvement of signs and symptoms according to the examiner’s decision. In some cases, after a temporary stop, tCSA was reintroduced during the follow-up (period C) if needed. This was done through a temporary compassionate use of this drug since this drug was not commercially available in France at that time.

We recorded the total duration of tCSA treatment (period A +B+C), the number of patients requiring a prolongation of tCSA after period A and the duration of the prolonged induction period A+B, the number of patients who needed a reintroduction of tCSA after period A+B and the number of patients till under tCSA at the end of the follow-up. We also recorded the number of additional treatments per day (before and during the follow-up), especially steroids, arti cial tear drops and ointments as well as the side effects related to all the treatments.

STATISTICAL ANALYSIS
We considered only one eye per patient for analysis. We used non-parametric tests because the variables were not normally distributed (D’Agostino and Pearson omnibus normality test). Descriptive statistics are given as the median (IQR) for continuous variables and number (percentage) for categorical variables. For comparisons, we used the paired Wilcoxon test for continuous variables and the Fisher exact test or χ2 test for dichotomous data. The Spearman test was used for non-parametric correlations. The Prism V5.01 software (Graphpad Software, La Jolla, California, USA) was used and p values <0.05 were considered statistically signi cant. All the tests were two tailed.

RESULTS
A total of 36 patients were included during period A, and 26 patients were ultimately followed for 10 years ( cure 1). The characteristics of the patients are displayed in the table 1.

Table 1 Patient characteristics at baseline

<table>
<thead>
<tr>
<th>Variable</th>
<th>Number of Patients</th>
<th>Median (IQR)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex (female)</td>
<td>25 (96.2)</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>60 (57–66)</td>
<td></td>
</tr>
<tr>
<td>Sjögren disease associated with dry eye</td>
<td>22 (64.6)</td>
<td></td>
</tr>
<tr>
<td>Duration of ocular dryness (months)</td>
<td>48 (29–56)</td>
<td></td>
</tr>
<tr>
<td>Schirmer I test (mm)</td>
<td>1 (0–4)</td>
<td></td>
</tr>
<tr>
<td>Fluorescein staining score (Oxford scale)</td>
<td>3 (3–4)</td>
<td></td>
</tr>
<tr>
<td>Lissamine green staining score (Oxford scale)</td>
<td>7 (6–8)</td>
<td></td>
</tr>
<tr>
<td>Break-up time (s)</td>
<td>3 (3–4)</td>
<td></td>
</tr>
<tr>
<td>Ocular surface disease index (OSDI) score</td>
<td>60.4 (50.0–73.4)</td>
<td></td>
</tr>
<tr>
<td>Global number of drops per day</td>
<td>6 (5–10)</td>
<td></td>
</tr>
</tbody>
</table>

Categorical variables are displayed as number (percentage). Continuous variables are displayed as median (IQR).

Twenty-two patients had Sjögren syndrome. One patient suffered from sarcoidosis and another one from primitive biliary cirrhosis. Two patients presented xerostomia and xerophthalmia with KCS but did not meet all the criteria for Sjögren disease according to Vitali et al.7 Sixty-three per cent of the patients presented associated meibomitis defined as mild (23.1%), moderate (30.2%) or severe (15.4%). At baseline, the median OSDI score was 60.4 (50.0–73.4) and 48.9 (33.3–64.6) after the 10-year follow-up (p=0.15). The Schirmer I test improved from 1.0 (0.0–4.0) mm at baseline to 2.5 (1.2–3.0) mm at 6 months without any statistical difference (p=0.12). At the end of the 10-year follow-up, the median Schirmer I test reached 3.0 (2.0–5.0) mm and was signi cantly improved compared with baseline (p=0.02). The t incic staining score was signi cantly lowered, from 3.0 (3.0–4.0) at baseline to 2.5 (1.2–3.0) after 6 months (p<0.001) and 2.0 (1.1–3.0) after the 10-year follow-up (p<0.001). The lissamine green conjunctival staining score decreased signi cantly from 7.0 (6.0–8.0) at baseline to 6.0 (3.5–7.0) at 6 months (p<0.01) and 5.0 (4.0–6.0) at 10 years of follow-up (p<0.001). The TBUT increased from 3.0 (3.0–
Clinical science

4.0) at baseline to 4.0 (3.0–5.0) seconds at 6 months (p=0.99) and rose to 4.0 (3.0–6.0) at the 10-year follow-up (p=0.03).

After a 10-year follow-up, the median duration of tCSA treatment was 23 (7–51) months (period A+B+C). All the patients still needed prolonged treatment with tCSA after the rst 6 months. The duration of prolonged induction treatment (A+B) without any tCSA discontinuation was 20 (8–41) months. After this initial tCSA treatment period (A+B), only 6.5% needed reintroduction of tCSA (period C), after a median delay of 40 (25–54) months. Only two patients were still taking tCSA after the 10-year follow-up. The median time without any tCSA treatment was 24 (11–53) months during the whole 10-year period.

The overall number of topical lubricant eye-drops per day was signi cantly lower during the tCSA treatment compared with baseline, 5 (3–6) vs 6 (5–10) eye-drops (p=0.001). The number of corticosteroid eye-drops per day decreased signi cantly from baseline to 6 months and 10 years, 1.5 (1–3), 0.5 (0–2.0) and 0 (0–0) eye-drops (p=0.001 and p<0.04, respectively). Less ointment application was used at the end of the initial 6-month tCSA treatment 0 (0–0) vs 1 (0–1) (p=0.01), but a statistical difference was no longer found after 10 years 0.5 (0–1) (p=0.19).

Local adverse effects were noted throughout the follow-up for 15% of patients: 10% presented with a burning sensation at instillation, 3% a stinging sensation at instillation and 2% red eyes immediately after instillation. Only two patients temporarily interrupted tCSA treatment because of local side effects, during a short period of 1.5 (1–2) months, but tCSA treatment was rather well tolerated after reintroduction in these patients.

DISCUSSION

DES is a common disorder that is on the rise and is responsible for a signi cant impact on quality of life even in people with good vision, and can be considered as a public health problem.59 The diagnosis is based upon symptoms of ocular discomfort and/or objective clinical signs. In 2007, the International Dry Eye Workshop (DEWS) categorised dry eye into four levels of severity and established treatment guidelines based on disease severity.10 11 The management of KCS encompasses two main approaches. The non-pharmacological approach includes eyelid hygiene, avoidance of exacerbating factors and punctal plugs in selected cases. The pharmacological approach includes arti cial tears, oral antibiotics (tetracyclines) in selected cases, topical corticosteroids and tCSA. Anti-in ammatory therapies are recommended beginning at level 2 (level 2 is characterised by both moderate symptoms and objective conjunctival and corneal staining).12 Several randomised studies have shown the ef cacy of tCSA in DED.5 13 Indeed, in ammation plays a key role in dry eye pathogenesis. The in ammatory cell in ltration of the ocular surface leads to an increased expression of adhesion molecules and in ammatory cytokines.14 tCSA was discovered in the early 1970s as an antifungal agent; it is a neutral, hydrophobic, cyclic metabolite of the fungi Tolypocladium in atum and Beauveria neuvei.15 Through its immunomodulatory effect, tCSA speci cally blocks T-cell activation and prevents synthesis of several proin ammatory cytokines.5 16 17 Therefore, it decreases ocular in ammation and restores ocular surface integrity, with increased tear production.18 In contrast to topical corticosteroids, which carry the main risks of secondary glaucoma and cataract,19 the good safety pro le of tCSA is appropriate for long-term therapy.20

In the present study, clinical tests showed an objective improvement after tCSA treatment. We found a signi cant improvement of uorescein and the lissamine green staining score during the tCSA treatment compared with baseline during the rst 6 months of use. When we compared baseline results and those obtained after the 10-year follow-up, with a median period of 23 months with tCSA treatment followed by a median period of 24 (11–53) months with no tCSA, we found a signi cant improvement for the Schirmer test, uorescein and lissamine green staining scores and TBUT. These results are in accordance with Sall et al., who, in their large multicentre randomised phase III clinical trial, found that tCSA was effective in the treatment of moderate-to-severe chronic dry eye. Wilson and Perry21 also suggested the long-term effect of tCSA after a 1-year follow-up in DES. In our study, only 6.5% of our patients had tCSA reintroduced after the prolonged induction phase. Indeed, almost all of our patients needed prolonged tCSA treatment with a median duration of 23 months. Other studies have demonstrated that tCSA might require several months to produce a therapeutic effect and up to 6 months for maximum improvement.19

In this series, the OSDI score decreased by 10.5% at the 10-year follow-up. In a large study with 5884 patients, Stonecipher et al.22 found similar outcomes with a statistically signi cant improvement of dry eye symptoms after tCSA treatment, with reductions in the impact of KCS in daily activities. The overall number of eye-drops used by our patients was signi cantly decreased during the tCSA treatment period and at the end of the 10-year follow-up. This was similar for topical corticosteroids. In our study, most of the patients presented associated meibomitis and showed a good clinical improvement with tCSA. In their study, Rubin et al.23 reported that tCSA should have a positive impact on meibomitis, with an improvement of the viscosity of meibomian gland secretion, TBUT and Schirmer scores compared with a control group (tobramycin 0.3%/dexamethasone 0.1%).

Overall in our series, tCSA was well tolerated with few patients presenting side effects. However, the primary Sjögren syndrome present in most patients can be associated with trigeminal sensory neuropathy, responsible for a decreased corneal and conjunctival sensation, which can explain this low side effect rate.24 Some patients reported non-speci c side effects such as burning, stinging and red eyes at instillation. Interestingly, this occurred only in a later period. This could be related to recovery of corneal sensation after tCSA treatment.25 Indeed, Liang et al.23 studied the safety of tCSA and concluded that tCSA was generally well tolerated, with no signi cant side effects. Considering the entire population treated initially, among the 36 patients, 4 needed to stop the treatment for a local side effect at an early stage and 2 patients secondarily experienced poor tolerance at instillation but did not stop tCSA for safety reasons.

These results suggest that tCSA with at least a 6-month initial treatment and afterwards a 23-month median treatment duration could have a sustained effect on dry eye symptoms and could slow down disease progression, without signi cant side effects. Although reported in the literature, this hypothesis needs to be con rmed with more robust ndings in further series. While most patients required a limited period of tCSA treatment, a few patients were still under treatment at the end of this long follow-up, although none of them needed continuous treatment for 10 years (range 7–51 months). To predict which patient will need prolonged treatment remains challenging, but early treatment with ciclosporin in moderate-to-severe cases resistant to well-conducted treatments before brosis occurs seems a good option.

The patients included in this study can be considered as having relatively severe DES. The baseline clinical presentation associated with well-identi ed underlying systemic disease (85% of patients presented Sjögren-associated DES) can be considered as


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worsening conditions. However, the fact that almost all our patients had Sjögren syndrome can also probably explain the good ef cacy of ciclosporin because the tissue damage is due to cell-mediated injury.27,28 The relevance of such an improvement of DED remains to be assessed from a clinical point of view. Indeed, DED is characterised by a poor correlation between signs and symptoms.29 In our series, the improvement of the oorescein score remains moderate, varying from 3 at baseline to 2 after 10 years, for a mean 20% improvement that can be considered as clinically relevant. The statistically non-signi cant improvement of symptoms through OSDI results can be partly explained by the frequent corneal hyposthesia found in severe DES. Therefore, a good indirect re ection of ef cacy was the decreased number of lubricant eye-drops and steroid use by the patients.

We acknowledge several limitations to this study. First, the small size of the sample and the retrospective design of the study undertaken in one centre are real weaknesses. Second, a potential confounding factor is the common use of systemic treatments such as steroids and immunosuppressive drugs that can interfere with symptoms. While these treatments were requested to be stable for at least 6 months at inclusion, it is likely that these treatments varied at least for certain periods during such a long follow-up. However, the impact of systemic treatments is dif cult to evaluate on local in amnatory signs. Third, the absence of a control group is another caveat and the question can be raised as to whether or not this improvement observed after such a long follow-up corresponded to spontaneous progression. However, all these patients belonged to the most severe group de ned by the Delphi score with a pre-existing optimal treatment and regular use of steroid eye-drops.12 The fact that these patients with severe ocular surface diseases should have improved spontaneously with a subsequently decreased number of lubricant eye-drops is quite unlikely, but due to the limitations of the design of our study, we cannot draw de nitive conclusions. Moreover, from an ethical point of view, it would have been impossible to leave these patients with such severe disease without treatment or with treatment such as steroids that have potential side effects for such a long time. Fourth, we have only considered patients with severe dry eye, thus limiting the number of include patients and the conclusions, which could have been drawn from other subgroups.

The strength of this study is the long follow-up. To our knowledge, this is the rst time that such a long follow-up has been reported with tCSA. Our aim was to assess the ef cacy of a tCSA treatment in durably improving ocular surface conditions. Another goal was to determine which information should be delivered to a patient su ering from severe DED and aiming to begin treatment with tCSA. This could be summarised as follows: the initial treat ment with tCSA should be longer than 6 months. Local side effects related with tCSA use should be explained in details to the patients in order to improve a long-term compliance.

In conclusion, tCSA treatment decreased the objective clinical signs of KCS and was associated in a long-term perspective with an improvement of symptoms on everyday activities and a reduction of articial tear use.

Contributors Design and conduct of the study (AMB, CC-G); collection, management, analysis and interpretation of the data (MS, AMB, AM-M, CC-G); preparation, review or approval of the manuscript (MS, AMB, AM-M, CC-G).

Competing interests AMB: consultant for Allergan, Bausch Lomb, Théa, research grant Honus; CC-G: consultant for Allergan, Bayer, Bausch Lomb, Novartis, Théa, research grant Honus.

Patient consent Obtained.

Ethics approval Local institutional review board (CPP Est).

Provenance and peer review Not commissioned; externally peer reviewed.

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