

Efficacy and Safety of Phospholipid Nanoemulsion-Based Ocular Lubricant for the Management of Various Subtypes of Dry Eye Disease: A Phase IV, Multicenter Trial

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Purpose: To evaluate the clinical efficacy and safety of propylene glycol/hydroxypropyl-guar (PG-HPG)-based nanoemulsion (Systane[®] Complete) lubricant eye drops in participants with dry eye disease (DED).

Participants and Methods: In this phase IV, open-label, single-arm, interventional, multicenter study, adult participants with DED – subtyped into aqueous deficient, evaporative, and mixed dry eye – were instructed to instill one drop of PG-HPG in each eye twice a day for 28 days. Endpoints included change from baseline in tear film break-up time (TFBUT) (primary) and ocular discomfort visual analog scale (VAS) score at Day 14 and TFBUT at Day 28 (secondary). Safety was assessed throughout the study. Data were analyzed for overall patient cohort and by DED subtypes.

Results: A total of 134 participants received treatment (mean age: 56.6 years; female: 75.4%). At Day 14, the mean (standard deviation [SD]) TFBUT increased from 2.6 (1.00) seconds at baseline to 4.2 (2.92) seconds (mean change: 1.5 [2.80] seconds); median change was 0.8 seconds (95% CI: 0.52, 1.19) for the overall cohort; the change was maintained at Day 28 (mean change: 1.4 (2.80) seconds, median change: 0.8 seconds (95% CI: 0.49, 1.17)). Subgroup analysis showed a mean change (SD) from baseline in TFBUT of 1.1 (2.41) seconds for aqueous deficient, 2.4 (3.17) seconds for evaporative, and 1.2 (2.63) seconds for mixed dry eye at Day 14, respectively. The ocular discomfort VAS score improved (decreased) from baseline at Day 14 (mean [SD] change: -17.3 [24.80]) for the overall cohort, -22.0 [21.73] for aqueous deficient, -17.6 [24.17] for evaporative and -13.1 [27.49] for mixed dry eye subgroups). Adverse events (AEs) were reported in 9 (6.7%) participants. No serious AEs were reported during the study.

Conclusion: Our study findings demonstrated that PG-HPG-based nanoemulsion lubricant eye drops were effective and well tolerated in participants with DED and all of its subtypes.

Keywords: aqueous deficient dry eye, dry eye disease, evaporative dry eye, propylene glycol/hydroxypropyl guar-nanoemulsion ocular lubricant, mixed dry eye

Introduction

Dry eye is a multifactorial disease of the ocular surface, characterized by loss of homeostasis of the tear film and accompanied by ocular discomfort and disturbed vision, thereby impacting quality of life.^{1,2} As a leading reason that patients seek professional eye care, it poses a significant economic burden.³

Artificial tear products or ocular lubricants that attempt to replace and/or supplement the deficient natural tear film are the initial mainstay treatment option in the management

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of dry eye disease (DED).⁴ Most of the available artificial tears currently on the market target either aqueous deficient or evaporative DED. Selection of an appropriate artificial tear to address the needs of a patient may be challenging for general eye care practitioners or pharmacists who may not be capable of performing an eye examination utilizing the appropriate series of tests to diagnose the exact type of DED.⁵ Furthermore, 30–70% of DED patients may suffer from symptoms of both aqueous deficient and evaporative type of DED (mixed type) and require treatment for both dry eye types for maximum relief.⁴ Thus, there is a need for a tear film ocular lubricant that can be used for managing all types of DED. Further, some of the significant challenges with ocular lubricants are the low retention time of active ingredients and blurred vision following instillation.^{4,6} In addition, a lipid-based ocular lubricant that can optimize the coverage area of lipids delivered to the ocular surface is an unmet need of patients with evaporative and mixed DED.

Systane[®] Complete lubricant eye drops (Alcon Laboratories, Inc. Fort Worth, TX, USA) is an innovative propylene glycol-hydroxypropyl-guar (PG-HPG) nanoemulsion, designed to replenish deficiencies in both the lipid and aqueous layers of the tear film.⁷ PG-HPG contains propylene glycol as an active demulcent, phospholipid nanoparticles to improve the coverage area of lipids delivered to the tear film and make the emulsion formulation less opaque (ie, translucent), and a higher amount of HPG to enhance the retention of lubricant on the ocular surface.

In a preclinical study, PG-HPG has been shown to provide effective moisture retention, protection, and improved cell barrier function and lubricity of corneal epithelium, which are required for management of all types of DED.⁸ However, the effect of this PG-HPG nanoemulsion-based eye drop in patients with all types of dry eye needed exploration.

The objective of this study was to evaluate the clinical effectiveness and safety of PG-HPG in adult study participants with all subtypes of DED.

Materials and Methods

Study Design

Between July 2018 and May 2019, participants with all subtypes of DED (aqueous deficient, evaporative, and mixed) were enrolled at six centers across North America and Europe in this phase IV, 28-day, open-label, single-arm, interventional study (NCT03492541). The study consisted of two phases: the screening phase (Days –7 to 0), which assessed the study participants' eligibility, followed by the treatment phase (Days 1 to 28; Figure 1). Eligible study participants received the first dose of PG-HPG on Day 1 and were required to self-administer twice-daily dosing for 28 days with scheduled visits on Day 14 and 28; additional doses could be administered as needed if the study participants experienced dry eye symptoms.

Written informed consent was obtained from each participant before entry into the study. The study was approved by

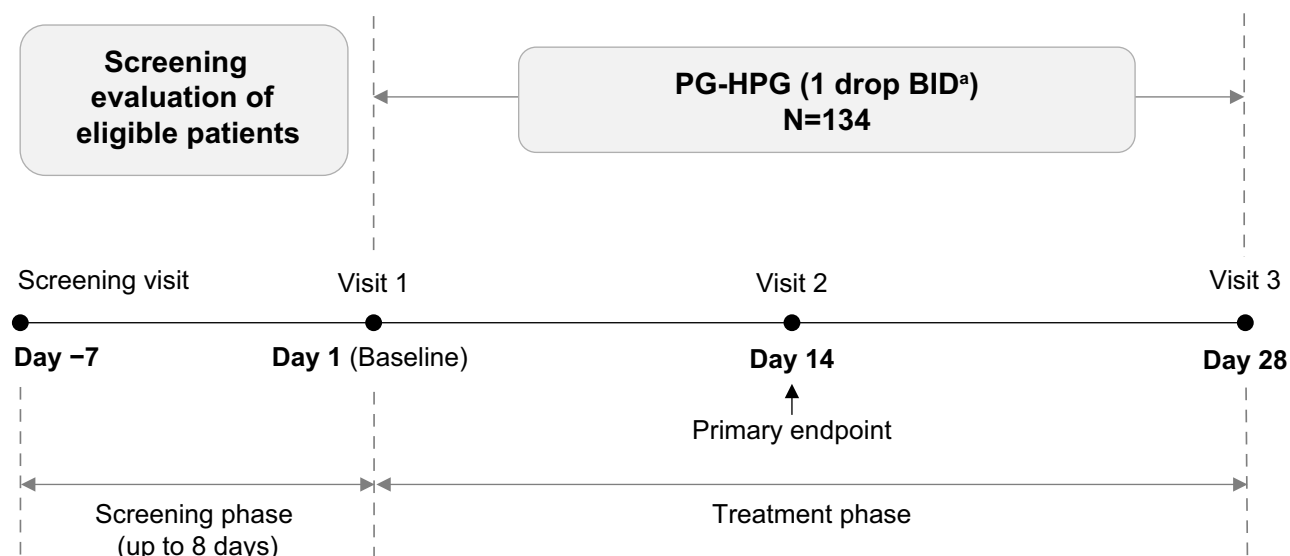


Figure 1 Study design. ^aOne drop was instilled twice-daily (morning and evening) in both eyes of eligible participants; additional doses were allowed in-between the two dosing intervals.

Abbreviations: BID, two times daily; PG-HPG, propylene glycol/hydroxypropyl guar-based lubricant eye drops.

the independent ethics committee or institutional review board at each participating center and was conducted in accordance with the Declaration of Helsinki, the International Conference on Harmonization, and Good Clinical Practice guidelines ([Appendix 1](#)).

Eligibility

Participants aged ≥ 18 years, having a tear film break-up time (TFBUT) of ≤ 5 seconds in at least one eye, and best corrected visual acuity of $\geq 20/80$ (or ≥ 55 letters score or ≥ 0.6 early treatment diabetic retinopathy study log of the minimum angle of resolution value) in both eyes at screening were eligible for inclusion. In addition to the above, participants who met at least one of the three criteria below in at least one eye at screening (randomization stratified based on the criteria below) were also included: (a) unanesthetized Schirmer I test score of ≤ 9 mm for a five-minute test duration (aqueous deficient dry eye stratum); (b) meibum quality score of ≥ 1 (on a 0 to 3 scale) or meibum expressibility score of > 1 (on a 0 to 3 scale) in either eyelid (evaporative dry eye stratum); and (c) met both entry criteria (a) and (b) (mixed dry eye stratum). (Information on meibum quality and expressibility scores is available in [Appendix 2](#)). Participants were required to discontinue the use of any other artificial tear supplements and use only the study product for the entire study duration. Key exclusion criteria were (a) history of hypersensitivity to the study drug or any of its excipients; (b) use of any topical ocular medication preserved with benzalkonium chloride or other products known to be toxic to the tear film lipid layer, or lid hygiene therapy or punctal plug insertion within 1 month prior to the screening visit; (c) contact lens use within a week prior to screening visit; (d) initiation of any topical ocular medication (with the exception of artificial tears/gels/lubricants) ≤ 2 weeks prior to the screening visit. Artificial tears/lubricants were prohibited from the beginning of the baseline visit until the patient was discontinued from the study.

Objectives and Endpoints

The primary objective was to evaluate improvement in tear film stability (ie, break-up) post-treatment with PG-HPG based on the change from baseline in TFBUT at Day 14. The secondary objectives of improvement of tear film stability, ocular discomfort, and improvement in corneal staining score with PG-HPG were assessed based on change from baseline in TFBUT at Day 28, in ocular

discomfort Visual Analog Scale (VAS) score at Day 14, and in cornea staining score at Day 28, respectively. Data on adverse events (AEs) were collected throughout the study.

Subgroup Analyses

Subgroup analyses by dry eye subtypes (aqueous deficient, evaporative, or mixed) were performed on the primary and secondary endpoints.

Assessments

Assessments during each visit were conducted at the same time of day within ± 1 hour of the start time of Visit 1 (baseline). TFBUT was measured 30 minutes following administration of study medication at screening, and Days 14 and 28 using an ocular staining dye, sodium fluorescein (NaFl). Two microliters of NaFl solution were instilled in the right eye, immediately followed by three consecutive TFBUT measurements. The same procedure was followed for the left eye. The time from the last blink until one or more black (dry) spots appeared in the pre-corneal tear film was measured.

Ocular discomfort was evaluated using patient-reported outcome (PRO) assessments using VAS at baseline and Day 14 as per the study protocol. Study participants indicated their level of discomfort/level of severity on a 0–100 scale; a higher score indicated more frequent discomfort/higher dry eye severity ([Appendix 3](#)).

Corneal staining score was assessed using NaFl, at screening, and Days 14 and 28 (or early exit) according to the procedure recommended by Bron et al.⁹

Details of the AEs and serious AEs (SAEs) were collected throughout the study.

Statistical Analysis

One hundred and twenty eligible patients were needed to estimate a 95% confidence interval (CI) with a precision of 0.38 and a standard deviation (SD) of 2.1 for the mean change from baseline at Day 14 in TFBUT (primary efficacy analysis). Considering a dropout rate of approximately 10%, up to 134 participants were scheduled to be enrolled.

Descriptive statistics (mean, median, SD, minimum and maximum values) were presented for continuous variables (TFBUT, and ocular discomfort VAS score). In addition, a 95% CI using Student's *t*-distribution was calculated for mean change from baseline at Day 14 in TFBUT. A 95% CI for the median using the Hodges–Lehmann estimator was

provided if the normality criteria were not met. Corneal staining was considered as a categorical measure.

Efficacy analysis was based on the full analysis set (FAS). Both FAS and safety analysis sets consisted of all participants who received at least one drop of PG-HPG lubricant eye drops.

The worse eye, defined as the eye with the lower TFBUT value at screening, of the participant was chosen for analysis. If both eyes had the same TFBUT value, then the right eye was considered the worse eye.

Data on AEs and SAEs were collected and analyzed for each eye of the participant.

All analyses were performed using SAS[®] statistical software (Version 9.4).

Results

A total of 183 participants were evaluated for eligibility, of whom 134 (73.2%) received treatment and 130 (97%) completed the study. The reasons for study discontinuations were adverse event (2 [1.5%]), withdrawal by the participant (1 [0.7%]), and other (1 [0.7%]) – this participant had a programmed cardiac procedure and was not available for study visits.

Demographic and Baseline Characteristics

The mean (SD) age of participants was 56.6 (14.78) years and the majority of the participants were female (75.4%). The numbers of participants for each of the dry eye subtypes (aqueous deficient, evaporative, and mixed) were similar (Table 1).

Primary Outcome

The mean TFBUT increased from 2.6 seconds at baseline to 4.2 seconds at Day 14 [mean (SD) change: 1.5 (2.80) seconds; the median change was 0.8 seconds (95% CI: 0.52, 1.19)] for the overall cohort. The results were consistent across the dry eye subgroups (Figure 2).

Secondary Outcomes

An increase in TFBUT at Day 14 was maintained through Day 28. In the overall cohort, the mean (SD) change from baseline in TFBUT at Day 28 was 1.4 (2.80) seconds and the median change was 0.8 seconds (95% CI: 0.49, 1.17). The results were comparable in the mixed-dry eye subgroup. The evaporative dry eye subgroup showed the largest change in mean (SD) TFBUT (2.5 [3.94] seconds). At

Table 1 Demographic and Baseline Characteristics (Full Analysis Set)

Characteristics	PG-HPG N=134
Mean (SD) age, years	56.6 (14.8)
Age category, n (%)	
<65 years	88 (65.7)
≥65 years	46 (34.3)
Gender, female, n (%)	101 (75.4)
Race, n (%)	
White	100 (74.6)
Black or African American	23 (17.2)
Asian	6 (4.5)
Other	4 (3.0)
Multiracial	1 (0.7)
Dry eye subtype - n (%)	
Aqueous deficient	41 (30.6)
Evaporative	44 (32.8)
Mixed	49 (36.6)
Mean (SD) ocular discomfort VAS score	45.9 (24.33)
Mean (SD) TFBUT, seconds	2.6 (1.01)

Notes: Ocular discomfort VAS score = square root (VAS frequency score x VAS severity score). All baseline characteristics were assessed before instillation. Percentages are based on the number of participants in the full analysis set. Full analysis set consisted of all participants who received at least one drop of PG-HPG lubricant eye drops.

Abbreviations: PG-HPG, propylene glycol-hydroxypropyl guar lubricant eye drops; SD, standard deviation; TFBUT, tear film break-up time; VAS, visual analog scale.

Day 28, the aqueous deficient subgroup showed a marginal improvement of 0.6 (1.47) seconds in mean TFBUT from baseline (median change 0.3 seconds, 95% CI –0.07, 0.88, Figure 3).

The ocular discomfort VAS score improved (decreased) from baseline at Day 14 (mean [SD] change –17.3 [24.80]) in the overall cohort. The dry eye subgroups followed a similar trend (Figure 4).

The cornea staining scores showed an improvement at Day 28 from baseline in the overall cohort as well as the dry eye subgroups (Table 2). Severe corneal staining was not reported during the study.

Safety Outcomes

Overall, 19 treatment-emergent AEs were reported in 9 (6.7%) participants. Ocular AEs were reported in 6 (4.5%) participants and are summarized in Table 3. One AE of blurred vision was reported; it was mild, transient, and was not considered to be related to the product per the investigator. Non-ocular AEs were

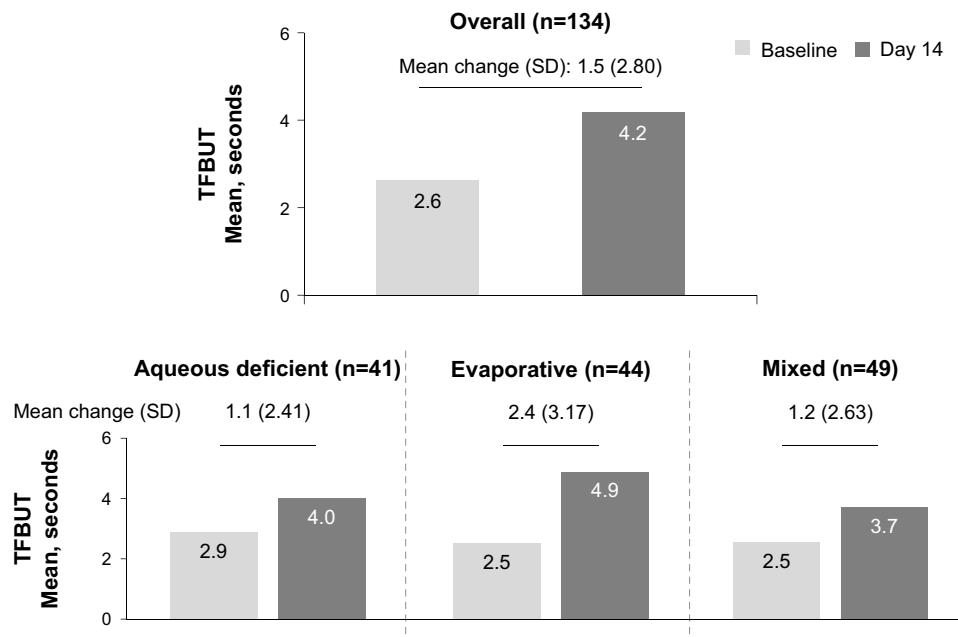


Figure 2 Change from baseline in TFBUT up to Day 14 in the overall cohort and dry eye subtypes (full analysis set). The 95% CI was calculated using Hodges–Lehmann estimator. Full analysis set consisted of all participants who received at least one drop of PG-HPG lubricant eye drops.

Abbreviations: CI, confidence interval; PG-HPG, propylene glycol-hydroxypropyl guar lubricant eye drops; TFBUT, tear film break-up time.

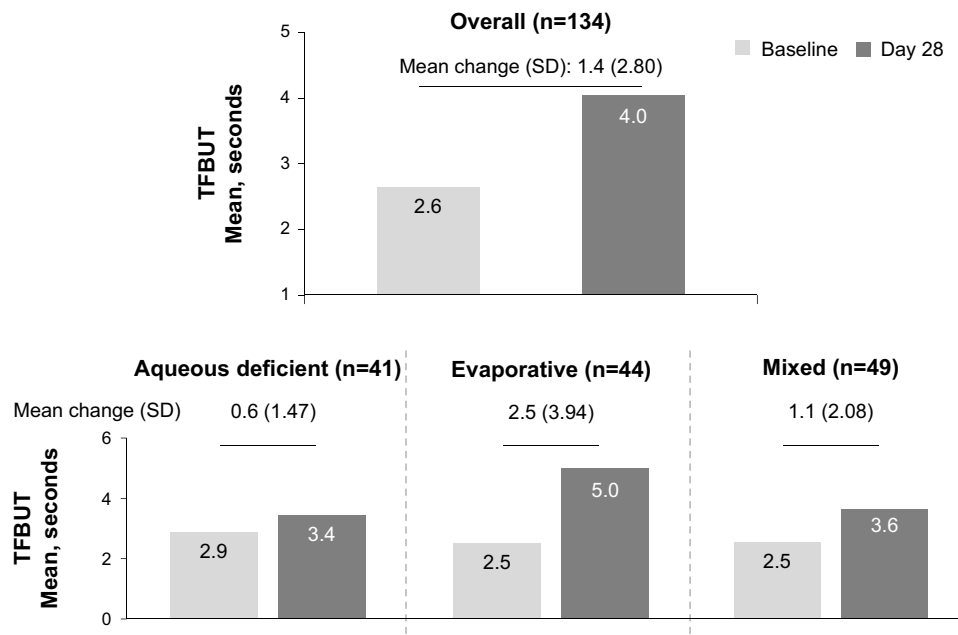


Figure 3 Change from baseline in TFBUT up to Day 28 in the overall cohort and dry eye subtypes (full analysis set). The 95% CI was calculated using Hodges–Lehmann estimator. Full analysis set consisted of all participants who received at least one drop of PG-HPG lubricant eye drops. Baseline was defined as the last available, non-missing, scheduled or unscheduled value collected prior to exposure to study treatment.

Abbreviations: CI, confidence interval; PG-HPG, propylene glycol-hydroxypropyl guar lubricant eye drops; TFBUT, tear film break-up time.

reported in 8 (6%) participants (headache in 3 [2.2%], nasopharyngitis in 2 [1.5%], influenza-like illness, arthritis and musculoskeletal chest pain in 1 (0.7%)

patient each). None of the ocular and non-ocular AEs reported were severe. Two participants discontinued the treatment owing to ocular AEs (eye pruritus and viral

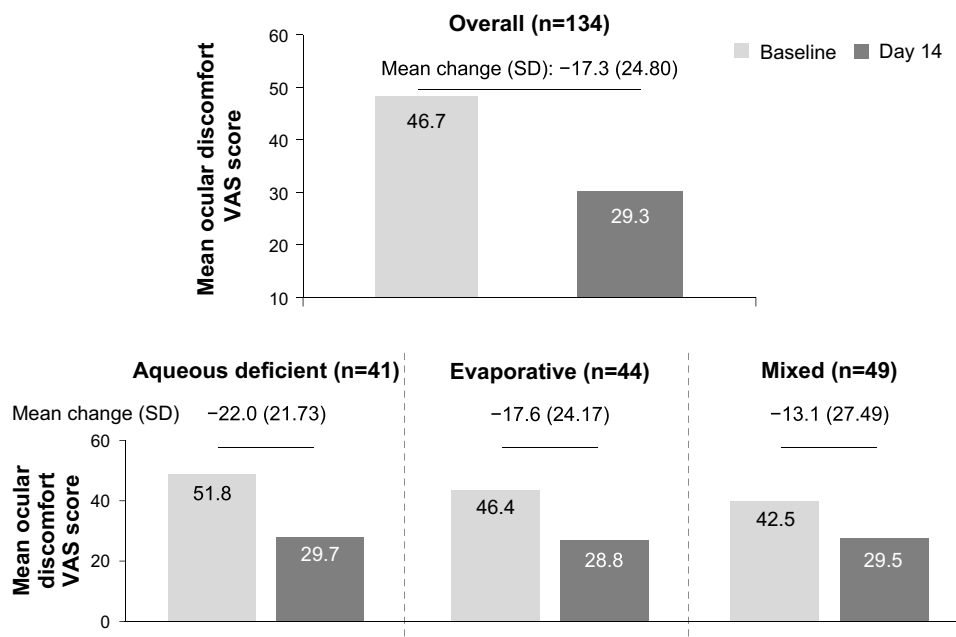


Figure 4 Change from baseline in ocular discomfort VAS score at Day 14 in the overall cohort and dry eye subtypes (full analysis set). Full analysis set consisted of all participants who received at least one drop of PG-HPG lubricant eye drops. Baseline was defined as the last available, non-missing, scheduled or unscheduled value collected prior to exposure to study treatment.

Abbreviations: PG-HPG, propylene glycol-hydroxypropyl guar lubricant eye drops; SD, standard deviation; VAS, visual analog scale.

conjunctivitis; 1 [0.7%] patient each). SAEs were not reported during the study.

Discussion

The results of this clinical study demonstrated that treatment with PG-HPG nanoemulsion-based lubricant eye drops enhanced the tear film stability in participants with DED, thereby meeting its primary objective. This improvement was sustained through Day 28. In addition, PG-HPG also improved the signs and symptoms of participants with DED, and was considered to be well tolerated throughout the study.

PG-HPG nanoemulsion-based lubricant eye drop is the first Systane[®] product that combines the benefits of HPG and nanoparticle technology (reduced lipid droplet size <100 nm) to replenish both the aqueous and lipid deficiencies of the tear film, potentially targeting all subtypes of dry eye. Following application of PG/HPG-nanoemulsion, HPG and borate form a cross-linked matrix which acts as a protective viscoelastic barrier on the ocular surface epithelium. This barrier improves the retention of the demulcent (PG) on the aqueous/mucin layer and moisture content of the deficient aqueous tear film, and also acts as a depot for slow release of lipid into the tear film, to supplement and stabilize any gaps due to lipid insufficiency. The anionic phospholipids in

PG-HPG can be beneficial for the proper structuring of the polar lipid layer and maintaining tear film stability by acting as an effective interface between the outer non-polar lipids and the underlying aqueous phase as evident from previous studies.^{8,10} Thus, PG-HPG-nanoemulsion restores the complete tear film structure to prevent exacerbations of DED and maintain a healthy ocular surface.⁷

The beneficial effect of HPG technology facilitating the replenishment of the deficient tear film is well reported.^{11–14} Systane[®] Balance (Alcon Laboratories, Inc. Fort Worth, TX, USA), which is an oil-based HPG emulsion product, helps to replenish the lipid layer in a lipid-deficient tear film.¹¹ Systane[®] and Systane[®] Ultra lubricant eye drops (Alcon Laboratories, Inc. Fort Worth, TX, USA) utilize the HPG technology and provide long-lasting relief from symptoms of dryness of the eye and sustained comfort.^{12–14}

PG-HPG utilizes the same active demulcent as Systane[®] Balance, ie, PG, but with increased concentration of HPG which allows for better crosslinking with borate. PG-HPG has been uniquely processed compared to Systane[®] Balance. The smaller nano-sized lipid droplets in PG-HPG optimize the lipid surface coverage and make the emulsion formulation less opaque compared with other Systane[®] lubricant eye drops, reducing blurred vision upon instillation of ocular lubricants.^{4,13,15}

Table 2 Change in Cornea Staining Score from Baseline to Day 28 (Full Analysis Set)

Baseline Score, n		PG-HPG (N=134) Post-Baseline (Day 28) Scores					
		Absent n (%)	Minimal n (%)	Mild n (%)	Moderate n (%)	Marked n (%)	Missing n (%)
Full cohort^a							
Absent	48	36 (75.0)	9 (18.8)	2 (4.2)	0	0	1 (2.1)
Minimal	43	14 (32.6)	24 (55.8)	4 (9.3)	0	0	1 (2.3)
Mild	34	5 (14.7)	12 (35.3)	14 (41.2)	1 (2.9)	1 (2.9)	1 (2.9)
Moderate	9	2 (22.2)	2 (22.2)	1 (11.1)	3 (33.3)	0	1 (11.1)
Marked	0	0	0	0	0	0	0
Total	134	57 (42.5)	47 (35.1)	21 (15.7)	4 (3.0)	1 (0.7)	4 (3.0)
Aqueous deficient^a							
Absent	10	6 (60)	3 (30.0)	1 (10.0)	0	0	0
Minimal	17	7 (41.2)	9 (52.9)	1 (5.9)	0	0	0
Mild	12	2 (16.7)	5 (41.7)	4 (33.3)	0	1 (8.3)	0
Moderate	2	0	1 (50.0)	1 (50.0)	0	0	0
Total	41	15 (36.6)	18 (43.9)	7 (17.1)	0	1 (2.4)	0
Evaporative^b							
Absent	23	18 (78.3)	4 (17.4)	1 (4.3)	0	0	0
Minimal	12	5 (41.7)	6 (50.0)	1 (8.3)	0	0	0
Mild	8	1 (12.5)	1 (12.5)	5 (62.5)	1 (12.5)	0	0
Moderate	1	0	0	0	0	0	1 (100)
Total	44	24 (54.5)	11 (25.0)	7 (15.9)	1 (2.3)	0	1 (2.3)
Mixed^b							
Absent	15	12 (80.0)	2 (13.3)	0	0	0	1 (6.7)
Minimal	14	2 (14.3)	9 (64.3)	2 (14.3)	0	0	1 (7.1)
Mild	14	2 (14.3)	6 (42.9)	5 (35.7)	0	0	1 (7.1)
Moderate	6	2 (33.3)	1 (16.7)	0	3 (50.0)	0	0
Total	49	18 (36.7)	18 (36.7)	7 (14.3)	3 (6.1)	0	3 (6.1)

Notes: ^aThere were no participants with severe cornea staining score at baseline and Day 28. ^bThere were no participants with marked or severe cornea staining score at baseline and Day 28. Percentages are based on the number of participants with worse eye in the respective baseline score category in the full analysis set. Full analysis set consisted of all participants who received at least one drop of PG-HPG lubricant eye drops.

Abbreviations: PG-HPG, propylene glycol-hydroxypropyl guar lubricant eye drops; TFBUT, tear film break-up time.

Tear film stability, which is essential for ocular surface homeostasis and functioning, is disturbed in DED.¹⁶ The TFBUT, constituting the time interval elapsed between a complete blink and the appearance of the first break in the tear film, is a direct test of tear film stability and a valuable diagnostic aid in DED.¹⁷ In the present study, treatment with PG-HPG nanoemulsion elicited an increase of 1.5 seconds (59% improvement) from baseline (2.6 seconds) in mean TFBUT at Day 14. A previous study has reported that a decrease in one second of TFBUT can have a major impact on ocular discomfort in >70% of DED patients.¹⁸ Thus, considering the low baseline value, the increase in TFBUT (>1 second) observed with short-term use of PG-HPG nanoemulsion, though modest, is yet meaningful for DED patients. However, the

variability observed in TFBUT outcomes may be attributed to factors (eg, residual tear volume, environmental conditions, etc.) that are known to influence TFBUT measurements.¹⁹ Earlier studies have reported that HPG in Systane[®] formulations interacts with the divalent ions and mucin in the natural tear film to form a gel-matrix, thus contributing to the tear film stability.^{12,13,20} The results of this study demonstrate the ability of PG-HPG to promote tear film stability in participants with DED, in agreement with earlier studies which showed an increased TFBUT with Systane[®] Balance in patients with evaporative DED and with Systane[®] lubricant eye drops in patients with DED.^{13,21}

The symptoms of ocular surface disorder improved after PG-HPG instillation – the ocular discomfort VAS

Table 3 Proportion (%) of Participants Reporting Ocular Adverse Events (Safety Analysis Set)

System Organ Class Preferred Terms	PG-HPG N=134
Eye disorders	5 (3.7)
Halo vision	1 (0.7)
Conjunctival hyperemia	1 (0.7)
Eye pruritus	1 (0.7)
Eyelid margin crusting	1 (0.7)
Vision blurred	1 (0.7)
Infections and infestations	1 (0.7)
Conjunctivitis viral	1 (0.7)

Notes: A patient with multiple AEs within a preferred term was counted only once for that preferred term. MedDRA Version 20.1 was used for the reporting of AEs. Safety analysis set consisted of all participants who received at least one drop of PG-HPG lubricant eye drops.

Abbreviations: AE, adverse event; PG-HPG, propylene glycol-hydroxypropyl guar lubricant eye drops.

score decreased at Day 14, and there was a reduction in corneal staining score at Day 28 relative to baseline. The improvement in dry eye symptoms may be attributed to the increase in tear film stability and the beneficial effect of the HPG technology – longer retention of the demulcent, thus reducing the friction between lid and ocular surface. Lipid nanoparticles in this formulation are designed for better lipid coverage of the tear film, contributing to a reduction in the ocular discomfort of patients suffering from DED. These results indicate that PG-HPG nanoemulsion can provide symptomatic relief in patients with DED and are consistent with those reported with other Systane® eye drops.^{11,14,22}

Findings in the subgroups based on the dry eye subtypes were comparable to that of the overall study cohort. The only exception was the change in TFBUT at Day 28 for the aqueous subgroup, which showed a nominal change from baseline based on 95% CI values (−0.07, 0.88). The effect on TFBUT was pronounced in the evaporative subgroup as compared to other subgroups, possibly because PG-HPG was formulated by processing of ingredients similar to that of Systane® Balance lubricant eye drops, which have a known efficacy in patients with evaporative DED. Previous studies have shown that anionic phospholipids can provide a stable interface between nonpolar lipids at the surface of the tear film and the aqueous layer, and polar lipids can enhance the lipid layer thickness.^{6,10} The reduced droplet size of nanoemulsion is expected to serve as a better delivery vehicle for lipids to the ocular surface to help optimize the evaporation rate of the tear film. However, studies involving

a larger population are warranted to further ascertain the efficacy across all dry eye subtypes.

Selection of an appropriate ocular lubricant is influenced by the cause of DED (aqueous deficiency, lipid deficiency, or both) and is critical for the management of DED.⁴ Patients with evaporative DED require ocular lubricants, which reduce the rate of tear evaporation. In addition to treating the root cause, for patients with meibomian gland dysfunction, replenishment of the tear film with lipids is important. PG-HPG nanoemulsion treatment acts by replenishing the lipid layer and leads to an increase in tear film stability, potentially reducing the rate of tear evaporation in patients with evaporative dry eye. Patients with aqueous deficient DED have low tear volume and require treatments that can sufficiently hydrate the ocular surface.⁴ The lubricant and HPG technology in PG-HPG nanoemulsion reduce the ocular discomfort in these aqueous-deficient patients. If only one of these two broad categories of ocular surface disease (evaporative dry eye and aqueous deficient dry eye) is addressed therapeutically, patients may continue to experience symptoms and report dissatisfaction with the prescribed treatment.⁴ PG-HPG nanoemulsion-based lubricant eye drops can serve as a total tear replacement technology in all types of DED.

Treatment of participants with PG-HPG nanoemulsion was well tolerated. The proportion of patients with blurred vision upon instillation of PG-HPG was low in comparison with other Systane® eye drops.^{13,15} HPG-based ocular lubricants usually cause blurred vision.^{13,15} However, in this study the investigator did not consider blurred vision to be treatment-related. This may be attributed to the small nano-sized droplets in PG-HPG nanoemulsion, which make it less opaque. Neither SAEs nor severe AEs were reported, similar to that known for other Systane® eye drops.^{11,23,24}

The strength of this study is that data were collected from six sites across different countries. The study is limited by the invasiveness of fluorescein TFBUT measurements typically used by clinicians. Further, this was a single-arm study; a comparison of PG-HPG with existing market products was not performed due to the unique combination of HPG and nanotechnology in this formulation. The effect of vehicle in PG-HPG nanoemulsion has been studied earlier in a preclinical study demonstrating better (32–40 times) hydration protection, lubricity, and polymer film break-up time (all $P < 0.05$) as compared to vehicle (nanoemulsion without HPG) in human corneal

epithelial cells.⁸ However, the inclusion of a comparator/placebo arm can be considered in future clinical studies.

Conclusions

Our study findings showed that PG-HPG nanoemulsion-based lubricant eye drops improved the tear film stability, signs and symptoms of DED, and were well tolerated in participants with DED irrespective of its subtypes.

Data Sharing Statement

Due to the varying rights of individuals and contractual rights of parties involved, Alcon does not make a practice of sharing datasets.

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Disclosure

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Alcon. The authors report no other conflicts of interest in this work.

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