

Comparative Analysis of the Osmoprotective Effects of Daily Disposable Contact Lens Packaging Solutions on Human Corneal Epithelial Cells

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Purpose: Contact lens (CL) wear challenges the balance of the ocular surface environment by increasing water evaporation and tear osmolarity. Maintaining ocular surface homeostasis during CL wear remains a goal of lens manufacturers and an important consideration for eye care professionals. The purpose of this study was to measure the metabolic activity and inflammatory responses of a transformed human corneal epithelial cell (THCEpiC) line under hyperosmotic conditions in the presence of CL packaging solutions.

Methods: CL packaging solutions sampled from seven daily disposable silicone hydrogel CL blister packages were prepared at 25% and made hyperosmolar (400 mOsm/kg) with NaCl. THCEpiCs were incubated with each solution for 24 hr, after which cell culture supernatants were collected. THCEpiC metabolic activity was determined by an alamarBlue assay. Concentrations in cell culture supernatants of inflammatory cytokine (interleukin [IL]-6) and chemokine (IL-8), as well as monocyte chemoattractant protein-1 (MCP-1), were quantitated by specific enzyme-linked immunosorbent assays.

Results: THCEpiC metabolic activity under hyperosmolar conditions decreased in the presence of somofilcon A and senofilcon A solutions ($p=0.04$ and 0.004 , respectively), but no other solution (all $p\geq 0.09$). Concentrations of IL-6 increased in the presence of delefilcon A, somofilcon A, narafilcon A, and senofilcon A solutions (all $p\leq 0.001$), but no other solution (all $p\geq 0.08$), while those of IL-8 increased in the presence of all solutions (all $p\leq 0.03$) but kalifilcon A ($p>0.99$), and those of MCP-1 increased in the presence of delefilcon A, verofilcon A, somofilcon A, and stenfilcon A solutions (all $p<0.0001$), but no other solution (all $p>0.99$).

Conclusion: CL packaging solutions differ in their capacity to inhibit epithelial inflammation. THCEpiC inflammatory response was less in the presence of a CL packaging solution containing osmoprotectants than in solutions lacking osmoprotectants under moderately hyperosmolar conditions in vitro. Clinical studies are warranted to further substantiate the benefit of osmoprotectants.

Keywords: contact lens, cornea, TFOS, hyperosmolarity, osmoprotection, homeostasis

Introduction

Contact lens (CL) is one of the most widespread and successfully used medical devices available today. The Centers for Disease Control and Prevention (CDC, Atlanta, GA) estimated that about 45 million people older than 17 wore CLs in the United States,¹ and the Tear Film and Ocular Surface Society (TFOS) International Workshop on Contact Lens Discomfort estimated that over 140 million people wore CLs worldwide.² Recent advances in lens design and manufacturing with the goal of maintaining ocular surface homeostasis have led to newer CLs with the potential to better maintain eye health and comfort during CL wear. This includes more benign interactions between the lens material, the tear film, and the ocular surface with which it makes intimate contact.

A 2017 report of the TFOS International Dry Eye Workshop II (TFOS DEWS II) defined dry eye disease (DED) as

a multifactorial disease of the ocular surface characterized by a loss of homeostasis of the tear film, and accompanied by ocular symptoms, in which tear film instability and hyperosmolarity, ocular surface inflammation and damage, and neurosensory abnormalities play etiological roles.³

The report attributes DED to reduced tear secretion leading to inflammation and peripheral nerve alteration, the former activating nociceptor nerve endings responsible for the sensation of dryness and pain.⁴ While the exact relationship between dryness-mediated cellular and molecular inflammatory mechanisms and peripheral nociceptor terminals is not completely understood, such receptors might respond to both the mechanical stress of tear breakup and the hyperosmotic stress of tears following tear film evaporation.⁴

DED symptoms can occur both in the presence or absence of CLs, the former including CL-induced dry eye (CLIDE, characterized by DED signs and symptoms that did not exist prior to CL wear) and CL-associated dry eye (CLADE, characterized by DED signs and symptoms during CL wear).⁵ The TFOS DEWS II report summarizes the consequences of CL wear that lead to CLIDE and CLADE, including a thinner, patchy lipid layer, poor wettability with impaired spreading, tear film instability, increased tear evaporation rate, lower basal tear turnover rate, increased tear osmolarity, and reduced tear film meniscus volume, as well as biochemical changes within the tear film.⁵ CL wear further challenges ocular surface homeostasis by disrupting the tear film, splitting it into distinct pre-lens and post-lens films,^{6,7} which can lead to undesirable physical and chemical interactions between the lens, the tear film, and the ocular surface. This manifests as evaporation from the pre-lens tear film and the deleterious consequences of tear film thinning.⁸ The TFOS DEWS II report additionally cites tear hyperosmolarity and inflammatory mediators as causes of DED symptoms and ocular cell damage and dysfunction.⁹ McMonnies proposed an amplifying evaporative dryness cascade that occurs during CL wear, culminating in hyperosmolar tears, CL dryness, and ocular surface inflammation;¹⁰ the latter occurs in response to hyperosmolar tears,¹⁰ but proteins sorbed on and denatured at the CL surface,¹¹ as well as mechanical irritation by a dehydrated and dewetted lens, can also play a role. CLs that best maintain water at the surface and throughout the lens to ameliorate hyperosmolarity if it does occur might help prevent or reduce ocular surface inflammation, as this may prevent the dryness cycle leading to loss of homeostasis.¹²

Tear hyperosmolarity was recognized as a core mechanism of DED prior to DEWS II,^{13,14} but DEWS II recognized loss of tear film homeostasis as the unifying element.³ Hyperosmolar tears are reported to disturb ocular homeostasis primarily by affecting ocular inflammation via corneal epithelial cell expression of inflammatory cytokines, chemokines, and matrix metalloproteinases (eg MMP-9).^{15–19} This occurs through activation of mitogen-activated protein kinase p38, c-Jun N-terminal kinase, nuclear factor- κ B, and activator protein-1.²⁰ Hyperosmolarity also leads to loss of barrier function,^{21,22} increases corneal epithelial transient receptor potential vanilloid type-1 channel activity,²³ affects cellular apoptosis,²⁴ and disrupts normal cellular metabolism.²⁵ Additionally, hyperosmolar stress is reported to disrupt neuroimmune homeostasis of the ocular surface by reducing the density of corneal intraepithelial nerves and terminals and sensitizing the ocular surface to hypertonicity,²⁶ as well as induce functional and structural alterations of corneal nerves.²⁷ Hyperosmolar stress can also inhibit neutrophil activity, reducing protection against corneal infection.²⁸

Cytokine interleukin (IL)-6 and chemokines IL-8 and monocyte chemoattractant protein-1 (MCP-1) are significant pro-inflammatory mediators whose concentrations are frequently reported to increase significantly in the tears of DED patients.^{29–31} However, clinical testing is not always feasible for evaluating novel ocular formulations, and cellular models have proven to be more efficient.

Further, while physical and chemical characterizations of many contact lens packaging and care solutions (eg, pH, osmolarity, surface tension, and viscosity) have been reported in literature,^{32,33} neither the clinical significance of each individual property nor the combination of all properties is completely understood. As such, *in vitro* cell line models are frequently employed for multi-purpose solution (MPS) cytotoxicity studies and for evaluating the potential effects of ocular formulation ingredients due to their availability and reproducibility. Numerous studies have demonstrated that human corneal epithelial cells exposed to hyperosmolar and desiccation stresses induced significant IL-6, IL-8, and MCP-1 responses,^{34,35} as such, these pro-inflammatory mediators were selected for measurement in this study.

Fortunately, several osmolytes have been found to be effective in osmoregulation and mitigation of the deleterious effects of tear hyperosmolarity;^{36,37} these include amino acids and their derivatives such as proline, betaine, and L-carnitine, as well

as polyols such as erythritol and glycerol, and sugars and their derivatives such as trehalose.^{16,23,24,36–51} Including these osmotic management agents in ocular solutions like artificial tears, CL MPSs, and CL rewetting drops may help ameliorate the effects of tear film evaporation and maintain ocular surface homeostasis. Studies in literature report decreased symptoms of eye dryness and improved comfort in CL wearers using rewetting drops containing osmoprotectants.^{50,51}

In many CL packaging solutions, buffering agents like sodium chloride and sodium salts, such as sodium borates and sodium phosphates, contribute the most to the total solution osmolarity. Substituting potassium for sodium reduces the concentration of sodium, which is the electrolyte whose concentration becomes elevated in lacrimal gland disease.⁵² Additionally, including organic osmolytes in the solution allows for further reduction in sodium concentration.

Lens manufacturers incorporate ingredients into their CLs, including daily disposable (DD) silicone hydrogel lenses,⁵³ through their respective solutions to support the lens,⁵⁴ including one that contains osmoprotectants. The delefilcon A and verofilcon A (both Alcon Laboratories, Inc., Ft. Worth, TX) solutions contain copolymers of poly-amidoamine and poly(acrylamide-acrylic) acid;^{55,56} the kalifilcon A (Bausch & Lomb Incorporated, Rochester, NY) solution contains poloxamer 181, poloxamine 1107, glycerin, and erythritol;⁵⁷ the somofilcon A and stenfilcon A (both CooperVision Inc., Pleasanton, CA) solutions contain poloxamer 407 and polysorbate, respectively;^{58,59} and the narafilcon A and senofilcon A (both Johnson & Johnson Inc., Jacksonville, FL) solutions contain methyl ether cellulose^{60,61} (Table 1). The objectives of this study were as follows: 1. To determine if ingredients for managing osmolarity affect metabolic activity and cytokine/chemokine levels of human corneal epithelial cells under hyperosmotic stress and 2. To measure and compare metabolic activity and cytokine/chemokine levels of the same cell line exposed to DD silicone hydrogel CL packaging solutions with and without osmoprotectants under hyperosmotic conditions.

Materials and Methods

Cell Line

A human corneal epithelial cell transformed with simian virus 40 to corneal epithelial cell line 2.040 pRSV-T (ATCC[®] CRL-11516[™]) (THCEpiC) was obtained from the American Type Culture Collection (ATCC, Manassas, VA). Such cell lines are commonly used in studies of corneal epithelium,^{62,63} as they do not vary phenotypically compared with primary cells. THCEpiCs were received at passage 22 and used between passages 25 and 40 for these experiments. Cells were grown in Gibco[™] EpiLife[™] Medium (Catalog number MEPI500CA, Thermo Fisher Scientific, Waltham, MA) with human corneal growth supplement (Catalog number S0095, Thermo Fisher Scientific) and 50 IU/mL penicillin/50 µg/mL streptomycin (Catalog number 15-140-122, Thermo Fisher Scientific).

Table 1 Contact Lenses Evaluated

Material	Water Content	ISO Group ⁵³	Manufacturer's Solution Description	Brand Name	Manufacturer
Delefilcon A ⁵⁵	≥80% (surface) 33% (bulk)	V-C	PBS solution with approximately 0.3% of polymeric wetting agents consisting of copolymers of polyamidoamine and poly (acrylamide-acrylic) acid	Dailies Total1	Alcon Laboratories, Inc. (Ft. Worth, TX)
Verofilcon A ⁵⁶	≥80% (surface) 51% (bulk)	V-B		Precision 1	
Kalifilcon A ⁵⁷	55%	V-B	PBS solution with potassium chloride, poloxamine, poloxamer 181, glycerin, and erythritol	Infuse/Ultra One Day	Bausch & Lomb Incorporated (Rochester, NY)
Somofilcon A ⁵⁸	56%	V-B	Borate buffered saline solution containing 0.005% w/v poloxamer 407	Clariti-1 day	CooperVision, Inc. (Pleasanton, CA)
Stenfilcon A ⁵⁹	54%	V-B	PBS solution with Tween	MyDay	
Narafilcon A ⁶⁰	46%	V-C	Buffered saline solution with methyl ether cellulose	1-day Acuvue TruEye with HydraClear	Johnson & Johnson Vision Care, Inc. (Jacksonville, FL)
Senofilcon A ⁶¹	38%	V-C		Acuvue 1-day Oasys with HydraLuxe	

Abbreviations: ISO, International Organization for Standardization; PBS, phosphate-buffered saline.

Cell Culture

Cells were cultured in a supplemented EpiLife medium in a 5% CO₂/95% humidity incubator at 37°C for 72 hr for all experiments. The medium was then replaced with a fresh medium, and the cells were incubated for additional 24 hours under the same conditions prior to incubation with test solutions.

Preparation of Hyperosmolar Test Media

Hyperosmolar media (400, 450, 500, and 530 mOsm/kg) with or without 0.9% erythritol + 0.9% glycerin osmoprotectants added were prepared from basal medium by addition of NaCl.

Preparation of Hyperosmolar Contact Lens Solutions

Preliminary experiments indicated that 50% CL packaging solutions significantly affected cell metabolic activity in 450 and 500 mOsm/kg hyperosmolar media. Therefore, CL packaging solutions were sampled from delefilcon A, verofilcon A, kalifilcon A, somofilcon A, stenfilcon A, narafilcon A, and senofilcon A blister packages (Table 1); for each lens, three to four packages of solution from same lot number were pooled, diluted to 25%, and made hyperosmolar [400 mOsm/kg] by addition of NaCl. Hanks' Balanced Salt Solution (HBSS) diluted to 25% was prepared as the control.

Experimental Procedure

THCEpiCs cultured to confluency in 96 well plates were incubated for 24 hr with 0.25 mL of normal osmolar (315 mOsm/kg) and hyperosmolar media (450, 500, and 530 mOsm/kg), with or without osmoprotectants to determine the effect of osmolarity and osmoprotection on metabolic activity. Note that 315 mOsm/kg tonicity of normal medium is slightly hypertonic relative to normal tears, in which a value of 316 mOsm/L was the threshold proposed as the "osmolarity referent" in the diagnosis of keratoconjunctivitis sicca, compared with an average value of 302 ± 9.7 mOsm/L for normal tears across 16 published studies;⁶⁴ subsequently, the TFOS DEWS report proposed 308 mOsm/L as a sensitive threshold for early-stage DED.²⁸ More recently, 316 mOsm/L in either eye, 308 mOsm/L in both eyes, or >8 mOsm/L inter-eye difference was proposed as a pathological cut-off.^{65,66} It should be noted that these values refer to osmolarity of tears sampled from the meniscus and that local, transient osmolarity in areas of high evaporation rate and tear break-up can be as high as 600–900 mOsm/L.^{67,68} In this study, isotonic refers to normal media rather than normal tears, regardless of the definition of "normal" tear osmolarity.

After determining the osmolarity at which cell metabolism decreases relative to normal medium (500 mOsm/kg), as well as the osmolarity at which osmoprotectants lessen the decrease in cell metabolism relative to unprotected media (530 mOsm/kg), the experiment was repeated using media ranging from normal (315 mOsm/kg) to hypertonic (400, 450, 500 mOsm/kg), ie, concentrations at which respective solutions with and without osmoprotectants do not affect cell metabolism differently. Cell culture supernatants were collected for assessment of cytokine/chemokine activation.

To evaluate the interactions between THCEpiCs and CL solutions, the experiment was repeated using moderately hypertonic (400 mOsm/kg) solutions to increase the likelihood that no CL solution would decrease metabolic activity.

Cell Metabolism and Cytokine/Chemokine Activity Assays

An alamarBlue assay was performed to evaluate THCEpiC metabolic activity incubation with the respective solutions.⁶⁹ Cell culture supernatants were quantitated with specific enzyme-linked immunosorbent assays (ELISAs) for the detection of inflammatory cytokine (IL-6) and chemokines (IL-8, MCP-1) (Quantikine ELISA, R&D Systems, Inc., Minneapolis, MN). IL-6, IL-8, and MCP-1 were chosen as biomarkers of corneal inflammation due to their respective roles in the pathogenesis of DED.⁷⁰ IL-6 affects differentiation of B cells into antibody producing cells,⁷¹ IL-8 affects neutrophil activation in response to inflammatory stimuli,⁷² and MCP-1 acts as a monocyte chemoattractant.⁷³ Hyperosmolar stress has been reported to increase corneal epithelial production of IL-6 and IL-8,^{25,34} as well as MCP-1.³⁴ While evidence that reducing these inflammatory markers improves clinical outcomes is lacking in literature, recent studies demonstrate that anti-IL-6 antibodies reduce inflammatory responses, both in vitro and in a murine model of fungal keratitis,⁷⁴ and alkali burn.⁷⁵

In this study, metabolic activity data were collected from four replicates per test group ($n=4$). Pro-inflammatory cellular response markers were collected from these same test groups.

Statistical Analysis

All test group data sets were first analyzed using descriptive statistics and calculated as the mean values and standard errors. Additional statistical analysis was evaluated for any statistical significance by two-way analysis of variance (ANOVA) followed by Tukey's or Dunnett's multiple comparisons.⁷⁶ Results were considered statistically significant at an alpha level of 5.0% and a p -value ≤ 0.05 . All test group data sets were analyzed using GraphPad Prism version 8.0 for Windows (GraphPad Software, San Diego, CA; www.graphpad.com).

Results

Metabolic activity of THCEpiCs incubated with hyperosmolar media with and without added osmoprotectants is shown in [Figure 1](#). Activity decreased with increased solution osmolarity, the difference from isotonic being significant at 500 and 530 mOsm/kg (both $p \leq 0.0001$). Activity is greater in the presence of osmoprotectants only in the higher osmolar medium (530 mOsm/kg; $p \leq 0.0001$).

Cytokine/chemokine concentrations measured in the supernatants of isotonic and hypertonic media are shown in [Figure 2A–D](#). The addition of glycerol and erythritol osmoprotectants significantly reduced pro-inflammatory cytokine/chemokine response at solution osmolarities of 400 mOsm/kg (MCP-1, $p=0.04$; [Figure 2D](#)) and 450–500 mOsm/kg (IL-6, both $p=0.02$, and IL-8, both $p < 0.0001$; [Figure 2B](#) and [C](#), respectively).

THCEpiC metabolic activity in all isotonic CL solutions was comparable to that in the HBSS control (all $p > 0.05$, data not shown). Metabolic activity in moderately hypertonic (400 mOsm/kg) diluted CL solutions, as well as supernatant cytokine/chemokine responses relative to control, are shown in [Figure 3](#). Activity was lower in hypertonic somofilcon A and senofilcon A CL solutions ($p=0.04$ and 0.004 , respectively), but not kalifilcon A ($p=0.84$), delefilcon A ($p=0.65$), verofilcon A ($p=0.97$), narafilcon A ($p=0.09$), or stenfilcon A ($p=0.45$; [Figure 3A](#)). IL-6 responses were greater in hypertonic solutions of delefilcon A ($p=0.001$), somofilcon A, narafilcon A, and senofilcon A (all $p < 0.0001$), but not kalifilcon A ($p=0.67$), verofilcon A ($p=0.26$), or stenfilcon A ($p=0.08$; [Figure 3B](#)). IL-8 responses were greater in hypertonic solutions of all CLs (delefilcon A, senofilcon A, narafilcon A, and somofilcon A (all $p < 0.0001$), verofilcon A ($p=0.0001$), and stenfilcon A ($p=0.03$) but kalifilcon A ($p > 0.99$; [Figure 3C](#))). MCP-1 responses were greater in

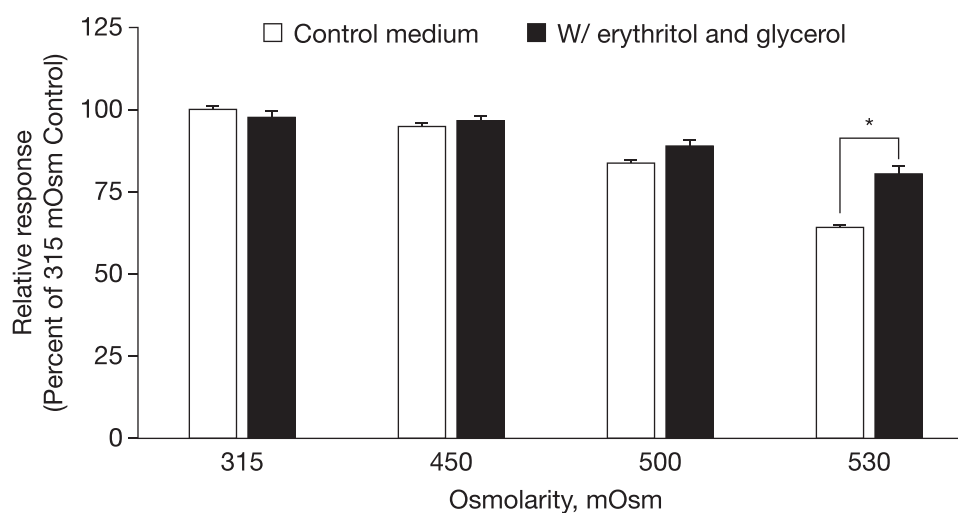


Figure 1 Metabolic activity relative to control medium of THCEpiC cultured in baseline EpiLife control medium until confluent, then incubated with hyperosmolar EpiLife medium in the absence (□ bars) or presence (■ bars) of 0.9% erythritol + 0.9% glycerol (mean \pm standard error). Two-way ANOVA versus equivalent hyperosmolarity without osmoprotectants using Tukey's multiple comparisons (* $p \leq 0.05$).

Abbreviations: ANOVA, analysis of variance; THCEpiC, transformed human corneal epithelial cell.

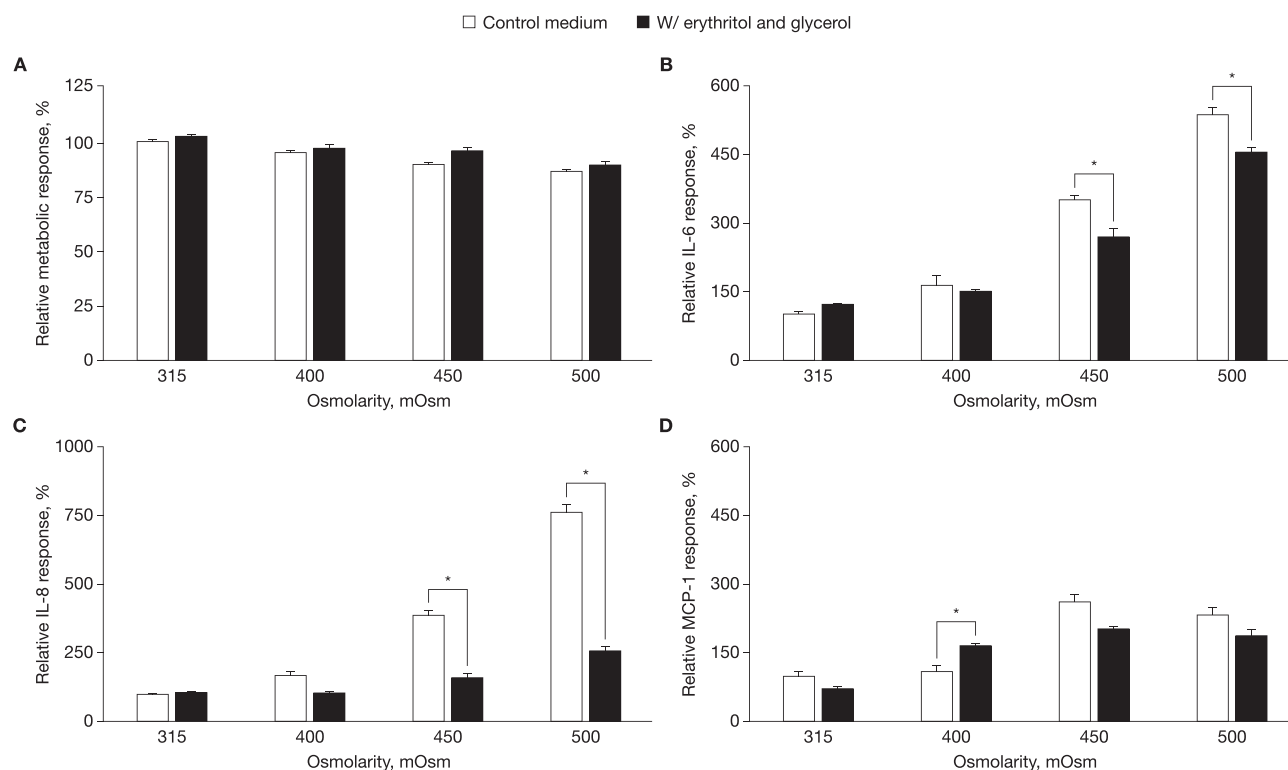


Figure 2 (A) Baseline metabolic activity of THCEpiC cultured in baseline EpiLife control medium until confluent, then incubated with hyperosmolar EpiLife medium in the absence (□ bars) or presence (■ bars) of 0.9% erythritol + 0.9% glycerol relative to 25% HBSS control (mean \pm standard error); (B) Relative IL-6 response; (C) Relative IL-8 response; (D) Relative MCP-1 response. For (A), two-way ANOVA followed by Dunnett's multiple comparisons finds no significant differences in metabolic activity with or without osmoprotection at equivalent osmolarity ($p > 0.05$). For (B–D), two-way ANOVA versus equivalent hyperosmolarity without osmoprotectants followed by Dunnett's multiple comparisons ($*p \leq 0.05$).

Abbreviations: ANOVA, analysis of variance; HBSS, Hanks' Balanced Salt Solution; IL, interleukin; MCP-1, monocyte chemoattractant protein-1; THCEpiC, transformed human corneal epithelial cell.

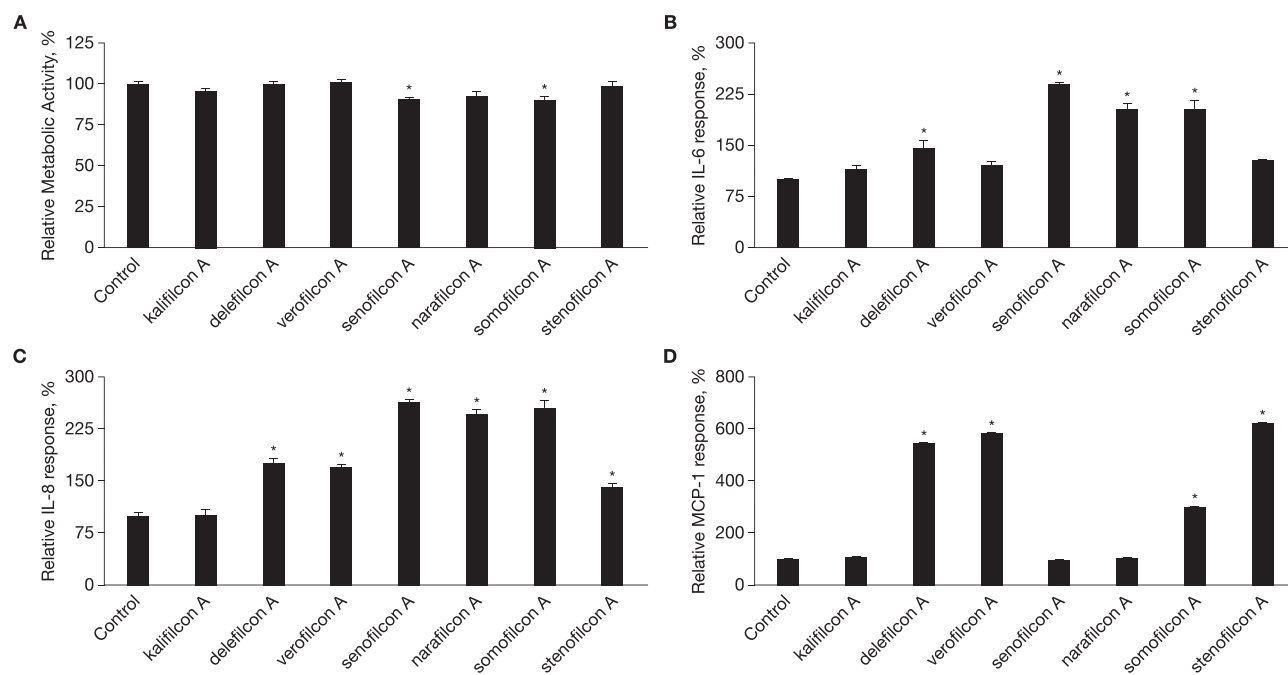


Figure 3 (A) Metabolic activity of THCEpiC cultured in baseline EpiLife control medium until confluent, then exposed to 400 mOsm/kg hyperosmolar CL solutions relative to 25% HBSS control (mean \pm standard error); (B) Relative IL-6 response; (C) Relative IL-8 response; (D) Relative MCP-1 response. Two-way ANOVA followed by Dunnett's multiple comparisons ($* indicates greater than control; all $p \leq 0.05$).$

Abbreviations: ANOVA, analysis of variance; CL, contact lens; HBSS, Hanks' Balanced Salt Solution; IL, interleukin; MCP-1, monocyte chemoattractant protein-1; THCEpiC, transformed human corneal epithelial cell.

hypertonic solutions of delefilcon A, verofilcon A, somofilcon A, and stenfilcon A (all $p < 0.0001$), but not kalifilcon A, narafilecon A, or senofilcon A (all $p > 0.99$, [Figure 3D](#)).

Discussion

Tear concentrations of certain inflammatory markers, notably IL-1 β , IL-6, IL-8, IL-17, LTB $_4$, and MMP-9, are reported to increase over the daily course of CL wear, more so with planned replacement compared with DD lenses.⁷⁷ A study of CL wearers found that in a controlled adverse desiccating environment (5% relative humidity, 23°C, and 0.43 m/s [mean velocity] localized airflow), those subjects whose tears contained the highest levels of inflammatory cytokines IL-4 and IL-6 were associated with increased CL wear symptoms including dryness and discomfort,⁷⁸ which is significant as these are the primary reasons for CL discontinuation.⁷⁹ A similar study found the increase in tear concentrations of some inflammatory mediators to differ between CL models,⁸⁰ possibly reflecting differences in lens polymers and components of their respective packaging solutions. In most CL wearers, ocular surface inflammation is not a major complication of lens wear and is often asymptomatic if it does occur.⁸¹ Nonetheless, it can contribute to discomfort at the end of the day.¹⁰ Most CL-mediated inflammation is reported to be subclinical with DD lens wear.⁸²

With the recognition that maintaining ocular homeostasis during CL wear has the potential to improve lens comfort and vision,⁸³ manufacturers have adopted technologies that support homeostasis. While improving lens water retention during wear represents one approach,^{84,85} infusion of beneficial ingredients into the lens is another.⁸⁶ All lenses evaluated in this study include humectant and/or surfactant wetting agents in the solution; the kalifilcon A lens additionally includes osmoprotectants also used in other ocular solutions,^{36,37} as well as potassium electrolyte.⁵⁷ While physical and chemical characterization of CL packaging solutions is beyond the scope of this study, published literature demonstrates broad ranges in pH, osmolarity, and surface tension of commercial CL blister package solutions, as well as variation in contact angle between different lens/solution combinations, reflective of the surfactants in the solutions and the specific lenses.^{87,88}

THCEpiCs were used in this study to determine the effects of glycerol and erythritol osmoprotectants, as well as CL packaging solutions, upon epithelial stress in hyperosmotic cell culture media. Experiments with isotonic and hyperosmolar cell culture media with and without osmoprotectants highlight two important phenomena. First, THCEpiC metabolic activity in isotonic medium was comparable to that in 450 mOsm/kg medium, while activity in 500 and 530 mOsm/kg media was significantly less; further, the osmoprotectants minimized loss of metabolic activity only in high osmolarity medium (530 mOsm/kg). Second, when cell culture supernatants were analyzed under conditions of equivalent metabolic activity and the same osmolarity, significantly lower cytokine/chemokine responses were observed with osmoprotectants at thresholds of 450 mOsm/kg (IL-6 and IL-8; [Figure 2B and C](#)) and 400 mOsm/kg (MCP-1; [Figure 2D](#)). These results are similar to those of Corrales et al, who reported that osmoprotectants L-carnitine and erythritol, alone or in combination, protected against hyperosmolar stress activation of primary corneal epithelial cells cultured in 400 mOsm/kg media;³⁸ and Hua et al, who reported that L-carnitine, erythritol, and betaine reduced expression of pro-inflammatory cytokines and chemokines by primary corneal epithelial cells cultured in 450 mOsm/kg media,⁴⁰ suggesting that osmoprotectants could play a role in reducing hyperosmotic stress in CL dryness.

Decreased metabolic activity in high osmolarity solutions suggests that investigations of cellular response to hyperosmolar solutions be conducted at osmolarities no greater than 450 mOsm/kg to avoid concomitant responses due to change in cellular metabolism. For this reason, THCEpiC interactions with CL solutions were investigated in a moderately hypertonic medium (400 mOsm/kg). Because the osmolarities of the evaluated CL packaging solutions vary from hypotonic to hypertonic, solutions were first diluted to 25%, then osmolarity was adjusted by the addition of NaCl. THCEpiC metabolic activity was lower in hypertonic (400 mOsm/kg) senofilcon A and somofilcon A CL solutions compared with HBSS control ([Figure 3A](#)), complicating interpretation of cytokine/chemokine response to these solutions. IL-6 concentrations were greater in hypertonic solutions of delefilcon A, somofilcon A, narafilecon A, and senofilcon A compared with control ([Figure 3B](#)). IL-8 concentrations were greater in hypertonic solutions of all CLs but kalifilcon A ([Figure 3C](#)). MCP-1 concentrations were greater in hypertonic solutions of delefilcon A, verofilcon A, somofilcon A, and stenfilcon A ([Figure 3D](#)).

All CL solutions except kalifilcon A elicit at least one cytokine/chemokine response under hyperosmotic conditions. Overall, the kalifilcon A solution elicited the least THCEpiC inflammatory response, likely reflecting in part the presence

of the osmoprotectants, consistent with the results in cell culture media. Previous *in vitro* studies similarly report solution-dependent responses to different CL packaging solutions⁸⁹ and MPSs.^{90,91} The present study demonstrates the potential for osmoprotectants to reduce corneal epithelial stress under hyperosmotic conditions that occur in some CL wearers, particularly at the end of the day.¹⁰

The clinical benefit of osmoprotectants in the packaging solution has not been widely studied. While CLs packaged in solutions lacking osmoprotectants have acceptable performance in asymptomatic wearers and wearers experiencing only minor symptoms while wearing lenses,^{92–96} a recent clinical study found that subjects who reported experiencing CL dryness symptoms with their habitual lenses noticed increased comfort while wearing the kalifilcon A lens.⁸⁶ In addition, a different study reported that CL wearers who used hypo-osmotic saline drops, which reduce tear osmolarity by dilution experienced improved end-of-day comfort and reduced dryness and burning.⁹⁷ We propose that while hyperosmotic stress leading to ocular inflammation in symptomatic patients results in discomfort, inclusion of osmoprotectants in CL packaging solutions can lead to decreased symptoms of discomfort.

While moisturizers and surfactants have previously been used in ophthalmic and CL solutions, the infusion of osmoprotectants into a commercial CL packaging solution is a more recent development. Components included in the kalifilcon A solution were selected based upon recommendations from the TFOS DEWS II report,³⁷ while maintaining ocular surface homeostasis in mind.⁸⁶ Surfactants promote tear film spreading⁹⁸ and stabilize protein structure.⁹⁹ Poloxamine 1107 is reported to be especially effective at suppressing aggregation of heat-denatured lysozyme.¹⁰⁰ Osmoprotectants balance osmotic pressure under osmotic stress, inhibiting inflammation. Glycerin boasts protein stabilizing¹⁰¹ and demulcent properties, while both glycerin and erythritol are included as ingredients in numerous ophthalmic solutions.^{37,102} Potassium electrolyte plays a role in epithelial surface homeostasis.¹⁰³ Further investigation to identify properties of other kalifilcon A solution ingredients and any synergistic effect between them is warranted.

The primary limitation of this study is the *in vitro* nature of the experiments, which do not necessarily predict clinical CL wear outcomes. The ocular surface, which includes multiple cell types, is dynamic and resists internal and external assaults more capably than do exposed cells. In addition, the osmolarities of the solutions evaluated in this study are higher than would typically be found in asymptomatic CL wearers, although transient, local osmolarity can exceed these levels up to three-fold.^{67,68} Moderately high osmolarity is required to elicit cellular inflammatory responses *in vitro* in order to evaluate the beneficial effects of CL packaging solutions and their respective ingredients. Also, packaging solutions were tested at 25% of their normal concentrations; it is not known if the solutions and their ingredients at full concentration would elicit the same cellular responses, *in vitro* or clinically.

Conclusion

Including osmoprotectants in CL packaging solution may protect corneal epithelium from inflammation under hyperosmotic conditions that occur in CL-related DEDs and that, in some wearers, worsen at the end of the day. Other components such as surfactants and potassium electrolyte might also contribute to preserving ocular surface homeostasis, and their potential roles remain to be determined. Overall, this study underscores the significance of exploring novel strategies, particularly osmoprotective agents, to enhance the safety and comfort of CL wearers and maintain ocular health.

Data Sharing Statement

All relevant data are within the manuscript. Clarification requests around the manuscript and its data can be made to the corresponding author.

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Author Contributions

All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure

All authors are employees of Bausch & Lomb Incorporated. Ms Catherine Scheuer reports a patent US 17/398,556 pending to Bausch+Lomb, a patent Taiwan 110129328 pending to Bausch+Lomb, a patent PTC/EP2021/072140 pending to Bausch+Lomb. Dr William Reindel has a patent WO2022034010A1 pending to Bausch+Lomb. The authors report no other conflicts of interest in this work.

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