Steady-State Diffusion of Water through Soft-Contact-Lens Materials

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Abstract

Water transport through soft contact lenses (SCL) is important for acceptable performance on the human eye. Chemical-potential gradient-driven diffusion rates of water through soft-contact-lens materials are measured with an evaporation-cell technique. Water is evaporated from the bottom surface of a lens membrane by impinging air at controlled flow rate and humidity. The resulting weight loss of a water reservoir covering the top surface of the contact-lens material is recorded as a function of time.

New results are reported for a conventional hydrogel material (SofLensTM One Day, hilafilcon A, water content at saturation $w_{10} = 70$ weight %) and a silicone hydrogel material (PureVisionTM, balafilcon A, $w_{10} = 36$ %), with and without surface oxygen plasma treatment. Also, previously reported data for a conventional HEMA-SCL ($w_{10} = 38$ %) hydrogel are reexamined and compared with those for SofLensTM One Day and PureVisionTM hydrogels. Measured steady-state water fluxes are largest for SofLensTM One Day, followed by PureVisionTM and HEMA. In some cases, the measured steady-state water fluxes increase with rising relative air humidity. This increase, due to an apparent mass-transfer resistance at the surface (trapping skinning), is associated with formation of a glassy skin at the air/membrane interface when the relative humidity is below 55-75%.

Steady-state water-fluxes are interpreted through an extended Maxwell-Stefan diffusion model for a mixture of species starkly different in size. Thermodynamic non-ideality is considered through Flory-Rehner polymer-solution theory. Shrinking/swelling is self-consistently modeled by conservation of the total polymer mass. Fitted Maxwell-Stefan diffusivities increase significantly with water concentration in the contact lens.

Keywords: evaporation-cell method, water diffusion-coefficient, soft contact lens, extended Maxwell-Stefan model, trapping skinning, glass transition.

Introduction

The performance of soft contact lenses (SCL) on the human eye depends strongly on the water content and water-transport properties of the contact lens [1]. There is much evidence to believe that *in-vivo* contact-lens dehydration is primarily caused by an evaporative-dehydration process [2-7]. Evaporation at the anterior surface of a partially dewetted SCL draws water from the post-lens tear film (PoLTF) and through the lens into the environment (air). If this outward water transport rate is sufficiently large, the PoLTF may deplete, causing corneal desiccation [8, 9], reduced on-eye movement [10, 11] and, in extreme cases, lens adherence to the cornea surface [7, 12]. Water loss may also lead to discomfort and to dryness symptoms [13], and may reduce oxygen transmissibility [14], as well as fitting and corrective power of the lens [7]. Transport of water through a SCL can also be induced by an osmotic gradient between pre-lens and post-lens tear films [15], or by mechanical lid stress [16]. Understanding the importance of these phenomena requires quantitative knowledge of the rate of water transport through a thin polymer membrane.

Previous work on water diffusion in soft contact lenses has focused primarily on measuring self-diffusion coefficients [17-22], which characterize water transport in the absence of a chemical potential gradient. A few studies report measurements of kinetic sorption/desorption [23-27] of water or steady-state water fluxes [8, 28] driven by a water-activity gradient through a SCL.

In this work, we report steady-state diffusion rates of water through soft-contact-lens materials under a water chemical-potential gradient. To measure water flux precisely, we have greatly improved a recently developed evaporation-cell method [28]. In this experimental method, a water activity-gradient is established by water evaporation from the bottom surface of a SCL thin membrane whose upper surface is maintained in contact with pure water. In a recent report [29], we have shown that our method gives reliable diffusivities by comparing evaporation-cell data with results from other techniques for a HEMA SCL. Here, we study a conventional hydrogel (SofLensTM One Day) and a silicon hydrogel (PureVisionTM) with (prefix ST-) and without (prefix NT-) surface plasma treatment. For comparison, we also reconsider the previous results obtained for a HEMA SCL [29] .

Two distinct frameworks are commonly used to describe solute transport through membranes: the pore-flow model and the solution-diffusion model [30]. In the pore-flow framework, the membrane polymer is viewed as a porous solid matrix through which the solute hydraulically flows, whereas in the solution-diffusion model the polymer/solute is viewed as a molecular mixture, and solute transport is via diffusion. According to Wijmans and Baker [30], in a solution-diffusion membrane, the free-volume elements are a consequence of statistical concentration fluctuations in the membrane. They appear and disappear in about the same time scale as that for the motion of the solute penetrant. Conversely, in a porous-medium membrane, the free volume elements (pores) are relatively stable and do not fluctuate in position and size on the time scale of penetrant transport. In contrast to others who adopt a porous-medium model to describe water transport through SCL materials [2, 31], we argue that the solution-diffusion model

provides a more consistent representation of the physics of water transport in SCL hydrogel materials. Further, the very small pressure-induced water permeabilities of HEMA [32, 33] and PureVisionTM SCL [32] support a diffusive rather than a pressure-driven viscous mechanism for water transport through these materials.

Within the solution-diffusion framework, Fick's law is generally adopted to describe diffusion in binary systems because of its simplicity. However, Fick's law does not explicitly account for thermodynamic nonideality and also cannot be readily applied to multicomponent systems. We view a water/hydrogel SCL as a highly nonideal molecular mixture. Moreover, we are interested in extending our present study to include simultaneous, multicomponent transport of other small molecules (e.g., salt or oxygen) through a contact lens. A self-consistent description of multicomponent transport for a simple fluid mixture is given by the generalized Maxwell-Stefan (GMS) framework that accounts both for thermodynamic nonideality and for diffusion coupling in multicomponent mixtures [34]. However, even in the binary case, difficulties are encountered when GMS is applied to transport through a cross-linked polymeric membrane because the unknown molecular weight (or, equivalently, a molar concentration) of the membrane must be specified [28]. This problem arises because GMS does not represent correctly the frictional resistance between molecules that differ greatly in size. Recently, we have proposed an extended Maxwell-Stefan model (EMS) to describe multicomponent transport for molecules of starkly different size [35]. When EMS is applied to solute transport in a membrane, there is no need to specify the unknown membrane molecular weight. In this work, we adopt the EMS model to analyze our experimental water fluxes through SCL materials. Swelling/shrinkage of the

membrane is included self-consistently by imposing conservation of the polymer mass in the membrane. For completeness, we report also Fickian diffusion coefficients for water in our SCL materials.

Experimental

Materials

Distilled/deionized water is obtained from a MilliQTM purification system; water resistivity is greater than 18 MΩ cm. Commercial SCL materials were kindly provided by Bausch & Lomb as flat, rectangular sheets (7 x 9.5 cm) synthesized between two glass plates. Two soft-contact-lens (SCL) materials were used in this study: a conventional hydrogel, SofLensTM One Day (hilafilcon A), and a last-generation silicone-hydrogel, PureVisionTM (balafilcon A). One batch of PureVisionTM sheets was subjected to surface oxygen plasma treatment, similar to the commercial product [36-38], whereas a second batch was not.

Table 1 gives properties of SCL membranes, including a previously studied HEMA (2-hydroxyethyl methacrylate) SCL [29]. Water content at saturation, w_{10} , reported by Bausch & Lomb, refers to weight percent of water in a wet contact lens that has been completely water saturated in a sterile isotonic solution at room temperature. The conventional hydrogel SofLensTM One Day has a water content at saturation of 70%, much larger than that of HEMA at 38%. The saturation water content of PureVisionTM is 36%, similar to that of HEMA.

Wet thickness, L_0 , of the SCL membranes was measured using a micrometer (Mitutoyo MDC 0-1" PF, accuracy $\pm 1 \mu\text{m}$) at different locations; the average value reported in Table 1 (with the standard deviation) is used for subsequent calculations.

From the small standard deviation, we conclude that the lenses have an essentially uniform thickness. The glass-transition temperature of each dry SCL material was measured by modulated differential scanning calorimetry (Modulated DSC 2920, TA Instruments) as described elsewhere [39]. Table 1 also reports the dry mass density, ρ_p , for each contact lens, as estimated from the water content and from the specific gravity of the saturated SCL (reported by Bausch & Lomb) by assuming ideal volume of mixing. Flory interaction parameters, χ , are calculated as discussed later.

Apparatus

The right side of Fig. 1 shows a schematic of the evaporation cell used to measure diffusivities of water through soft-contact-lens membranes [28]. The SCL membrane supports a small reservoir of water (10 g) on the top surface while the bottom surface is exposed to impinging-air jet flow at fixed relative humidity (RH) and flow rate. The so-established chemical-potential gradient for water between the two membrane surfaces (i.e. the water content difference) induces an evaporative water flux across the lens that is measured gravimetrically. The evaporation cell, which sits atop an electronic balance (Sartorius, Model CP 343 S) is similar to that of Hoch et al. [28]. A detailed description of the modifications may be found in the thesis of Fornasiero [40].

A brass stand pipe is located below the membrane surface to provide an air supply of known humidity. The ratio between the pipe diameter and the SCL-exposed diameter (2.29 cm) is chosen to provide uniformly accessible mass transfer across the membrane surface [41]. Turbulent air flow is desired because measured water evaporation fluxes at finite air-flow Reynolds numbers, Re , must be extrapolated to infinite Re (see next section). Large air-flowrates are required to reach the turbulent regime in an empty pipe.

Further, when directly blown against the SCL membrane, such high flow rates cause damage and a large, lift-induced scattering of weight measurements. To overcome these difficulties, a 12-element Kenics® static mixer is inserted into the brass air-supply stand pipe. The Kenics® static mixer produces turbulent flow at Re as low as 500 (supplier specifications and [42]). As illustrated on the left of Fig. 1, dry, compressed air (relative humidity $RH = 5\text{-}7\%$) from the laboratory line is first filtered to remove dust and oil droplets. The air supply is then split into two lines. One line feeds the dry air to a humidification column ($H = 100$ cm; OD = 10 cm) where it is water saturated by countercurrent contact with a water stream in a packed column (Berl saddles, nominal size $\frac{1}{2}$ in, 40-cm height). The other line bypasses the humidification column and is recombined with the wet-air line in a ratio required to reach the desired RH .

A mass-flow sensor (Sensirion, Model ASF 1400) and a humidity/temperature sensor (Sensirion, Model SH 11) allow real-time monitoring of the total air flow rate, temperature, and relative humidity. Together with the electronic balance, they are interfaced to a computer through a dedicated LabView® program for automatic data collection.

Before measuring steady-state water fluxes through the membranes, we verified that no other water loss occurs in the evaporation cell. Instead of a SCL membrane, we replaced the lens material with an impermeable aluminum foil in the evaporation cell and measured the weight loss as in a typical water-flux experiment. The measured water flux was less than 1% of the lowest observed flux with a SCL material in place. We conclude that our evaporation cell is tightly sealed.

Method

SCL membranes are removed from the original ReNu Multiplus® (Bausch & Lomb) disinfecting solution and rinsed thoroughly with distilled/deionized water. Then, the membranes are soaked in distilled/deionized water for at least three days. Water is renewed every 24 h to ensure complete removal of the disinfecting solution. Upon completion of the cleaning procedure, the membranes are immersed in fresh distilled/deionized water and stored in a refrigerator to prevent bacterial growth.

After choosing the specific type of lens to be studied, a sheet of the contact-lens material is sandwiched between two supporting Delrin® rings, and 10 g of distilled/deionized water are placed on the top surface of the sandwiched membrane. Two layers of SaranTM wrap are then fastened over the top ring, such that the water-containing chamber is sealed and undesired evaporation from the top surface is minimized. The evaporation cell is then placed on the balance. Once the desired air flow rate and relative humidity are attained, measurements of mass, flow rate, relative humidity, temperature and time are recorded every 15 s for at least 4 h with a LabView® program. Water mass-loss measurements are averages over this 15-s time interval.

Steady-state conditions are reached when the water flux through the lens is constant, as verified by comparing the slope of mass-versus-time data at various intervals of time. The last-hour slope is used to calculate the steady-state water mass flux, n_l . Typically, n_l differs by less than 1% from the last two-hours slope.

For each lens material, experimental data are collected for seven different air-flow rates in the range 6-50 L/min and for three different relative humidities at 24.0 ± 0.5 °C. Chosen air-flow rates are 6, 8, 10, 14, 20, 30 and 50 L/min, while relative humidities are

20, 50, and 80%. Air flow and relative humidity are maintained constant within $\pm 0.5\%$ and 1% of the set points, respectively. Each experiment is repeated at least twice to assure data reproducibility. Repeat experiments yield water fluxes that typically agree to within 2 – 4%. Additional details on the experimental apparatus and measurement procedures are available elsewhere [40].

In general, a mass-transfer resistance for water is present at the membrane/air interface. This interfacial resistance is in series with the diffusive resistance of the membrane and must be accounted for. For impinging-jet flow, the mass-transfer coefficient scales with the square root of the air-flow rate [41]. Therefore, a plot of $1/n_l$ as a function of $Re^{-1/2}$ should be linear. The intercept on the ordinate of this plot gives the inverse water flux for an infinite air-flow Re , i.e. in the absence of mass transfer resistance at the membrane/air interface [28]. This extrapolated water flux, n_l^∞ , is used here to determine the water diffusivity in the SCL material. Because of excellent reproducibility of measured fluxes, the experimental error for extrapolated water fluxes is small, as demonstrated in column 3 of Table 2.

Extended Maxwell-Stefan Framework

To interpret the water-evaporative flux results, we use the recently proposed extended Maxwell-Stefan (EMS) model [35]. Because the water head in the evaporation-cell reservoir is very small, we assume that our system is isobaric. Further, we assume isothermal diffusion with no volume change upon mixing. Water transport occurs essentially only in the direction orthogonal to the membrane surface. Under these conditions, EMS reduces to

$$c_1 \frac{d\mu_1}{dx} = RT \frac{\phi_1 \phi_p}{v D_{lp}} (v_p - v_1) \quad (1)$$

where subscripts 1 and p denote water and polymer, respectively; μ_1 is the local chemical potential of water in the membrane; x is the distance from the upper surface of the membrane; c_1 is molar concentration of water in the water/polymer mixture; ϕ_i is volume fraction of i ; v_i is species i velocity; v is the polymer segment molar volume; and D_{lp} is the binary EMS diffusion coefficient for water in the membrane.

Since EMS regards a membrane/solvent system as a nonideal polymer/solvent mixture, the chemical potential of water in the membrane is calculated from the polymer-gel/solution theory of Flory and Rehner [28, 43-45]:

$$\mu_1 = \mu_1^0 + RT \left[\ln \phi_1 + \phi_p + \chi \phi_p^2 + \frac{v_1^0 G}{RT} \left(\phi_p^{1/3} - \frac{1}{2} \phi_p \right) \right] \quad (2)$$

where the superscript 0 denotes a pure species, χ is the Flory solvent-polymer interaction parameter, v_1^0 is the molar volume of pure water, and G is the membrane bulk modulus. The first three terms within the square brackets of Eq (2) correspond to Flory-Huggins [46] theory for a polymer with an infinite molecular weight, whereas the last term provides the contribution of membrane elasticity. Here, the Flory interaction parameter χ is assumed constant and is obtained from the measured membrane water content when equilibrated in a sterile isotonic solution [28], as reported in Table 1 (the difference by water contents in distilled/deionized water and isotonic solution is minimal for the nonionic lenses studied). Precise values of G for all SCL's are not available. Fortunately, calculations are not sensitive to the value of G . In our calculations, we use $G=1.0$ MPa for all SCL materials, close to that reported for PureVisionTM [36, 38, 47].

Appendix I gives calculation details for fitting EMS diffusivities $\bar{D}_{lp}$ while accounting for shrinking/swelling of the SCL membranes, and lists the boundary conditions for integration of Eq (1). For completeness, Appendix I also demonstrates the calculation of Fickian diffusion coefficients, D_{lp} , for our water/SCL membrane systems.

Results

In Figs. 2-5, we plot the inverse of the measured steady-state mass fluxes of water $1/n_1$ as a function of air-flow $Re^{-1/2}$ for our contact-lens materials. Here, the reference diameter for Re is that of the empty air-supply line. Best linear fits are also drawn for comparison. Clearly, the experimental data are well represented by fitting straight lines for all SCL membranes and for all relative humidities, indicating that our experiment obeys the impinging-jet flow conditions. Therefore, we can confidently extrapolate our measured water fluxes to infinite air-flow rate to obtain the steady-state diffusive mass fluxes n_1^∞ through the SCL membranes in the absence of any external mass-transfer resistance. In these plots, the nonzero slope of the straight lines shows that the external mass transfer resistance cannot be neglected. This slope increases with rising air relative humidity in agreement with theoretical predictions [48].

Figs. 4 and 5 show measured steady-state fluxes for non-treated (NT) and surface-treated (ST) PureVisionTM lenses, respectively. $RH = 50$ and 80% results for ST-PureVisionTM are almost indistinguishable from those for NT-PureVisionTM. Because the two materials present identical water fluxes at these RH s, we did not repeat the experiments for ST-PureVisionTM to verify data reproducibility after investigating NT-PureVisionTM. At $RH = 20\%$, water fluxes for ST-PureVisionTM are 10% lower than those for NT-PureVisionTM at the same air-flow rate. Extrapolated water fluxes for ST-

PureVision™ differ less than 10% from those for NT-PureVision™ at the same RH (see Table 2). The oxygen-plasma surface treatment creates small, hydrophilic silicate islands on the surface of ST-PureVision™ contact lens [37, 38, 49]. These islands are separated by the nascent, hydrophobic silicone-hydrogel material (balafilcon-A), and are elevated by about 10-50 nm [36-38]. Because only a partial area of the lens surface is affected by plasma treatment and only for a shallow thickness, material bulk properties such as diffusivity are expected to be, at most, marginally affected by surface treatment, consistent with our experimental findings. Because ST-PureVision™ and NT-PureVision™ contact-lens materials have essentially the same equilibrium and transport properties for water, we refer to these materials simply as PureVision™ in the remaining discussion.

Table 2 lists the extrapolated fluxes n_1^∞ . In this table columns 2 and 3 reported measured quantities, whereas column 4-9 list calculated values from EMS or Fick's laws, as described in Appendix I. Comparison of n_1^∞ at the same relative humidity for HEMA and SofLens™ One Day materials shows that the water flux is significantly larger, by an order of magnitude, for the high-water-content SofLens™ One Day lens. Similarly, Martin [8] reported a positive correlation between steady-state evaporative loss and water content for conventional soft-contact-lens materials. This result is also consistent with clinical and experimental observations that conventional hydrogel contact lenses with higher water content at saturation tend to dehydrate on eye more quickly and in a greater amount [1, 5, 50, 51]. Silicone-hydrogel PureVision™ has a water content at saturation that is slightly lower than that for HEMA. However, the measured water fluxes are twice as large. To our best knowledge, there are no previous published data for water fluxes

through silicone hydrogel materials driven by a chemical-potential gradient. Some data have been reported for tritium-labeled water diffusion rates, which show a somewhat (10%) higher water transport for PureVision™ compared to that in HEMA [16, 36].

Discussion

The calculated steady-state membrane thickness for SofLens™ One Day is nearly half of those for HEMA and PureVision™ lens materials (see Tables 1 and 2), whereas the steady-state water fluxes for SofLens™ One Day are 10 and 5 times larger than those for HEMA and PureVision™, respectively. Thus, the large variation in water fluxes must be primarily related to the physico-chemical properties of the lens membranes. SofLens™ One Day is a copolymer of HEMA and N-vinyl-pyrrolidone (VP). The latter monomer confers augmented hydrophilicity to the SCL material, increasing saturation water content from 38 (HEMA homopolymer) to 70 %. Moreover, polyvinylpyrrolidone exhibits water diffusivities greater than those in HEMA [52-55]. Therefore, VP enhances water transport in SofLens™ One Day by increasing both water uptake and diffusivity in the SCL material. PureVision™ has a more complex chemical composition with the exact formulation proprietary. However, literature indicates that PureVision™ is a copolymer of tris-(trimethylsiloxy)-silyl-propylvinyl carbamate (TRIS-VC), N-vinyl-pyrrolidone [56], a vinyl carbonate functional polydimethylsiloxane (PDMS) macromer, and a vinyl carbamate derivative of alanine [47]. Although water uptake for PureVision™ is essentially the same as that for HEMA, water diffusivity is most likely augmented by the presence of both silicone [57, 58] and VP groups.

To interpret our experimental fluxes with EMS or Fick's law, we first take the EMS ($\bar{D}_{lp}$) or Fickian (D_{lp}) diffusivities as concentration independent, and we fit $\bar{D}_{lp}$ or

D_{lp} separately for each RH condition, as explained in Appendix I. Table 2 lists the resulting diffusivities, average steady-state water-content ($\langle w_1 \rangle$) and fractional thickness variation calculated both with the EMS model and with Fick's law. At fixed RH , the largest diffusivity is obtained for SofLensTM One Day followed by PureVisionTM and HEMA. This order is consistent with the previous analysis based on membrane chemistry. Clearly, however, the values of D_{lp} and D_{lp} both increase significantly with average water content for all SCL membranes. The starting assumption of concentration-independent diffusivities is, therefore, not satisfied.

Concentration dependent diffusivities

In an attempt to account for the strong water-content dependence of the diffusivities, we assumed an exponential variation with water volume fraction, $D_{lp} = a \exp(b\phi_1)$, where a and b are constants; we fit the measured water fluxes to obtain parameters a and b according to the procedure outlined in Appendix I. Unfortunately, fitting results were poor. To resolve this discrepancy, we first questioned the assumption of isothermal diffusion as outlined in Appendix II. Our calculations show that evaporative cooling at the membrane/air interface has little impact on the measured water fluxes (see Table 3). Hence, our systems can be effectively considered isothermal.

To explore further the reasons for our inability to describe the RH dependence of our extrapolated water fluxes, n_1^∞ , we examined the trend of measured water fluxes as a function of RH . Because a higher air RH corresponds to a lower driving force for water flux, one expects lower fluxes at higher RH . For HEMA n_1^∞ indeed decreases (or remains unchanged within experimental uncertainty) with increasing RH . However, the opposite

trend and a minimum are observed with SofLens™ One Day and PureVision™ materials, respectively. Crank [59, 60] reports similar increases in steady-state diffusive transport for increasing gas concentration of an evaporating solvent from polymer membranes. Such increase of solvent flux with a smaller driving force is commonly observed in drying technology and is termed “trapping skinning” [61].

In EMS, the water flux at steady state can be written in the form

$$n_1^\infty = \frac{1}{L} \int_{\phi_L}^{\phi_0} \frac{D_{lp}}{RT} \frac{\phi_1}{\phi_p \hat{v}_1^0} \frac{d\mu_1}{d\phi_1} d\phi_1. \quad (3)$$

where $\hat{v}_1^0$ is the specific volume of pure solvent. The derivative of the chemical potential with composition is positive for a stable mixture. For any non-negative concentration-dependent D_{lp} , the integrand in Eq (3) is non-negative. Since the upper limit of integration in Eq (3) is fixed and the lower limit, ϕ_L , decreases by lowering the air RH , EMS predicts a steady-state flux that always increases when the air RH decreases. Therefore, our model cannot predict trapping skinning for any D_{lp} that is a non-negative function of concentration. Arguments presented in Appendix III show that accounting for shrinking/swelling does not relieve the discrepancy in the EMS model. Note, also, that the same result is encountered in Fickian diffusion models [60, 62].

Although trapping skinning was first observed several decades ago, there is no clear physical understanding of this phenomenon. It has been hypothesized [61, 63-65] that trapping skinning may occur when the surface of the polymeric material exposed to the atmosphere undergoes a rubber-to-glass transition during drying. A recent study using atomic force microscopy [66, 67] showed that such transition occurs in some HEMA-based SCLs when environmental relative humidity drops below 50-60%. We have also

investigated the possibility of glassy-skin formation for our SCL materials by measuring glass transition temperatures as a function of water content with modulated differential scanning calorimetry [39]. We find that, for the formation of a glassy skin, a local water content is required of less than 12, 15, and 7 weight % for HEMA, SofLensTM one day and PureVisionTM materials, respectively. The corresponding air RH ($\sim$ water activity) can be estimated from equilibrium water-activity versus water-content data obtained with a gravimetric sorption technique at 35 °C [68]. Our results indicate that the rubber-to-glass transition occurs at estimated RH of 74%, 64% and 57% for HEMA, SofLensTM One Day and PureVisionTM, respectively. Therefore, it is likely that all three SCL materials exhibit a glassy skin at the membrane/air interface when the experimental RH is equal to 20% and 50% in Table 2, which may explain the existence of trapping skinning. For HEMA, the appearance of a glassy skin apparently compensates exactly the effect of increased driving force for water flux when the RH is reduced from 50% to 20%. For HEMA, no substantial variation with RH is observed in the measured water fluxes. For SofLensTM One Day material, however, trapping skinning overpower the chemical potential gradient driving water transport leading to a decreasing water flux with decreasing RH . Therefore, consistent with on our bulk T_g measurements and with the surface T_g experiments of Opdahl et al. [66] and Koffas et al. [67], trapping skinning is the most likely explanation for the unsuccessful fitting of our experimental fluxes with a diffusivity increasing with membrane water-concentration. A concentration-dependent diffusivity cannot explain the water flux data for the studied contact lens materials.

Only a few models [61, 63-65] have been proposed to explain trapping skinning, mainly for the drying of polymeric coatings. One model invokes a viscoelastic diffusion

process for a polymer/solvent mixture near its glass-to-rubber transition temperature [61, 63, 64]; another model assigns its capability of predicting trapping skinning to incorporation of a non-ideal volume change in the glassy region of the polymer/solvent mixture [65]. Unfortunately, we cannot directly apply these models to our systems because many of the required parameters are not available or cannot be easily obtained for our SCL materials. However, we believe that it is unlikely that the incorporation of a non-ideal volume change can explain our results. Indeed, Appendix III shows that the steady-state water flux must increase upon reduction of external RH , except in the unlikely instance that the partial specific volume for the SCL polymer is negative. We conclude that accounting for non-ideal volume of mixing in a glassy skin cannot represent trapping skinning in our systems.

Other models may be proposed to represent trapping skinning. For example, we may consider a surface resistance on the SCL - surface exposed to air which decreases upon raising the external relative humidity. However, the number of parameters required to fit our extrapolated water fluxes is too large to obtain unambiguous values. Because EMS (and Fick's law) do not account for a rubber-to-glass transition and cannot represent trapping skinning, we can only report the concentration-independent diffusivities listed in Table 2; these reported diffusivities must be considered as effective diffusivities when the RH is below the critical RH for the glass-to-rubber transition.

Soft contact lens wear

At the beginning of a blink cycle, a soft contact lens on-eye is normally bathed in the tear film. The pre-lens tear film (PrLTF) on a soft contact lens is generally unstable and ruptures several seconds after lid opening [69-71]. Some regions of the SCL surface

are then exposed directly to external air [72]. Upon PrLTF rupture, an evaporative-dehydration process commences that draws water through the contact lens and out of the post-lens tear film toward the environment [2-7]. A subsequent blink deposits a new PrLTF on the SCL surface, and the contact lens partially rehydrates. This process repeats cyclically at each blink. After some time, the soft contact lens reaches a periodic steady state [2, 48]. In this condition, the SCL has constant water content when averaged over a blink cycle. However, there remains a non-zero water flux from the PoLTF through the contact lens toward the external air. The amount and the rate of dehydration depend on a large number of factors including water diffusivity in the SCL material, SCL thickness, PrLTF stability, SCL surface area exposed to air upon PrLTF rupture, blink rate, external *RH*, and wind velocity. Because water diffusivity and water content are higher for SofLensTM One Day, we expect SofLensTM One Day to be more prone to *in-vivo* dehydration and PoLTF collapse than HEMA and PureVisionTM contact lenses. From this point of view, high water transport rates through SCLs are undesirable. Consequently, formation of a glassy skin and a water diffusivity that increases with water concentration may be beneficial for a soft contact lens worn on the eye because they limit lens dehydration in low humidity environments. Conversely, physical interaction of the eyelid with a glassy skin may lead to discomfort [73].

Conclusions

We have investigated water transport through typical soft-contact-lens (SCL) materials with an evaporation-cell method. Measured steady-state water fluxes are largest for SofLensTM One Day ($\sim 50\text{-}70 \times 10^{-6} \text{ g cm}^{-2} \text{ s}^{-1}$), an order of magnitude greater than those for HEMA membranes ($\sim 4\text{-}5 \times 10^{-6} \text{ g cm}^{-2} \text{ s}^{-1}$). For PureVisionTM, the surface

oxygen-plasma treatment does not affect water transport through the soft contact lens. For SofLensTM One Day and (in some cases) for PureVisionTM materials, water fluxes increase with rising environmental relative humidity. This phenomenon is associated with formation of a glassy skin at the membrane/air interface. Independent measurements of the glass-transition temperature for these SCL-polymers as a function of water content strongly suggest that our SCL materials develop a glassy skin when $RH = 20\%$ and $RH = 50\%$, whereas our lenses remain rubbery at $RH = 80\%$.

Steady-state water fluxes are interpreted with an extended Maxwell-Stefan (EMS) model for a homogeneous mixture of molecules of starkly different size. Fick's law diffusion coefficients are also reported. Fitted (effective) diffusivities increase with rising water concentration. EMS diffusivities are in the range $2.6\text{-}7.3$, $8.2\text{-}35$, and $5.0\text{-}17 \times 10^{-7} \text{ cm}^2 \text{ s}^{-1}$, respectively, for HEMA, SofLensTM One Day and PureVisionTM; corresponding Fickian diffusivities are in the range $1.7\text{-}3.4$, $5.3\text{-}16$ and $3.3\text{-}8.4 \times 10^{-7} \text{ cm}^2 \text{ s}^{-1}$. The faster water transport and higher water content of SofLensTM One Day suggest that this lens material is more prone to *in-vivo* dehydration and PoLTF depletion than HEMA and PureVisionTM.

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Appendix I. Fitting Diffusivities to Evaporation-Cell Water Fluxes: Mathematical Details

Eq (1) in the text gives the EMS diffusion equation for an isothermal, isobaric membrane/water system. At steady state, the membrane polymer species velocity is zero and the water velocity (or flux) is a constant. Because the chemical potential in Flory-Rehner theory is expressed as a function of volume fraction, it is convenient to rewrite Eq (1) as

$$\frac{d\mu_1}{d\ln\phi_1} \frac{d\phi_1}{dx} = -\frac{RT}{D_{lp}} \phi_p \hat{v}_1^0 n_1^\infty \quad (\text{I.1})$$

where n_1^∞ is the extrapolated mass water flux through the SCL material at steady-state. Eq (I.1) is a first-order nonlinear differential equation with an unknown parameter, the diffusivity D_{lp} . We want to obtain D_{lp} by fitting to our measured water fluxes, n_1^∞ , at steady state.

Unfortunately, integration of Eq (I.1) is not straightforward since the steady-state thickness L of the membrane is unknown because the membrane swells or shrinks in response to the water-content profile. To account self-consistently for swelling/shrinking of the membrane, we impose conservation of the total polymer mass [28, 74]. Thus, we define a variable η representing the mass of polymer per unit area between $x = 0$ and x ,

$$\eta(x) = \int_0^x \frac{\phi_p}{\hat{v}_p} dx \quad (\text{I.2})$$

where $\hat{v}_p$ is the partial specific volume for polymer. Polymer mass conservation requires that $\eta(L)$ is a constant $(= L_0 \phi_{p0} / \hat{v}_p)$. It is convenient to write Eq (I.2) in differential form and to solve it simultaneously with Eq (I.1) [74]. We define a dimensionless

distance $\xi = x/L$. After replacing x with ξ in Eq (I.2), we reduce the integral equation to a first-order differential equation or

$$\frac{d\eta}{d\xi} = L \frac{\phi_p}{\hat{v}_p} \quad (\text{I.3})$$

To now solve for the membrane thickness we note that L is a scalar independent of position so that

$$\frac{dL}{d\xi} = 0 \quad (\text{I.4})$$

In general, the EMS diffusivity is a function of composition. Since this dependence is not known, we first assume that D_{lp} constant and then, if needed, relax this constraint. We express the constancy of D_{lp} as

$$\frac{dD_{lp}}{d\xi} = 0 \quad (\text{I.5})$$

After rewriting Eq (I.1) in terms of the variable ξ , we solve numerically the set of coupled, nonlinear, first-order ordinary differential Eqs (I.1), and (I.3)-(I.5).

Boundary Conditions

Four boundary conditions are required. Since there is no mass-transfer resistance at the air/membrane interface at infinite air-flow rate, the volume fraction of water at the bottom surface of the membrane, ϕ_{1L} , is set by thermodynamic equilibrium with water vapor in air at the bulk relative humidity. Thus, the boundary condition at the membrane surface exposed to air is

$$\phi_1 = \phi_{1L} \quad \text{at } x = L \quad (\text{I.6})$$

where $\phi_{1,L}$ is calculated as explained earlier [28]. The other surface of the membrane is in equilibrium with pure water. Therefore, we have that

$$\phi_1 = \phi_{10} \quad \text{at } x = 0 \quad (\text{I.7})$$

where ϕ_{10} is calculated from the membrane water content at saturation [28]. The associated boundary conditions for Eqs (I.3) and (I.4) are, respectively

$$\eta(0) = 0 \quad (\text{I.8})$$

$$\eta(1) = L_0 \phi_{p0} / \hat{v}_p \quad (\text{I.9})$$

For each relative humidity, we solve simultaneously the set of coupled nonlinear differential equations using a finite-difference scheme with Newton iteration of the nonlinearities (subroutine BAND(J) [75]).

EMS Diffusivities

As shown in the Discussion section, the resulting D_{lp} values vary with relative humidity. In particular, D_{lp} increases significantly with the average water content in the lens. Therefore, we relax the hypothesis of constant D_{lp} and assume an exponential concentration dependence for D_{lp}

$$D_{lp} = a \exp(b\phi_1) \quad (\text{I.10})$$

where a and b are constants. We use a Levenberg-Marquardt method [76] to fit these constants to the measured steady-state water fluxes in Table 2.

Fickian Diffusivities

Because most researchers report Fickian diffusivities for small penetrants in polymeric membranes, we calculate here the diffusion coefficient of Fick's law for our

systems. Under the assumption of a concentration-independent D_{lp} , the Fickian water-concentration profile in the membrane is linear at steady state. Steady-state thickness L can be determined analytically by integration of Eq (I.2). Fickian diffusivities are then given by

$$D_{lp} = \frac{N_1^\infty L}{c_{i0} - c_{iL}} \quad (\text{I.11})$$

where c indicates a molar concentration and N_1^∞ is the extrapolated molar flux of water.

We do not attempt to fit concentration-dependent Fickian diffusivities.

Appendix II. Impact of Evaporative Cooling on Steady-State Water Fluxes

During an evaporation-cell experiment, cooling occurs on the bottom SCL surface because of water evaporation. We want to assess the importance of evaporative cooling in our evaporation-cell experiments by comparing calculated water steady-state fluxes for isothermal and nonisothermal diffusion. The non-isothermal case requires solving simultaneously heat transfer (HT) and mass transfer (MT) equations in a membrane/air system. At steady state, the equations and boundary conditions for HT and MT in the membrane are, respectively [77]:

$$\frac{dn_1}{dx} = 0 \quad w_1(x=0) = w_{10}, \quad w_1(x=L) = w_{1L}; \quad (\text{II.1})$$

$$\frac{de}{dx} = 0 \quad T(x=0) = T_0, \quad T(x=L) = T_L, \quad (\text{II.2})$$

where

$$n_1 = \frac{\rho D_{1p}}{1 - w_1} \frac{dw_1}{dx}, \quad (\text{II.3})$$

and

$$e = -k_T \frac{dT}{dx} + n_1 c_p (T - T_0). \quad (\text{II.4})$$

In Eqs (II.1)-(II.4), T indicates temperature, and w_1 is the water weight fraction; subscripts 0 and L indicate membrane/water and membrane/air interfaces, respectively; k_T is the thermal conductivity for the water/membrane mixture; c_p is the specific heat capacity for liquid water; ρ is the wet-SCL mass density; n_1 is the water steady mass flux, and e is a combined energy flux defined by equations (19.3-4) and (19.3-6) in reference [77]. To derive Eq (II.3), we used Fick's law to express the water diffusive

flux. In Eq (II.4), the first term on the right represents heat conduction, while the second term corresponds to the enthalpy flux associated with mass transport [77].

The water weight fraction, w_{1L} , and the temperature, T_L , at the membrane/air interface are unknown. We obtain these quantities by setting the fluxes n_1 and e in the membrane equal to the corresponding fluxes on the airside of the membrane/air interface:

$$n_1 = k_1 \rho_{air} \left(\frac{w_{1L}^{air} - w_1^{air}}{1 - w_{1L}^{air}} \right) \quad (II.5)$$

$$e = h_T (T_L - T_0) + n_1 c_{pv} (T_L - T_0) + \lambda_1 n_1 \quad (II.6)$$

where superscript air indicates air side; k_1 and h_T are mass transfer and heat transfer coefficients at the SCL membrane/air interface, respectively; ρ_{air} is air mass density; c_{pv} is specific heat capacity for water vapor, and λ_1 is the enthalpy of vaporization for water. In Eq (II.6), the first term represents the heat flux in absence of simultaneous mass transfer, while the second and third terms give the heat flux associated with water-vapor transport and water vaporization, respectively. We assume that the temperature in bulk air is equal to that of the water reservoir ($T_0 = 25$ °C), and that w_{1L}^{air} and w_{1L} are related by local equilibrium.

To obtain T_L and w_{1L} , we use an iterative procedure. For the isothermal case, only Eq (II.5) is required, and every quantity appearing there is evaluated at temperature T_0 . Details on parameter estimates and calculation procedure are given elsewhere [40].

Table 3 summarizes the results of calculations for coupled HT and MT equations. Table 3 shows that cooling at the membrane/air interface does not exceed 3-4 °C, and that it is larger for smaller RH . Water flux is reduced by evaporative cooling. The effect is the

greatest for SofLensTM One Day at the lowest RH , because the evaporative fluxes are the largest for this SCL material. Despite conservative assumptions for D_{lp} and k_T , water fluxes for nonisothermal diffusion differ little from those obtained in the isothermal case. These results indicate that evaporative cooling at the membrane/air interface is not important for interpreting our water flux measurements.

Appendix III. Predicted Dependence of Steady-State Water-Flux on External Relative Humidity for Fickian or Maxwell-Stefan Diffusion Models.

According to a recently proposed drying model [65], trapping skinning in unsteady-state drying of polymer films can be explained by nonideal volumetric mixing behavior exhibited by a glassy polymer-solvent mixture. We show here that, under the assumption that the solvent diffusivity in a binary solvent/polymer system is a non-negative function of concentration only, accounting for nonideal volumetric behavior is insufficient to capture trapping skinning in an isothermal permeation experiment at steady-state conditions.

As example, consider a membrane one surface of which is in equilibrium with pure water (located at $x = 0$) and the other is in equilibrium with air at a $RH < 100\%$ (located at $x = L$), respectively. We investigate how the steady-state flux changes by varying the RH . The mass flux of solvent 1 is

$$n_1 = \rho_1 v_1 = \rho_1 v^\square + j_1^\square \quad (\text{III.1})$$

where $v^\square$ is the volume average velocity, $v^\square = \rho_1 \hat{v}_1 v_1 + \rho_2 \hat{v}_2 v_2$, $j_1^\square$ is the diffusive flux with respect to the volume average velocity, ρ_i is a mass concentration and $\hat{v}_i$ is a specific partial volume for species i . The diffusive flux $j_1^\square$ can be expressed as a product of a non-negative diffusivity $D(\rho_1)$, a function of composition only, and a gradient of composition:

$$j_1^\square = -D(\rho_1) \frac{d\rho_1}{dx} \quad (\text{III.2})$$

Because $D(\rho_1)$ can be any non-negative function of composition, Eq (III.2) includes Fick's law, EMS or the diffusion models used in reference [65]. At steady state, the polymer velocity is zero or

$$n_1 = j_1^\square / (1 - \rho_1 \hat{v}_1) = - \frac{D(\rho_1)}{(1 - \rho_1 \hat{v}_1)} \frac{d\rho_1}{dx}, \quad (\text{III.3})$$

and the water flux, n_1 , is constant. After multiplying both sides of Eq (III.3) by $\rho_2 dx$, integration of Eq (III.3) results in

$$n_1 = \frac{1}{M} \int_{\rho_{1L}}^{\rho_{10}} \frac{D(\rho_1)}{\hat{v}_2} d\rho_1 \quad (\text{III.4})$$

where ρ_{10} and ρ_{1L} are the solvent mass concentrations at the membrane surfaces at $x = 0$ and $x = L$, $(1 - \rho_1 \hat{v}_1) = \rho_2 \hat{v}_2$, and M is the total polymer mass per unit area given by

$$M = \int_0^L \rho_2 dx. \quad (\text{III.5})$$

In Eq (III.4), ρ_{10} is the solvent mass concentration in equilibrium with a pure solvent phase and ρ_{1L} is the solvent mass concentration in equilibrium with air at a given RH . Since the upper limit of integration is fixed and the lower limit decreases by lowering air RH , the range of integration rises when air RH falls. For any non-negative concentration-dependent $D(\rho_1)$, n_1 must increase by widening the range of integration, except when the partial specific volume of the polymer is negative. A negative $\hat{v}_2$ is unlikely. Therefore, except in the rare event of a negative $\hat{v}_2$, taking into account the non-ideal volumetric behavior in a glassy polymer-solvent mixture is not sufficient to explain a decreasing steady-state solvent flux when air RH falls.

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Figure Captions

Fig. 1 Schematic of evaporation-cell apparatus (right) along the humidified air supply (left). The drawing is not to scale.

Fig. 2 Measured steady-state water fluxes as a function of air-flow Reynolds number for HEMA at three relative humidities. Open circles correspond to experimental data at $RH = 80\%$, filled squares at $RH = 50\%$, and open triangles at $RH = 20\%$. Fitted straight lines are also shown. The dotted line is for $RH = 80\%$, the continuous line for $RH = 50\%$, and the dashed line for $RH = 20\%$.

Fig. 3 Steady-state water fluxes as a function of air-flow Reynolds number for SofLensTM One Day at three relative humidities. Symbols and lines have the same meaning as in Fig. 2.

Fig. 4 Steady-state water-fluxes as a function of air-flow Reynolds number for NT-PureVisionTM at three relative humidities. Symbols and lines have the same meaning as in Fig. 2.

Fig. 5 Steady-state water fluxes as a function of air-flow Reynolds number for ST-PureVisionTM at three relative humidities. Symbols and lines have the same meaning as in Fig. 2.

Table 1 Soft-contact-lens (SCL) properties

SCL	Material	ρ_p (g cm ⁻³)	L_0 (μ m)	w_{10} (%)	χ	T_g (°C)
HEMA	HEMA	1.274 ^b	195 (± 3)	38	0.820	116
SofLens TM One Day	Hilafilcon A	1.28	189 (± 9)	70	0.544	143
PureVision TM ^a	Balafilcon A	1.11	181 (± 3) ^c	36	0.886	124 ^e
			183 (± 2) ^d			

^a Both surface plasma treated (ST) and not treated (NT)

^b From reference [78]

^c For NT-PureVisionTM

^d For ST-PureVisionTM

^e Estimated by extrapolating T_g vs. w_1 data to $w_1 = 0$. Direct measurement was not possible because PureVisionTM lens material tends to decompose above 100 °C [39].

Table 2 Steady-state water-fluxes and diffusivities at 24.0 °C in soft-contact-lens materials in the absence of any air-side resistance. $\langle w_1 \rangle$ is the average water content in the SCL membrane at steady state; L_0 and L are initial and steady state membrane thicknesses; $\bar{D}_{lp}$ and D_{lp} are EMS and Fickian diffusivities, respectively. Fluxes are in (10^{-6} g cm $^{-2}$ s $^{-1}$) and diffusivities in (10^{-7} cm 2 s $^{-1}$) units.

Soft contact lens	RH (%)	n_1^∞	EMS			FICK		
			$\langle w_1 \rangle$	$\bar{D}_{lp}$	$\frac{L}{L_0}$	$\langle w_1 \rangle$	D_{lp}	$\frac{L}{L_0}$
HEMA	20	4.66 (± 0.06)	0.16	2.59	0.702	0.20	1.66	0.733
	50	4.70 (± 0.06)	0.20	3.70	0.743	0.23	2.12	0.769
	80	4.45 (± 0.04)	0.27	7.25	0.822	0.28	3.38	0.841
SofLens TM One Day	20	47.1 (± 1.0)	0.27	8.24	0.367	0.34	5.28	0.418
	50	65.7 (± 4.3)	0.32	16.0	0.404	0.38	9.23	0.450
	80	70.4 (± 17.2)	0.42	34.5	0.486	0.47	15.9	0.529
NT- PureVision TM	20	8.89 (± 0.14)	0.16	5.46	0.750	0.19	3.56	0.779
	50	7.95 (± 0.11)	0.20	6.94	0.789	0.22	4.08	0.812
	80	9.49 (± 0.21)	0.26	17.3	0.860	0.27	8.38	0.873
ST- PureVision TM	20	8.12 (± 0.22)	0.16	5.04	0.750	0.19	3.29	0.777
	50	8.05 (± 0.37)	0.20	7.11	0.789	0.22	4.18	0.811
	80	8.55 (± 0.23)	0.26	15.8	0.861	0.27	7.63	0.873

Table 3 Effect of evaporative cooling at the membrane/air interface on water/membrane diffusion rates. $\Delta T = T_L - T_0$ is the difference of interfacial (T_L) and bulk temperatures (T_0), and $\Delta n_1 = n_{1,iso} - n_1$ is the difference of water fluxes for the isothermal ($n_{1,iso}$) and non-isothermal case (n_1).

Soft contact lens	RH (%)	ΔT (°C)	$\Delta n_1 / n_{1,iso}$ (%)
HEMA	20	-0.81	2.67
	50	-0.32	1.06
	80	-0.16	0.52
SofLens TM one-day	20	-3.69	9.64
	50	-1.62	3.26
	80	-0.58	0.77
NT-PureVision TM	20	-1.48	4.70
	50	-0.53	1.67
	80	-0.29	0.84

Fig. 1

Fig. 2

Fig. 3

Fig. 4

Fig. 5