Effect of ambient temperature on the human tear film

Ali. A. Abusharha¹, PhD, E. Ian Pearce², PhD, Raied Fagehi,¹ PhD
¹Department of Optometry and Vision sciences King Saud University, Saudi Arabia
²School of health and life sciences, Glasgow Caledonian University, United Kingdom.

Correspondence
Ali. A. Abusharha
King Saud University, Riyadh 12372
FAX: 01413313387
Tel: 00966114693543
Email: aabusharha@ksu.edu.sa

E. Ian Pearce
70 Cowcaddence Road
Glasgow G4 0BA
FAX: 01413313387
Tel: 01413318201
Email: E.I.Pearce@gcu.ac.uk

Raied Fagehi
King Saud University, Riyadh 12372
FAX: 01413313387
Tel: 00966114693543
Email: rfagehi@ksu.edu.sa

Acknowledgement
The authors extend their appreciation to the college of Applied Medical Sciences Research Centre and the Deanship of Scientific Research at King Saud University for its funding for this research.
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Abstract

Purpose: During everyday life the tear film is exposed to a wide range of ambient temperatures. This study aims to investigate the effect of ambient temperature on tear film physiology.

Method: A Controlled Environment Chamber (CEC) was used to create different ambient temperatures (5°C, 10°C, 15°C, 20°C, 25°C) at a constant relative humidity of 40%. Subjects attended for two separate visits and were exposed to 25°C, 20°C and 15°C at one visit and to 10°C and 5°C at the other visit. The subjects were exposed to each room temperature for 10 minutes before investigating tear film parameters. The order of the visits was random. Tear physiology parameters assessed were tear evaporation rate, non-invasive tear breakup time (NIBUT), lipid layer thickness (LLT), and ocular surface temperature (OST). Each parameter was assessed under each condition.

Result: A three-fold increase in tear evaporation rate was observed as ambient temperature increased to 25 °C ($p=0.00$). The mean evaporation rate increased from 0.056 µl/min at 5°C, to 0.17 µl/min at 25°C. The mean NIBUT increased from 7.31 sec at 5°C to 12.35 sec at 25°C ($p=0.01$). A significant change in LLT was also observed ($p= 0.00$), LLT median ranged between to 20 to 40nm at 5 and 10 °C and increased to 40 and 90nm at 15, 20 and 25°C. Mean reduction of 4°C OST was observed as ambient temperature decreased from 25 to 5°C.

Conclusion: Ambient temperature has a considerable effect on human tear film characteristics. Tear evaporation rate, tear lipid layer thickness, tear stability and ocular surface temperature were considerably affected by ambient temperature. Chronic exposure to
low ambient temperature would likely result in symptoms of dry eye and ultimately ocular surface disorders.

**Keywords**: Tear film, dry eye, environmental factors, ambient temperature.
Introduction

Tear film function and structure can be altered by many different factors. Some of these factors are internal disorders such as insufficient tear production, meibomian gland dysfunction, hormonal changes and autoimmune disease. In addition, as the tear film is directly exposed to the ambient environment, tear film thinning and instability can be caused by external factors such as ambient temperature, humidity, air flow and pollution. Recently, researchers have shown an increased interest in the effect of the ambient environment on human body tissues. Symptoms including ocular itching, stuffy nose, dry throat, breathlessness, dry skin and headaches have been frequently reported by individuals working in adverse environmental conditions.

Ambient temperature is an external environmental factor that could affect the tear film. In addition to outdoor ambient temperature, the thermal conditions of commercial buildings and indoor workplace are not-well controlled. Unfortunately, to date, little attention has been paid to the relationship between the change in ambient temperature and tear film behaviour. One previous study has suggested that a relationship exists between tear film and atmospheric temperature. A significant difference in tear stability and production was found in normal subjects living in places characterized by a warm climate compared with those living in cold places. Another in vitro study investigating the effect of temperature on tear evaporation found that there was a threefold increase in evaporation rate as ambient temperature increased from 25 to 34°C. The relationship between ocular surface temperature and room temperature has been well documented. There has been little discussion about the effect of ambient temperature on tear film parameters. Most researchers to date tended to focus on the relation between OST and ambient temperature ignoring its effect on tear film parameters and the inter-relationship between these parameters. Therefore, the aim of this study was to determine
the effect of ambient temperature on ocular surface temperature and its effect on the ocular surface and tear film parameters. The inter-relationship between tear parameters and OST will be monitored at across range of different temperatures.

Method

Ethical approval was obtained from the Glasgow Caledonian University Human Ethics Committee. Subjects were healthy normals with no evidence of dry eye. Inclusion criteria was Ocular Surface Disease Index (OSDI) score of less than 12 and NITBUT of more than 10 seconds using HIRC AL grid. Twelve healthy normal subjects (3 female, 9 male, mean ± Sd 29.4±2.4 years) with no current ocular diseases or surgery were enrolled in this study.

As the control of ambient temperature and humidity was required during the experiment, a controlled environment chamber (CEC) (Weiss-Gallenkamp Ltd, Loughborough, UK) was used. The room temperature was set at 5 and 10°C on one visit. On the other visit, the CEC was set at temperatures of 15, 20 and then 25°C.

The subjects were divided into two groups (Group A and Group B). The order of visits was randomised in a cross-over design where the subjects in group A were exposed to a temperature of 25, 20 and 15°C and then exposed to 10 and 5°C ambient temperatures. Group B subjects were exposed to ambient temperature of 10 and 5°C, then at their second visit they were exposed to 25, 20 and 15°C. The relative humidity inside the CEC was maintained at 40% during the assessment of the tear film during the two visits. In order to minimize the possible implications of two visits, tear parameters assessments were conducted late morning and afternoon (after 11am) in both visits to avoid the effect of diurnal variation in ocular surface temperature.
A previous study investigating tear film parameters under different environmental conditions found a 6 to 10 minutes adaptation time was needed for the subjects before conducting ocular surface investigations. In this study, for the purpose of room temperature adaption, the subjects were exposed to each environmental condition for 10 minutes before starting investigations tear parameter.

The parameters assessed in this study were lipid layer thickness (LLT), tear evaporation rate (EVAP) non-invasive tear break-up time (NITBUT) and ocular surface temperature (OST). Tear break-up time and tear lipid layer thickness were assessed non-invasively by Keeler Tearscope Plus (Keeler Ltd, Windsor, UK). The Guillon and Guillon grading system was utilized to estimate the thickness of the tear lipid layer. Tear evaporation rate was measured using a Servo-Med Evaporimeter (Servo Med, Varberg, Sweden). A change in ocular surface temperature during exposure to the different ambient temperature was monitored with FLIR System ThermaCAM P620 (FLIR Systems, Surry, UK). A circle of approximately 4 mm diameter was placed at the estimated centre of the cornea and the mean temperature of this area was calculated. Thermal images of ocular surface were continuously recorded for one minute at frame rate of 30Hz. All temperature measurements were then exported to an Excel spreadsheet and 600 thermal values were selected with exclusion of the reading recorded immediately post-blink.

Variables were tested for normality using a Kolmogorov-Smirnov test. A repeated measure ANOVA and Tukey’s post-hoc test were applied for normally distributed data while the ordinal and data with non-normal distribution were analysed using Friedman’s test and post-hoc Wilcoxon rank-sum test. Correlation between tear parameters was assessed using Pearson’s test and Spearman’s test for the data with normal and non-normal distribution respectively.
Result

Lipid layer thickness

Changes in lipid layer thickness were found as temperature was altered (Figure 1A). The median grade of lipid layer thickness observed was grade 2 (20 - 40 nm) at 5 and 10°C and increased to grade 3 (40 - 90 nm) at 15, 20 and 25°C. Wilcoxon rank-sum test has shown that the lipid layer thickness was significantly thinner at 5°C in contrast to other ambient temperatures (p<0.05). Also a significant difference was observed at 10°C compared to lipid thickness at 20 (p=0.006) and 25°C (p=0.007). No significant change in lipid layer thickness was noticed when room temperature was changed from 20 to 25°C.

Tear evaporation rate

The box plot of tear evaporation indicates that, as the ambient temperature was increased, the tear evaporation rate also increased (Figure 1B). A three-fold increase in tear evaporation was observed as the ambient temperature increased from 5 to 25°C. The mean evaporation rate was 0.056 µl/min (20.11 g/m²/h) at 5°C, but increased dramatically to 0.17 µl/min (62.62 g/m²/h) when the room temperature was raised to 25°C. Tukey’s post-hoc test has been done for evaporation rate data. Statistically significant differences were observed in evaporation rate at 5°C when compared to 20°C (p=0.013) and 25°C (p=0.001).

NITBUT

Changes in tear film stability are shown in Figure 1C. At a room temperature of 5°C a significant reduction was found in NITBUT compared to those values obtained at all other temperatures (10, 15, 20 and 25°C), (p<0.05). The mean NITBUT values were 7.31 sec and 12.35 sec at 5°C and 25°C respectively. Measurements of NITBUT were statistically
analysed using Tukey’s post-hoc test. A significant change in NITBUT was observed at 10°C when compared to 20 (p=0.002) and 25°C (p=0.001).

**Ocular surface temperature**

Ocular surface temperature also showed a significant change (p<0.05). From the data in figure 1D it is apparent that there is a clear trend of decreasing OST as ambient temperature decreased. A reduction of 4°C OST was observed as ambient temperature decreased by 20º C (from 25 to 5 ºC).

Correlation tests were applied to determine the relationship between tear parameters as temperature was altered. Data from all temperatures were combined and analyzed using either Persons’s (R) for normally distributed data or Spearman’s (rho) correlation tests for ordinal and data with non-normal distribution. In Figure 2A a scatter plot shows a positive relationship between ocular surface temperature and evaporation rate (r = 0.45, p<0.05). Also, a significant correlation noted between evaporation rate and tear break-up time (r = 0.32, p<0.001) (Figure 2B). Tear evaporation was also showed a negative relationship with lipid layer thickness (rho = 0.43, p<0.001) (Figure 2C). Also a significant correlation between lipid layer thickness and tear film stability was found (rho= 0.49, p <0.001) (Figure 2D).

**Discussion**

The purpose of this study was to examine the effect of ambient temperature on tear film and the ocular surface. Tear stability, lipid layer thickness, evaporation rate and ocular surface were measured over a range of temperatures to evaluate the relationship between these parameters and temperature. In addition, the inter-relationship between these parameters was
examined to see how these are linked. This is an attempt to understand the relationship between these parameters in normal physiological conditions.

The evidence from this study suggests that the tear film lipid layer is affected significantly at low temperatures. It shows interference patterns characteristic of a thinner tear film after exposure to a temperature of less than 10°C. Hydrocarbon chains make up most of the lipid mass. It has been shown that lipid hydrocarbon structural order is affected by temperature.\textsuperscript{20} Moreover, a relationship between lipid hydrocarbon chain order and meibomian lipid delivery to the lid margin has been found.\textsuperscript{21} The melting range of meibomian lipids has been shown to be between 32.50 and 35°C.\textsuperscript{22,23} Previous work has shown that meibomian lipids become thicker as the ocular surface and eyelid temperature drops below 33°C, which may impede the normal delivery of meibomian lipids to the ocular surface.\textsuperscript{24} In addition to delivery, it has been shown that spreading over the tear film of meibomian lipids is also affected by temperature.\textsuperscript{23}

In the current study ocular surface temperature of less than 32 °C was observed after exposure to a room temperature of less than 10°C. Therefore, the changes in thickness and appearance of the tear film lipid layer observed in this study are likely to be due to variation in delivery and spread of meibomian lipid over the ocular surface.

A change in tear evaporation rate was observed when the ambient temperature changed. The evaporation rate increased as the temperature increased. The mean evaporation rate at 25°C was double that observed at 10°C, and triple the values recorded at 5°C. Box plots of evaporation rate at 25°C show that a quarter of participants had an evaporation rate characteristic of dry eye patients (evap>0.23µl/min\textsuperscript{25}). The lipid layer plays a critical role in controlling tear film evaporation. However, although a thin lipid layer was observed at 5 and 10°C, the evaporation rate was low in the cold environment, meaning that there may be
another factor apart from the meibomian lipid that could affect the evaporation of tears.

Decreased ambient temperature and OST result in decreasing the water molecule energy at the ocular surface, thus fewer molecules will be able to leave the ocular surface, which leads to a decrease in the evaporation rate.$^{26}$ Also, water-holding capacity of the atmosphere decreases with decreasing ambient temperature, therefore the atmosphere can hold very little moisture.$^{27}$ These two factors working together may explain the reduction in evaporation rate in cold conditions and the negative relationship observed between lipid layer thickness and evaporation rate at low temperatures.

This study shows significant changes in tear stability at low temperatures. The mean NITBUT value at 5°C was 7.31 seconds. It is accepted that tear film with a break-up time of less than 10 seconds is considered an unstable tear film (moderate dry eye – grade 2).$^1$ Moreover, recent work has demonstrated that the mean interblink interval (IBI) is 7.5 seconds.$^{28}$ In the current work, a NITBUT value of less than the typical IBI was recorded at 10 and 5°C, that may result in possible repeated exposure of the ocular surface. Consequently, frequent ocular surface exposure could lead to the development of the signs and symptoms of dry eye and visual disturbances. Previous studies have shown that tear stability is changed as tear film exposed to different ambient temperature.$^8$ Several studies have revealed that tear stability is the function of lipid layer integrity.$^{29-32}$ A change in the tear film lipid layer in a cold environment described earlier could be the reason behind the instability of tear film noted in this study.

In this study, it was found that ocular surface temperature decreased significantly as the ambient temperature decreased in 5°C steps. Pairwise comparison showed a significant difference in OST in all environmental conditions with the exception of that between 15 and 20°C ($p=0.093$). In the current study the mean change in central ocular surface temperature
was 0.18±0.40°C for each 1°C change in room temperature. A drop between 0.15 and 0.21°C in human ocular surface temperature for each 1°C reduction in ambient temperature has been reported.\textsuperscript{33,34} It has been suggested that a decrease in ambient temperature could result in significant reduction in corneal metabolic activity, which could be the reason behind the reduction in corneal temperature.\textsuperscript{35} However, it should be considered that metabolic rate may be not affected immediately with decreasing ambient temperature, therefore immediate change in OST could be resulted from another factor such as increasing evaporation rate.

**Conclusion**

This study has shown that ambient temperature influences tear film parameters. The current findings add to our understanding of the inter-relationship between tear film parameters and the physiological linkage between these parameters. The stability of tear film and lipid layer thickness were adversely affected at room temperatures of less than 10°C. NITBUT of less than the suggested IBI was recorded which may lead to exposure and to ocular surface pathology and a visual disturbance.

The evaporation rate showed a significant increase when the tear film was exposed to a room temperature of more than 20°C, and values of higher than the cut-off value of normal tear film were observed at a room temperature of 25°C.

Therefore, the effect of low or high ambient temperature on the tear film should not be neglected by people who work outdoors and spend a long time in adverse climate conditions or individual who play winter sports such as skiing and ice skating. In the same way, indoor workplaces need to be able to be maintained at a healthy and comfortable temperature range to ensure ocular comfort and well-being.


**Figures legend**

**Figure 1.** Comparison of the distribution of lipid layers pattern over range of ambient temperature. Thicker lipid pattern was observed in higher room temperature (A). A box plot showing tear evaporation rate (B) measured at 5, 10, 15, 20 and 25 °C. Significant increase in tear evaporation was found as ambient temperature was increased. NITBUT (C) became shorter when tear film was exposed to lower temperature. The tear break up time at 5°C was significantly shorter than all other room temperatures. Significant difference was seen in ocular surface temperature (D) between all ambient temperatures with exception of that between 15 and 20°C (*n*=12, *p*=0.093). The box represents the interquartile range that contains 50% of the values. The whiskers are lines that extend from the box to the highest and lowest values, excluding outliers (O) that are 1.5 to 3 box lengths from the upper and lower edge of the box and extremes (*) that are more than 3 box lengths. The line across the box indicates the median value. Pairwise significant differences are shown (*).

**Figure 2.** Scatter plot showing correlation between ocular surface temperature and evaporation rate (A), evaporation rate and tear break up time (B), evaporation rate and lipid layer thickness (C), tear break up time and lipid layer thickness(D).
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Figure 2. Scatter plot showing correlation between ocular surface temperature and evaporation rate (A), evaporation rate and tear break up time (B), evaporation rate and lipid layer thickness (C), tear break up time and lipid layer thickness (D).