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A novel ex vivo model of equine corneal wound healing

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A novel *ex vivo* model of equine corneal wound healing

by

**Rita Fay Wehrman**

A thesis submitted to the graduate faculty

in partial fulfillment of the requirements for the degree of

**MASTER OF SCIENCE**

Major: Veterinary Clinical Sciences (Veterinary Medicine)

Program of Study Committee:
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Iowa State University

Ames, Iowa

2017

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I dedicate my thesis work to my husband Steve Sanders. You make every day better and I cannot thank you enough for going on this journey with me.
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ABSTRACT

Corneal disease is a significant cause of decreased vision or blindness in the horse worldwide. Numerous topical and systemic drugs over a prolonged period of time may be used to treat an equine corneal ulcer. Despite the high prevalence of corneal disease and ulceration in the horse, and the complexity of corneal wound healing, research is limited and more heavily devoted to in vivo studies whose limitations include small sample sizes and ethical considerations. Bench top laboratory in vitro studies are limited in number and overlook the complexity of corneal wound healing by assessing each corneal layer separately from its neighbor.

The objective of this study was to develop a novel ex vivo model of equine corneal wound healing that maintained structural integrity, and is physiologically relevant, mimicking in vivo events.

In this randomized and controlled study, fourteen equine corneas were harvested within two hours of humane euthanasia for reasons unrelated to this study. All donors were screened and corneas were included if they were free of ocular and systemic disease. Corneas were immediately processed and corneoscleral rims excised 2 mm posterior to the limbus. Corneas were randomly assigned to one of the following groups: wounded (n=8) or unwounded (n=6) controls. Each pair of eyes was divided such that corneas from the same horse were assigned to a different group. In the event that only one of the two eyes was included, the eye was assigned to the wounded group. In the ‘wounded’ group, the axial cornea was wounded for 60 seconds
with a 6 mm disk of filter paper soaked in 1N sodium hydroxide (NaOH). Epithelial ulceration was confirmed employing fluorescein stain. Subsequently, all corneas were cultured using an air-liquid interface model in complete media. The corneal cups were placed on a rocker set at a 6 degree incline and incubated at 37 °C with 5% CO₂ humidity. The media bathed the cornea 8 times per minute to simulate normal horse blinking. Evaluation of corneal healing was performed daily and included fluorescein staining and fluorescein retention (i.e. ulcer measurements). Corneas from both groups were randomly assigned to undergo further processing via histopathology at designated time points of 24 hours (T24; n=5), T48 (n=5), and T72 (n=4) post wounding, and stained with hematoxylin and eosin. Structural integrity and pathologic changes to the corneal epithelium, stroma, Descemet’s membrane and endothelium were assessed. Interleukin-6 (IL-6) expression was then evaluated by means of RNAscope in situ hybridization.

All wounded corneas healed within 72 hours. Histologically, normal corneal architecture was observed including intact epithelium, minimal stromal edema, and presence of endothelium. Increased IL-6 expression was noted in wounded corneas compared to unwounded controls and was predominantly in the metabolically active basal epithelial cell layer. As the corneal wound healing progressed, IL-6 expression decreased.

The equine air-liquid interface, ex vivo, corneal wound healing model is an effective and physiologically relevant model. This model can be utilized to reduce or
eliminate the use of live horses for corneal wound healing studies, while providing an effective platform for corneal wound healing research in this species.
CHAPTER 1. GENERAL INTRODUCTION

Structure of the Equine Cornea

The cornea and sclera are part of the fibrous tunic of the eye and their junction is known as the limbus. The cornea makes up approximately 20% of this outer fibrous tunic and due to its transparency, curvature, and air liquid interface with the outer precorneal tear film (PTF) and the inner ocular media, is the most refractive part of the eye.\(^1,2,3\) Because of its anterior location, its purpose is also to protect the inner contents of the eye.\(^4\) This elliptically shaped structure is comprised of four main layers in the horse, including the epithelium, stroma, Descemet’s membrane, and the endothelium (Fig. 1-1).\(^1,2\) In the horse, the average total thickness of the cornea is between 700 and 900 µm.\(^5,6,7\)

Epithelium

The corneal epithelium is the anterior most layer of the cornea and is a continuation of the conjunctival epithelium (Fig. 1-1). It is in contact with the PTF superficially, which also aids in light refraction. The main function of the epithelium is to serve as a barrier against infection, thereby protecting the rest of the eye.\(^2\) As the cornea is maintained in a state of relative dehydration, any loss of this hydrophobic layer (i.e. ulceration) may compromise the cornea’s transparency via an influx of water from tears.\(^1\) Clinically this is appreciated as corneal edema and experimental removal of this layer results in a 200% increase in corneal thickness within 24 hours.\(^8\) In the horse, this layer is up to 15 cell layers thick and consists of three cell
types, which from anterior to posterior are known as superficial epithelial cells, wing cells, and basal cells.\textsuperscript{9,10,11} These different cell types communicate with themselves and one another using different intercellular junctions including desmosomes, gap junctions, and zonula occludens.\textsuperscript{1,12} These modes of communication contribute to expedient wound healing.\textsuperscript{13} As the superficial epithelium is constantly being sloughed off, or desquamating, the deeper wing cells proliferate and move towards the corneal surface, while the basal cells then differentiate into wing cells. Additionally, corneal stem cells are housed in the limbus, or corneaoscleral junction, and as the three cell types push anteriorly, the stem cells migrate centripetally to ‘re-stock’ the basal and superficial layers.\textsuperscript{14,15}

**Stroma**

The stroma is deep to the epithelium, comprises about 90% of the total corneal thickness and is 78% water (Fig. 1-1).\textsuperscript{1} This layer is made up of collagen fiber bundles known as lamellae which run parallel to, and across the length of the cornea. The stroma is further subdivided into the anterior stroma and posterior stroma. The anterior portion is considered to be less orderly and the lamellae are smaller in both thickness and width compared to the posterior stroma.\textsuperscript{1} They are arranged in a more oblique, or woven, orientation. Deep to the anterior stroma is the posterior stroma, whose broad sheets of lamellae are lined up in a more parallel, uniform, nonwoven orientation. In general, the stroma’s collagen fibrils are more precisely arranged with fewer spaces between the fibrils in the central zone versus the peripheral. This specific organization of the corneal stroma also plays a large
role in maintaining corneal clarity.\textsuperscript{16,17} Distributed in between the lamellae is the predominant stromal cell type, known as the keratocyte, which communicate via tight and gap junctions. Other cells that may reside normally within the stroma include macrophages, dendritic cells and lymphocytes.\textsuperscript{1,18}

\section*{Descemet's membrane}

The next corneal layer is Descemet’s membrane, which is the basement membrane of the endothelium (Fig. 1-1). It is also composed of collagen similar to the stroma; however, this collagen is more layered, which gives it an elastic property that is more resistant to trauma. Descemet’s membrane is approximately 10 µm thick and it thickens with age. This thin layer is thought to have regenerative properties.\textsuperscript{15}

\section*{Endothelium}

The inner most layer of the cornea is the endothelium (Fig. 1-1). In this delicate, 5 µm monolayer, cells are joined to one another via tight and gap junctions. This layer’s function includes actively pumping aqueous humor via Na\textsuperscript{+}/K\textsuperscript{+}-ATPase pumps back into the anterior chamber, and it is most responsible for maintaining a state of relative dehydration in the cornea.\textsuperscript{1,8} This has been demonstrated experimentally, in which removal of the endothelium resulted in an increase of greater than 500% in overall corneal thickness.\textsuperscript{8} Most mammals are born with their total, maximum amount of endothelial cells, which slowly decreases with time. In the
horse, this endothelial cell density has been established at 3155+/-765 cells/mm\(^2\).\(^7\) As this layer cannot generally undergo mitosis, it ‘heals’ itself by cellular spreading, causing changes in cell shape (pleomorphism) and size (polymegethism). Should the endothelial cellular density decrease by too large an index to pump fluid efficiently, severe, diffuse corneal stromal edema will result.\(^{15}\) In the dog, this has been established at less than 500 to 800 cells/mm\(^2\) and is known as corneal decompensation.\(^1\) To date, this threshold has not been determined in the horse.

While the endothelium has the most active job in maintaining corneal transparency, compromise of any of the aforementioned layers will result in corneal pathology and an impaired visual axis. Corneal clarity relies upon many factors including the presence of each layer, appropriate architectural integrity, and the individual cell’s ability to communicate with itself and its neighbors. These factors are the cornerstone of corneal wound healing.

**Corneal Wound Healing**

Corneal injuries are vision and eye threatening across species. Corneal healing is a complex process that involves all cellular layers and their ability to communicate via mediators with one another. Each layer responds differently to insult.

**Epithelial healing**

The corneal epithelium has stem cells located within the limbus of the eye. These corneal limbal stem cells are able to provide the cornea with a lifetime of
The objective of epithelial wound healing is two-fold: first, to trigger migration of the epithelial cells to the wounded area, and second to upregulate cell proliferation and differentiation, all with the end goal of filling the defect and therefore healing the wound. This is done via autocrine and paracrine cellular mediated signaling with a significant amount of ‘cross talk’ between the corneal epithelium, limbal stem cells, and the underlying stromal cells. Initial factors that are released following epithelial wound healing include interleukin-1 (IL-1) and tumor necrosis factor alpha (TNF-α) and growth factors, that trigger the epithelial cells migration and then proliferation. Interleukin-6 (IL-6) also plays a large role, and has been shown to significantly increase epithelial wound healing rates in a rabbit in vivo model and to stimulate epithelial cell migration in rabbit in vitro models. Additionally, matrix metalloproteinases are released, which cause the cells to disengage from one another and to release their cell matrix adhesions.

**Stromal healing**

Due to this layer’s thickness, it goes without saying that the organized healing of the stroma is of paramount importance. Should this layer heal without the precision it requires, there will be a reduction in corneal transparency (i.e. fibrosis, or scar, formation), irregular light refraction, and therefore impaired vision. Damage to the stroma causes a large release of cytokines, that trigger apoptosis of underlying keratocytes. Once cleared of potentially damaged keratocytes, the remaining activated keratocytes transform into migratory fibroblasts and myofibroblasts. Fibroblasts secrete actin, a contractile cytoplasmic element, while myofibroblasts express a contractile cytoskeleton element known as alpha-smooth muscle actin (α-
It is well known that the myofibroblast is only detected following stromal corneal wounding and its expression of α-SMA is credited with wound contracture. However, this spindeloid myofibroblast cell and its activity with the extracellular matrix (ECM), is at the heart of fibrosis formation within the cornea as the myofibroblast deposits new extracellular matrix and aberrant proteins to fill up the defect. With time, these proteins can form scar tissue and as the ECM undergoes very slow turnover (years in humans) scar formation may persist.

As such, the “well meaning” myofibroblast has been linked to corneal haze and scar formation following wounding. Inflammatory cells including monocytes, macrophages, polymorphonuclear leukocytes, natural killer cells, and T cells also have a role in stromal wound healing. It is thought these cells enter the cornea at the limbus or through the circulatory system and tear film. The remodeling of the stroma is both complex and lengthy as the anterior and posterior stroma will take months to regain their normal, unique collagen fibril orientation.

**Descemet’s membrane and endothelial healing**

Descemet’s membrane has regenerative properties, but similar to the endothelium, has not been studied extensively due to its more interior location relative to the outer stromal and epithelial layers. Compared to epithelial and stromal wound healing, relatively less is known about endothelial wound healing, again due to this layer’s inner location within the cornea. However, unlike the epithelium, the endothelium undergoes little to no proliferation, instead closing wound gaps by migration and spreading. In the rabbit, a transient morphology to
a myofibroblast-type, α-SMA expressing migratory cell has been noted.⁡²⁵,⁵³,⁵⁴ To
date, this has not been demonstrated in the horse.

**Cellular communication**

There is a significant amount of ‘cross talk’ between the corneal layers, and
while each layer heals uniquely, the overall tissue architectural organization lends
itself to cellular communication.⁤⁵⁵ These corneal cells release mediators secondary
to injury that are distributed throughout the cornea using gap junctions, tight
junctions, and zonula occludens.¹,¹³,¹⁸ Therefore, models of corneal wound healing
that include all corneal layers and that maintain normal corneal architecture are of
utmost importance.⁤⁵⁶ While in vitro assessment of cell lines is certainly important and
not to be overlooked due to its relatively low cost and high speed, this type of over
simplified model is not able to mimic the complexities of the interactions both at the
corneal cellular level, nor at the corneal organ level.⁤⁵⁵ Further exploration of the
interlayer communication in corneal wound healing may further advance our
understanding of this complex process.

**Interleukin-6 Cytokine Activity**

Interleukin-6 is a potent immunologic mediator and it is involved in acute
phases of inflammation.⁤⁷ It is considered a myokine, or a cytokine produced from
muscle, and when released by myocytes, serves a protective, anti-inflammatory role
within the musculature.⁤⁵⁸,⁵⁹ However, it is also a proinflammatory cytokine⁤⁷ and its
secretion from T cells and macrophages leads to inflammation and tissue damage
especially following burn trauma. \cite{58-60} Interleukin-6 has been extensively studied within the context of corneal wound healing and its expression is upregulated following corneal injury in humans, as well as in animal and \textit{in vitro} models.\cite{60-64}

While the indirect suppression of IL-6 has been shown to have a negative effect on \textit{in vivo} corneal epithelial cell migration in mice\cite{63} and on \textit{in vitro} immortalized human corneal epithelial cell migration,\cite{61,63} increases in IL-6 expression actually expedite corneal wound closure in scratch assays employing human\cite{62,65} and rabbit corneal epithelial cells,\cite{66} and in \textit{in vivo} rabbit corneal wound healing models.\cite{67} Interleukin-6 has also been shown to increase in equine disease processes such as dermatologic epidermal wound healing,\cite{68} pneumonia,\cite{69} and endometritis.\cite{70} It is likely that IL-6 is one of the more systemically important equine cytokines. Although IL-6 expression has been demonstrated in rabbits,\cite{66,67} mice,\cite{60,62} and humans\cite{61,64,71} post corneal wounding, its role in equine corneal wound healing has not been studied to date.
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Figure 1-1. Histologic section of a normal equine cornea. Note the superficial epithelial layer, consisting of approximately 15 epithelial cell layers, the well-organized, thick, stromal layer, Descemet’s membrane, and the single-cell, inner-most, endothelial layer (black arrow).
CHAPTER 2. LITERATURE REVIEW

In vivo Corneal Wound Healing Studies

Ocular disease is one of the most common ailments of the horse\(^1\) and it is one of the most significant causes of decreased vision or blindness in the horse world-wide.\(^2\) This is likely due to several factors including a prominent globe, the lateral location of the globe within the skull, and the equid’s exuberant and head throwing behavior.\(^1\) As such, the horse is predisposed to corneal injury. Although equine corneal disease and the subsequent wound healing cascade is a complex process whose mechanisms are still not fully understood,\(^3\) it is known that inflammatory cells and their secretory products such as cytokines and growth factors are involved, as discussed in Chapter 1. In addition, underlying genetics may also play a role\(^3\) and it has been suggested that the aggressiveness of the corneal destruction may be linked to the equine host’s immunophenotype.\(^4\) This includes the local innate response of Pattern and Damage Recognition Receptors (PRRs, DRRs) that reside within the corneal epithelium and their role in cytokine recruitment.\(^4\) Hypothesis driven, in vivo, corneal wound healing studies have been conducted extensively in mice, rats, and rabbits,\(^5-10\) and included evaluation of corneal regeneration and healing processes, corneal epithelial cell migration, and IL-6 expression.\(^11\) Additionally, the role of IL-6 in rabbit,\(^12,13\) mouse,\(^5,14,15\) and human\(^16\) corneal wound healing has also been studied in vivo.

Despite the high prevalence of corneal disease and ulceration in the horse, the importance of a clear visual axis in this species, and the complexity of corneal
wound healing, few in vivo studies are hypothesis driven. Instead, most studies are heavily devoted to retrospective studies, reviewing patients’ medical records.\(^{17-24}\) Only few hypothesis driven corneal wound healing studies have been conducted in vivo to date.\(^{25,26}\) The limitations of such studies include small sample sizes, high cost, ethical concerns, and large variation amongst few subjects which often results in a lack of statistical significance.\(^{27}\) An ex vivo equine corneal wound healing model can overcome these limitations, and provide the platform for hypothesis driven studies of corneal wound healing in the horse.

**In vitro Corneal Wound Healing Studies**

In the laboratory setting, in vitro work possesses its own set of limitations. Cell culture lines that are purchased from commercial sources may be contaminated with other cell lines, may not be validated for the cellular phenotype in question, and may be over-passaged.\(^{28-31}\) Growing primary corneal cell cultures can prove to be a challenge as well. For example, while Mathes et al successfully established a reproducible cell culture model of equine keratocyte and endothelial cells, the epithelial cells grew slowly, did not survive multiple passages and were seen to suffer from keratocyte or fungal organism contamination.\(^{27}\) Additionally, by experimenting on individual corneal layers separately from neighboring layers,\(^{32,33}\) in vitro corneal research may be an overly simplified model which lacks the dynamic, physiological events of the in vivo corneal model.\(^{32-34}\) In vitro cell culture lines are not able to simulate the cornea’s complex molecular processes nor the interactions and ‘cross talk’ amongst the corneal layers, especially the epithelium and stroma. For
example, it is well known that epithelial and stromal interactions are mediated by paracrine expression and that autocrine signaling is likely occurring as well. By studying and experimenting on one corneal layer at a time, it is likely the cell to cell and cell to extracellular matrix interactions (the ‘cellular microenvironment’) are overlooked. These intercellular interactions are paramount to the normal routine of the cell, and tissue injury can result in disruption of these connections and therefore cellular communication. Therefore, the cells’ ability to respond to its local environment is impeded, and cell migration, differentiation and healing are diminished as the individual cell is not able to communicate the signaling information on to groups of other cells. As such, the in vitro researcher may be faced with the challenge of obtaining clinically applicable results.

**Current Ex vivo Studies**

The inherent challenge comes in the desire to learn more about corneal wound healing without either harming healthy animals in vivo or struggling with clinical applicability in vitro. The ex vivo model of corneal wounding has been developed in other species including the human, dog, pig (authors unpublished data), and rabbit, and represents a logical stepping stone between the equine in vitro ocular bench top models and the in vivo clinical studies, all while providing physiologically-based, clinically applicable conclusions. Recent publications have revolutionized the ex vivo approach, demonstrating superior epithelial preservation using an air-liquid interface human ex vivo model. Deshpande et al refined the ex vivo corneal wound healing model, by ensuring that media rolled over the cultured
rabbit corneas. The technique resulted in an extended corneal lifespan of up to 4 weeks, versus 2 weeks in static conditions. This modified technique allowed them to evaluate the inflammatory role of the cytokine interleukin-17A on their rabbit corneas over an extended period of time. Previously, Lin et al had evaluated the shorter term effects of interleukin-33 in a modified human ex vivo model. However, to our knowledge, IL-6 expression has not been studied utilizing the ex vivo corneal wound healing model.

A species specific, reproducible model of corneal wound healing is of great importance. To date, no ex vivo models of equine corneal wound healing have been established. Therefore, the purpose of this study was two-fold. First, to establish an ex vivo model of equine corneal wound healing, and second, to investigate the physiologic relevance of this model. This study is at the heart of improved animal welfare and represents a logical stepping stone between the in vitro bench top models and the in vivo clinical studies.

The successful development of this model could serve to decrease in vivo corneal wound healing studies in the horse, while providing an effective platform for equine corneal wound healing research.
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CHAPTER 3. A NOVEL EX VIVO MODEL OF EQUINE CORNEAL WOUND HEALING

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ABSTRACT

**Purpose:** To establish a physiologically relevant *ex vivo* equine corneal wound healing model.

**Methods:** Fourteen globes were harvested from healthy horses within two hours of humane euthanasia for reasons unrelated to this study and immediately processed. Corneas were randomly assigned to one of the following groups: wounded (n=8) or unwounded (n=6) controls. In the wounded group, the axial cornea was ulcerated with a 6 mm filter paper disk soaked in 1 N sodium hydroxide (NaOH) at Time 0. Corneoscleral rims were excised 2 mm posterior to the limbus and corneal cups cultured in complete medium using an air-liquid interface model. The medium bathed the corneal cup eight times per minute to simulate normal horse blinking. Evaluation of corneal healing was performed daily and included fluorescein staining and a modified Hackett-McDonald ocular scoring system which included the calculated ulcer area. Corneas were randomly assigned to undergo further processing via histopathology at 24 hour (T24; n=5), 48 hours (T48; n=5), and 72 hours (T72; n=4) post wounding. Additionally, RNAscope *in situ* hybridization assay was performed according to the manufacturer’s instructions to evaluate IL-6 expression.

**Results:** All wounded corneas healed within 72 hours. Histologically, normal corneal architecture was seen including intact epithelium, minimal stromal edema and the presence of endothelium. Interleukin-6 expression increased in wounded equine corneas compared to unwounded controls. The greatest IL-6 expression was in the basal epithelial layer, close to the wound margin. As the corneal wound healing progressed, IL-6 expression decreased.
Conclusions: The equine air-liquid interface, *ex vivo*, corneal wound healing model is an effective and physiologically relevant model. This model can be utilized to reduce or eliminate the use of live horses for corneal wound healing studies, while providing an effective platform for corneal wound healing research in this species.

INTRODUCTION

Corneal ulcers are very common in horses, and with a lack of appropriate treatment, they present an imminent risk for the eye and for vision.\(^1\) If not identified early or if secondary infection occurs, keratomalacia caused by various proteases acting on corneal collagen, may develop. This condition may lead to perforation of the eye. However, despite the horse being predisposed to corneal injury (large eye size, lateral skull placement, exuberant behavior) and the frequency of ophthalmic presentations to the veterinarian,\(^1\) our knowledge base of equine corneal ulceration resolution is lacking.

Corneal wound healing is a complex process that involves two basic steps – cellular migration to cover the wound followed by mitosis of the epithelial cells to fill in the defect.\(^2,3\) Physical damage to the cornea’s outer most layer, the epithelium, is the most common cause of wounding and it is associated with signs of ocular pain including blepharospasm and lacrimation. As the injured eye runs the risk of permanent, vision altering consequences or even perforation, it is not surprising that prompt and aggressive treatment is of paramount importance. Treatment of these corneal disorders can be time consuming, costly and frustrating for both the owner
and the patient, and should a superficial, uncomplicated erosion not be healed within 72 hours, it should be considered complicated and is therefore in danger of perforation. In these cases, painful inflammation is likely involved, and clinical signs of reflex uveitis may be present and include photophobia, corneal edema and aqueous flare.

Despite the high prevalence of corneal disease and ulceration in the horse, the importance of a clear visual axis in this species, and the complexity of corneal wound healing, few in vivo studies are hypothesis driven. Instead, most studies are heavily devoted to retrospective studies, reviewing patients’ medical records. Only a few prospective corneal wound healing studies have been conducted in vivo to date. The limitations of such studies include small sample size, high cost, ethical concerns, and large variation amongst few subjects that often results in a lack of statistical significance.

In the laboratory setting, in vitro work possesses its own set of limitations including obtaining clinically relevant results. By experimenting on individual corneal cell types separately from neighboring layers, in vitro research may be an overly simplified model that lacks the dynamic, physiologic events of the in vivo model. In vitro pure cell culture lines are not able to simulate the cornea’s complex molecular processes nor the interactions and ‘cross talk’ amongst the corneal layers, especially the epithelium and stroma. For example, it is well-known that epithelial and stromal interactions are mediated by paracrine expression and that autocrine signaling is likely occurring as well. Therefore, the in vitro researcher may be faced with the obstacle of obtaining clinically applicable results.
The inherent challenge comes in the desire to learn more about these involved processes without either harming healthy animals in vivo or struggling with clinical applicability and physiologic relevance in vitro. The ex vivo model has been developed in other species,\textsuperscript{20,21} and represents a logical stepping stone between the equine in vitro ocular bench top models and the in vivo clinical studies, all while providing physiologically-based, clinically applicable conclusions.\textsuperscript{17,22} While effective, species specific, reproducible models of corneal wound healing are of paramount importance, the development of an equine ex vivo model could serve to decease in vivo corneal wound healing studies in the horse, all while maintaining the clinical relevance of the results.

Interleukin-6 (IL-6) is a potent immunologic mediator and it is involved in acute phases of inflammation.\textsuperscript{23} It is considered a myokine, or a cytokine produced from muscle. Upon its release from myocytes, it serves a protective, anti-inflammatory role within the musculature.\textsuperscript{24,25} However, it is also a proinflammatory cytokine and its secretion from T cells and macrophages leads to inflammation and tissue damage especially following burn trauma.\textsuperscript{24,25} Additionally, IL-6 has been shown to increase in septic conditions in both humans and horses\textsuperscript{26} and high levels of IL-6 are expressed in equine disease processes such as endometritis\textsuperscript{27} and pneumonia.\textsuperscript{28} Previous equine dermatologic studies have shown significant increases in IL-6 mRNA levels following pathogen-associated molecular pattern (PAMP) exposure to gram negative bacteria\textsuperscript{29} and in an epidermal wound healing model.\textsuperscript{30} Equine neutrophils express significantly more IL-6 than other mediators.\textsuperscript{31} Given this, it is likely that IL-6 is one of the more systemically important equine
cytokines and may play a large role in equine corneal wound healing as well. Although IL-6 expression has been demonstrated in rabbits, mice, and humans post corneal wounding, its expression during corneal wound healing in the horse has not been studied to date.

Therefore, the purpose of this study was two-fold. First, to establish an ex vivo model of equine corneal wound healing. And second, to investigate the physiologic relevance of this model by evaluating the rate of epithelial wound healing, anatomical structure, and IL-6 expression in wounded and unwounded corneas. This study is at the heart of improved animal welfare and represents a logical stepping stone between the in vitro bench top models and the in vivo clinical studies.

MATERIALS AND METHODS

Corneas

Corneas were obtained from horses that were euthanized at the Iowa State University College of Veterinary Medicine Lloyd Veterinary Medical Center for reasons unrelated to this study. Fourteen eyes from eight horses were included. The median age and weight of the donors was 19.25 years (range 8-30 years) and 370 kilograms (range 160-540 kgs), respectively. Of these eight horses, three were geldings and five were mares and breeds represented included three Quarter Horses/Quarter Horses crosses, 3 ponies, one Paint and one Peruvian Paso. All
horses were free of systemic and ocular disease. Two eyes (one right and one left) were excluded due to ophthalmic disease.

**Corneal wounding**

Immediately following humane euthanasia, globes were enucleated, rinsed in 2% betadine (Select Medical Products, Jacksonville, FL, USA), and placed in 1x phosphate buffered saline (PBS; Life Technologies, Grand Island, NY, USA). All further processing occurred within two hours of humane euthanasia. Globes were randomly assigned to wounded (n=8) or unwounded (n=6) control groups. If one of the eyes was excluded the fellow eye was assigned to the wounded group. Wounded corneas included 4 right eyes and 4 left eyes, while unwounded corneas included 3 right eyes and 3 left eyes. During all processing steps, a delicate task wiper (Kimwipes, Kimberly-Clark, Roswell, GA, USA) wet with 1X PBS was used to facilitate holding of the eyeball. For the eight wounded corneas, an alkali burn model was employed in which a 6 mm filter paper disc soaked in 1 N sodium hydroxide (NaOH) was applied to the axial cornea for 60 seconds (Fig. 3-1). After wounding, 1X PBS was used to gently rinse the globe. Fluorescein stain (fluorescein sodium, Akron Inc., Lake Forest, IL, USA) was applied to confirm epithelial erosion and stromal exposure, and baseline ulceration measurements were obtained using a millimeter scale bar (Fig. 3-2). Wounding the cornea with the globe intact allowed for a crisp, well demarcated corneal ulceration. Once wounded, a #20 Bard Parker blade was used to score and then excise the sclera tangentially, approximately 2 mm posterior to the limbus, freeing the corneoscleral (“corneal”) cup (Fig. 3-3). The
non-wounded globes were processed in a similar fashion, excluding the alkaline burn.

**Ex vivo cultures**

Each corneal cup was placed in an individual 100 mm petri dish (Fischer Scientific, Waltham, MA, USA) along the dishes’ edge. Placement of the corneal cup along the side of dish minimized motion of the corneal cup while it was on the rocker. Additionally, great care was taken to ensure that the corneal cup retained its convex shape while introduced to the petri dish as no conformers were used. Once placed in the petri dish, approximately 25 mL of medium was added such that the medium came to the limbus but did not float the corneal cup off the floor of the petri dish. The medium consisted of Dulbecco’s modified Eagle’s medium and Ham’s F12 medium (DMEM/HamF12 + GlutaMAX™, Gibco Life Technologies, Grand Island, NY, USA) in a 1:1 ratio. The medium was supplemented with 10% fetal bovine serum (Fischer Scientific, Waltham, MA, USA), 2.5µg/mL Amphotericin B (Life Technologies, Grand Island, NY, USA), 100 IU/mL penicillin (Life Technologies, Grand Island, NY, USA), 100 µg/mL streptomycin (Life Technologies, Grand Island, NY, USA), 10 ng/mL epidermal growth factor (Gibco Life Technologies, Frederick, MD, USA) and 5 µg/mL bovine pancreatic insulin. The petri dishes were placed on a rocker (Benchmark Scientific, Edison, NJ, USA) that was set at a 6 degree incline and 8 rocks per minute to actively bathe the corneal cup in medium in order to simulate blinking (Fig. 3-4). All corneas were incubated at 37 °C with 5% CO₂ humidity and monitored daily. Petri dishes and medium were changed out daily.
Corneal wound healing assessment

All daily evaluations for ulcer size were performed using slit-lamp biomicroscopy (Keeler SL4 PSL Classic, Keeler Instruments Inc., Windsor, UK) and a modified Hackett-McDonald ocular scoring system.\(^3\) Ulcer size was measured daily utilizing a millimeter scale, following application of fluorescein stain (Fig. 3-5, 3-6), and the ulcer area was calculated \((A = \pi r^2)\) and averaged for each time point. Three and two corneas from the wounded and non-wounded group respectively, were submitted for RNAscope in situ hybridization and histopathology analysis 24 (T24) and 48 (T48) hours post wounding, and two corneas from each group were submitted at 72 hours (T72) post wounding.

Modified Hackett-McDonald Ocular Scoring System

The modified Hackett-McDonald scoring system is used frequently in ophthalmology toxicology studies to describe ocular changes,\(^3\) and was employed in this study to describe corneal changes:

a) Fluorescein intensity staining was graded 0-4 with 0=no fluorescein uptake; 0.5=no true uptake/epithelial erosion; 1=faint intensity stain; 2=mild intensity stain; 3= moderate intensity stain; 4= marked intensity stain.

b) Ulcer location and description were noted.
c) Corneal edema was graded 0-4: 0=normal cornea with no edema; 1=slight loss of corneal transparency, all localized around the edge of ulceration; 2=mild loss of corneal transparency, with edema extending outward from the edge of ulceration; 3=moderate loss of corneal transparency with further extension of edema from ulceration's edge; 4= complete loss of corneal transparency with marked edema diffusely.

**Histopathology**

Corneal histopathology was performed on all corneas to assess structural integrity and pathologic changes to the corneal epithelium, stroma, Descemet’s membrane and endothelium. Each sample was fixed in 10% neutral buffered formalin (NBF) for 24 hours and then routinely stained with hematoxylin and eosin. All samples were evaluated using standard bright field microscopy.

**IL-6 RNAscope in situ hybridization**

RNA in situ hybridization (RNAscope, Hayward, CA, USA) was used in which single RNA IL-6 molecules within individual cells were visualized. All recommended RNAscope assay procedures were followed according to manufacturer's instructions for formalin fixed paraffin embedded (FFPE) sections using RNAscope FFPE Reagent Kit 2.0 (Advanced Cell Diagnostics, Hayward, CA, USA). Once pulled from culture, corneas were placed in 10% NBF for 24 hours and then paraffin-embedded in preparation for IL-6 gene expression (RNAscope Target Probe C1, 16ZZ Ec-IL6, Advanced Cell Diagnostics, Hayward, CA, USA). The embedded tissue was then cut
into 5 µm sections, mounted on slides and allowed to air dry overnight at room
temperature. The following morning, slides were baked for one hour at 60°C,
deparaffinized using a xylene immersion twice for 15 minutes and then dehydrated
in an ethanol series. Once dry, the slides underwent three pretreatment steps, for 30
minutes each, to ensure access to corneal tissues. After the first pretreatment, the
buffer was rinsed off the slides with deionized water and the slides were boiled at
98-104°C for 30 minutes on a hot plate with pretreatment 2 buffer. The slides were
again rinsed in deionized water and treated with pretreatment 3 buffer for 30 minutes
at 40°C in the HybEZ™ hybridization oven (Advanced Cell Diagnostics, Hayward,
CA). This allowed for target probe access to the corneal tissue and blocking of
endogenous peroxidase activity. The 16ZZ Ec-IL6 target probe was then applied to
each 5 µm tissue section and the slides placed back in the HybEZ™ hybridization
oven for 2 hours at 40°C. After washing, a series of amplifiers was applied to each
slide and they were then placed back in the oven. This included treatment with Amp
1 solution for 30 minutes, Amp 2 solution for 15 minutes, Amp 3 solution for 30
minutes and Amp 4 solution for 15 minutes. Each of these amplifying steps was
baked at 40°C. Amp 5 and 6 solutions were applied for 30 and 15 minutes,
respectively, at room temperature. After each of the six amplifying steps, slides were
washed with wash buffer for two minutes at room temperature. In order to detect the
RNA signal, the red reagent working solution was next applied to the tissue sections
and incubated for 10 minutes at room temperature. Once rinsed with deionized
water, the slides were counterstained with 50% hematoxylin, rinsed and dried.
Positive and negative controls were run simultaneously to confirm the assay’s
validity and the corneal samples’ RNA quality and included a PPIB housekeeping gene for a positive assay control and a dabP bacterial gene for a negative control. The tissue sections were evaluated using standard bright field microscopy.

**RNAscope *in situ* hybridization evaluation criteria**

First, all slides from all time submissions were examined utilizing a standard bright field microscopy microscope under 4x, 10x, 20x, and 40x magnification for tissue and cell morphology and for 16ZZ Ec-IL6 target probe signal. The number of IL-6 mRNA transcripts was counted using ACD semi-quantitative scoring recommendation guidelines. Ten slides were randomly selected from each submission time point from each cornea and scored. These ten scored slides were used and those slides used to generate an average score for that particular corneal submission. All slides were scored at 400x. Scores were assigned 0-4 as described previously\(^\text{39}\) in which a 0 was noted if no staining was seen or <1 dot/10 cells was counted, a score of 1 was denoted if 1-3 dots/cell were seen, a 2 was noted if there were 4-9 dots/cell and few clusters were seen, a 3 was marked if 10-15 dots/cell were seen and <10% of those dots were in clusters, and a 4 was assigned if >15 dots/cell were noted and >10% of dots were in clusters (Advanced Cell Diagnostics). All fields were scored by one investigator who was masked to the wounded or unwounded nature of the cornea and the time period of submission.
**Microscopic imaging**

Images were acquired using an Olympus BX53 standard bright field microscope (Olympus, Center Valley, PA, USA) and a DP73 microscope camera.

**RESULTS**

**Corneal ulcer resolution**

All wounded corneas healed within 72 hours and no unwounded corneas experienced any ulceration over the duration of this study. The area of re-epithelialization \( A = \pi r^2 \) via fluorescein stain was documented and averaged over time. The average corneal ulcer areas over time were \( T0=27.7 \pm 6.1 \), \( T24=18.6 \pm 6.3 \), \( T48=7.9 \pm 11.1 \) and \( T72=0 \) mm\(^2\) (Fig. 3-7).

**Modified Hackett-McDonald Ocular Scoring System**

a) Fluorescein intensity (0-4)

All wounded corneas received maximum intensity stain scores at T0 which subsequently decreased until no stain uptake was seen at T72. Specific average intensity scores (\( \pm SD \)) included \( T0=3.9 \pm 0.2 \), \( T24=2.9 \pm 0.2 \), \( T48=1.6 \pm 1.0 \) and \( T72=0 \).

b) Fluorescein location and description

All ulcers were located on the axial cornea and were circular in shape.
c) Corneal edema (0-4)
One wounded cornea experienced a slight loss of corneal transparency (score of 1) and paraxial corneal edema immediately adjacent to the ulceration was noted at T0. By T24 the edema had resolved and no other corneas experienced edema at any time point within this study duration.

Histopathology
All corneas displayed normal corneal architecture. Wounded corneas (Fig. 3-8) showed a lack of epithelium but minimal stromal edema and intact endothelium. Unwounded corneas displayed epithelium of normal thickness, minimal stromal edema and adhered, intact endothelium (Fig. 3-9). The endothelium can be seen in both wounded (Fig. 3-10) and unwounded T72 submissions (Fig. 3-11).

RNAscope in situ molecular detection analysis
The highest expression of IL-6 was seen in the basal epithelial cell layer adjacent to the wound bed in T24 wounded corneas, while the lowest IL-6 expression was seen nearest the limbus. By T72 all wounded corneas had healed and expressed minimal IL-6. Average RNAscope semi-quantitative scoring for each time submission group was calculated and included wounded RNAscope semi-quantitative scoring for each time point, including wounded T24=1.7 ± 0.7, T48=0.9 ± 0.7, T72= 0.2± 0.4 and unwounded T24=0.3 ± 0.5, T48=0 and T72=0 groups (Figs. 3-12- 3-14).
Decreased IL-6 expression was seen in the wounded corneas nearest the limbus (Fig. 3-15). Minimal diffuse IL-6 expression was seen in T24 unwounded corneas while T48 and T72 unwounded corneas expressed no IL-6 mRNA transcripts (Fig. 3-16).

**DISCUSSION**

*Ex vivo* corneal wound healing models are becoming popular in recent years, as they maintain the dynamic, complex, physiological events of the *in vivo* model, while decreasing the need for live animals for this type of studies. In this study, we established an *ex vivo* model of equine corneal wound healing, and demonstrated its physiological relevance. The anatomical integrity of the different corneal layers was maintained throughout the study period. Apart from the ulcerated site, all corneas maintained a normal epithelial cell layer, and exhibited a minimal amount of stromal edema (Fig. 3-10). In addition, a healthy endothelial cell layer and Descemet's membrane were noted in all corneas in our study (Fig. 3-10). It is noteworthy that in other *ex vivo* corneal wound healing models, as well as in our previous experience (not published), the endothelium is usually damaged or missing where it was contacted by the conformer. Some studies do not mention the integrity of their endothelial layer. Due to the unique anatomical features of the equine cornea (i.e. greater diameter and thickness), a conformer was not needed in our model, leading to a healthy, intact endothelium. As a result, minimal corneal stromal edema was present, compared to models utilizing conformers.
All corneal ulcers in our study healed within 72 hours from wounding, consistent with previous publications.\(^1\) Corneal epithelial wounds heal by cellular migration and proliferation. Although the mechanism of corneal wound healing is not fully understood, studies have demonstrated the importance of IL-6 in this complex process.\(^3\) Interleukin-6 is upregulated following corneal injury in humans, as well as in animal and \textit{in vitro} models.\(^{34,36,37,43}\) The increase in IL-6 expression facilitates, and expedites, epithelial wound closure.\(^{36,44,45}\) Fibronectin has been shown to induce corneal epithelial cells migration, a process that is enhanced by IL-6 \textit{in vivo}.\(^46\) In addition, IL-6 reduces the production of matrix metalloproteinase-2, which may have deleterious effect on wounded cornea.\(^47\) Furthermore, the suppression of IL-6 expression has negative effect on corneal epithelial cell migration, both \textit{in vitro} and \textit{in vivo}.\(^43\)

The results of our study demonstrate increased IL-6 expression in injured corneas, in agreement with other studies. Peak IL-6 expression has seen reported at one day, or 24 hours post corneal surgery in human patients,\(^37\) and 7 days after corneal alkali burn in mice\(^34\) and rabbits.\(^33\) The peak IL-6 expression in our study was noted 24 hours after corneal wounding, similar to the finding of Resan \textit{et al}.\(^37\) However, since in our study the first time point for IL-6 expression was 24 hours post injury, we cannot rule out an earlier peak of IL-6 expression. While we and others\(^33,34\) used an alkali burn model, we used equine corneas, while Sotozono \textit{et al} used mouse corneas\(^34\) and Imanishi \textit{et al} used rabbit corneas.\(^33\) The interspecies differences may account for the discrepancy in IL-6 peak expression time in these studies.
The RNAscope in situ hybridization technique was used in our study. This technique is a robust, highly specific and highly sensitive method for analyzing RNA biomarkers, and allows for molecular visualization of gene expression all while maintaining tissue integrity. Utilizing this modality, we showed that IL-6 expression was the greatest adjacent to the wound bed, in the basal epithelial cell layer, similar to the findings of others. Interestingly, Sumioka et al found that transient receptor potential cation channel subfamily V member 1 (TRPV1) is present mainly in the basal layer of mouse and rat corneal epithelium, and that TRPV1 is required for the upregulation of IL-6.

This study presents a novel ex vivo equine corneal wound healing model. In this model, corneal architecture (including corneal endothelium) was maintained, unlike other ex vivo corneal wound healing models. Hence, this model better mimics in vivo events as seen by timely wound healing and IL-6 upregulation. In conclusion, this novel ex vivo corneal wound healing model is physiologically relevant, and may be utilized for future studies of corneal wound healing, while decreasing or negating the use of live, healthy animals for such studies.

CONCLUSIONS

In summation, this study established an air-liquid interface with medium bathing ex vivo model of equine corneal wound healing. Ulcer resolution was noted to be consistent and timely, mimicking the in vivo response of an uncomplicated, superficial ulceration that heals within 72 hours. Histologically, intact epithelium,
minimal stromal edema and presence of the endothelium were seen. Further, IL-6 analysis via RNAscope quantification displayed an upregulated response of this pro-inflammatory cytokine in wounded corneas versus unwounded controls. This finding parallels a previous in vivo rabbit study which demonstrated that the metabolically active basal epithelial layer expressed the greatest amount of IL-6. The expression and elucidation of this important cytokine, coupled with the timely macroscopic ulceration resolution and the maintenance of corneal architecture histologically, validates the physiologic relevance of this model. Therefore, this model can be utilized to reduce or eliminate the use of live horses for corneal wound healing studies, while providing an effective platform for corneal wound healing research in this species.

Conflicts of Interest

The authors have no conflicts to declare.
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**Figures for Chapter 3**

Figure 3-1: Equine globe showing application of the 6 mm filter paper disk soaked in NaOH applied to the axial cornea.

Figure 3-2: Freshly wounded cornea showing circular area of ulceration that is fluorescein positive.

Figure 3-3: Corneoscleral cup excision employing a Bard Parker blade placed approximately 2 mm posterior to the limbus. The anterior portion that contains the cornea (ie “corneal cup”) will then be placed in the incubator cornea side up.
Figure 3-4: Rocker setup including petri dishes with corneas. Note the 6 degree incline that aided in bathing the corneas eight times per minute. The rocker was placed in the CO$_2$ incubator.

Figure 3-5. Representative wounded cornea at T24. Note the axial, circular fluorescein uptake highlighted by cobalt blue light and the millimeter scale bar.

Figure 3-6: Representative wounded cornea at T48. Note the pinpoint axial fluorescein uptake highlighted by cobalt blue light versus the much larger fluorescein positive area shown in the above T24 figure 3-5.
Figure 3-7: Values for the average ulcer area over time including T0=27.7±6.1, T24=18.6±6.3, T48=7.9±11.1 and T72=0 mm².

Figure 3-8. Histopathology of representative T48 wounded cornea stained with hematoxylin and eosin. Note the lack of epithelium (black arrow) superficially and the occasional keratocyte (blue arrow) in the stroma. 400x.
Figure 3-9: Histopathology of representative T48 unwounded cornea stained with hematoxylin and eosin. Note the robust 16 cell layer thick epithelium with the basal cell layer (*), wing cell layer (**) and superficial epithelial layer (***) . Additionally, note the lack of edema within the stromal layer. 400x.

Figure 3-10: Representative wounded corneal submission at T72 displaying intact endothelium in a nonwounded zone. Note the robust epithelial layer, the minimal stromal edema, the thickened Descemet's membrane (abbreviated DM) and the intact endothelial layer (thick blue arrow). 100x.
Figure 3-11. Representative unwounded corneal submission at T72 displaying intact endothelium. Note the robust superficial epithelial layer, the minimal stromal edema, the thickened Descemet’s membrane, and the intact endothelial layer (black arrow). 40x.
Figure 3-12: RNAscope semi-quantitative scoring for each time point including wounded T24=1.7 ± 0.7, T48=0.9 ± 0.7, T72= 0.2± 0.4 and unwounded T24=0.3 ± 0.5, T48=0 and T72=0 groups.

Figure 3-13: RNAscope in situ hybridization analysis of a representative T24 wounded cornea immediately adjacent to the alkaline burn. Note the increased IL-6 expression in the basal epithelial cell layer (*). 400x.

Figure 3-14: RNAscope in situ hybridization analysis of paraxial corneal section of a representative T24 wounded cornea. Note the increased IL-6 expression in the basal epithelial cell layer (*) but that the number of IL6 mRNA transcripts are fewer than the above Figure 3-13 as this image is closer to the limbus. 400x.
Figure 3-15: RNAscope *in situ* hybridization analysis of a representative, perilimbal area of T24 wounded cornea. Note the lack of IL-6 expression in the basal epithelial layer (*) as this is the farthest point from the wound. 400x.

Figure 3-16: RNAscope *in situ* hybridization analysis of a representative, axial area of T72 unwounded cornea. No IL-6 expression is seen in the basal epithelial layer (*). 400x.
CHAPTER 4. GENERAL CONCLUSIONS

Summary

Corneal disease is one of the most significant causes of impaired vision in the horse world-wide. This is likely due to several factors including a prominent globe, the lateral location of the globe within the skull and the equines’ exuberant and head throwing behavior. As such, the horse is predisposed to corneal injury. Although equine corneal disease and the subsequent wound healing cascade is a complex process whose mechanisms are still not fully understood, it is well known that inflammatory cells and their secretory products such as cytokines and growth factors are involved. However, despite the high prevalence of corneal disease and ulceration in the horse, the importance of a clear visual axis in this species, and the complexity of corneal wound healing, research is limited and more heavily devoted to in vivo studies. Limitations of these in vivo studies include ethical concerns, cost, small sample sizes, and large variation amongst subjects. In vitro research may be an overly simplified model that lacks the dynamic, physiologic events of the in vivo model as these cell culture lines are not able to replicate ‘cross talk’ across corneal layers. This novel ex vivo model bridges the gap between in vivo and in vitro research and displayed physiologic relevance via timely ulcer resolution, histologically intact corneal layers and IL-6 expression in wounded versus unwounded controls. As such, this model can be utilized to reduce or eliminate the use of live horses for corneal wound healing studies, while providing an effective platform for corneal wound healing research in this species, including, but not limited
to, investigation of corneal wound healing physiology, and evaluation of different treatment modalities.

**Concluding Remarks**

The *ex vivo* equine corneal wound healing model that was described is not only a novel model but one that has shown physiologic relevance when comparing results to *in vivo* ulcer resolution healing times, maintenance of anatomic structure as seen histologically, and IL-6 expression that mimics previous *in vivo* studies. Hence, this model can be utilized in order to improve our knowledge of equine corneal wound healing while reducing the use of live horses for corneal wound healing studies.