HUMAN ADENOVIRUS TYPE 37 AND THE BALB/C MOUSE: PROGRESS TOWARD A RESTRICTED ADENOVIRUS KERATITIS MODEL (AN AMERICAN OPHTHALMOLOGICAL SOCIETY THESIS)

BY James Chodosh MD

ABSTRACT

Purpose: To establish a mouse model of adenovirus keratitis in order to study innate immune mechanisms in the adenovirus-infected cornea.

Methods: Balb/c 3T3 fibroblasts were inoculated with human adenovirus (HAdV) serotypes 8, 19, or 37 and observed for cytopathic effect. Viral growth titers were performed, and apoptosis was measured by TUNEL assay. Viral and host cytokine gene expression was assessed by RT-PCR in cultured Balb/c 3T3 fibroblasts and in the corneas of virus-injected Balb/c mice. Western blot analysis was performed to detect cell signaling in the virus-infected cornea.

Results: Only HAdV37 induced cytopathic effect in mouse cells. Viral gene expression was limited, and viral replication was not detected. Apoptotic cell death in HAdV37-infected Balb/c cells was evident 48 and 72 hours postinfection (P < .01). MCP-1, IL-6, KC, and IP-10 mRNA levels were increased maximally by 8.4, 9.6, 10.5, and 20.0-fold, respectively, at 30 to 90 minutes after HAdV37 infection. Similar cytokine elevations were observed in the corneas of Balb/c mice 4 hours after stromal injection of HAdV37, when viral gene expression for the viral capsid protein IIIa was not detected. Western blot showed increased phosphorylation of ERK1/2 at 4 and 24 hours after corneal infection.

Conclusions: Despite limited viral gene expression, HAdV37 infection of Balb/c 3T3 fibroblasts results in increased proinflammatory gene expression. A similar pattern of cytokine expression in the corneas of HAdV37-infected Balb/c mice suggests the mouse adenoviral keratitis model may be useful for the study of early innate immune responses in the adenovirus-infected corneal stroma.


INTRODUCTION

Ocular infection by subgroup D adenovirus (HAdV) serotypes 8, 19, or 37 causes epidemic keratoconjunctivitis (EKC), manifest by acute pseudomembranous conjunctivitis, punctate and macro-epithelial corneal erosions, and delayed-onset subepithelial corneal stromal infiltrates. Subepithelial infiltrates, the hallmark of epidemic keratoconjunctivitis, cause photophobia, foreign body sensation, and reduced vision, and may persist and/or recur for months to years. It has been shown that HAdV19 infection of human corneal fibroblasts in vitro induces expression of the potent chemoattractants interleukin-8 (IL-8) and monocyte chemoattractant protein-1 (MCP-1), and that this expression appears to be transcriptionally regulated by an intracellular signaling cascade initiated by viral capsid binding to the cells. 2-5 These data suggest that adenoviral subepithelial corneal infiltrates result when infection of superficial keratocytes induces a specific intracellular signal transduction cascade leading to expression of proinflammatory mediators in the corneal stroma. However, a suitable animal model is lacking in which to confirm these observations. We hypothesize that the mouse cornea, while not intrinsically susceptible to infection with human adenoviruses, can serve as a suitable tissue site with which to study host-derived pathogenic mechanisms in adenovirus keratitis. Our long-term goal is to understand the interplay between adenoviruses and mechanisms of immunity in the human cornea. The specific purpose of this study was to determine whether mouse cells and/or the mouse cornea can be induced to express proinflammatory mediators upon infection with an ocular subgroup D adenovirus.

BACKGROUND AND SIGNIFICANCE

Ocular Adenovirus Infection

Originally isolated in 1953 from human adenoids, the adenovirus is nonenveloped with a linear double-stranded DNA genome. The virus protein capsid forms a regular icosahedron with 20 triangular surfaces composed of 240 hexons, and 12 vertices each composed of a single penton. Each penton contains a penton base and a projecting fiber. DNA restriction enzyme analysis and genome typing has resulted in the classification of six adenovirus species or subgroups (A through F) containing a total of 51 distinct serotypes.

Most adenoviral eye disease presents clinically as one of three classic syndromes: simple follicular conjunctivitis, pharyngoconjunctival fever, or EKC. Serotypes 8, 19, and 37 (subgroup D) are the major etiologic agents of EKC, the only adenoviral syndrome with significant corneal involvement. In EKC, severe pseudomembranous conjunctivitis and punctate epithelial keratitis develop 1 week to 10 days after exposure, followed by multifocal subepithelial (stromal) corneal infiltrates 7 to 10 days thereafter. Subepithelial infiltrates, the sine qua non of EKC, cause photophobia and reduced vision and may persist for months to years. 7 Available evidence suggests that adenovirus infection of corneal cells plays a central role in the corneal manifestations of EKC. Others...
have shown that adenoviruses infect corneal epithelium and keratocytes in vitro and in vivo, suggesting that adenoviral epithelial keratitis represents the clinical manifestation of viral cytopathic effect in the corneal epithelium, and that corneal subepithelial infiltrates in EKC might represent the effects of infection of keratocytes within the stroma.

Neutrophils are the first inflammatory cells in the tears of patients with EKC and the first cells to infiltrate the corneas of experimental animals with adenovirus keratitis. Rare histopathologic specimens removed at surgery from patients with long-lasting subepithelial infiltrates due to EKC showed mononuclear cell infiltrates in the superficial corneal stroma. Adenoviral particles were not apparent by electron microscopy, nor was adenoviral antigen evident by immunofluorescence. These anecdotal data are most consistent with the presence of a persistent chemotactic signal in the superficial corneal stroma long after viral replication has ceased and viral antigen has been consumed. In addition, the stromal location of infiltrates in the EKC-affected cornea suggests the possible participation by infected keratocytes in the inflammatory cascade. We hypothesize that keratocyte-derived leukocyte chemoattractants play a role in the pathogenesis of the stromal keratitis in EKC.

**Human Keratocytes: Biology and Function**

As the primary resident cells of the corneal stroma, keratocytes maintain the cornea in a precisely organized and transparent state. Within the cornea, keratocytes are distinguishable as three distinct populations; those just beneath Bowman’s membrane (subepithelial keratocytes) form a particularly dense cellular network, contain twice as many mitochondria and considerably more heterochromatin than keratocytes in the mid and posterior regions. Keratocytes capably produce mediators of inflammation, including IL-6, G-CSF, MCAF, MDNCF, RANTES, MIG, I-TAC, IP-10, MCP-1, GRO-α, ENA-78, and IL-8, and have been implicated in necrotizing stromal inflammation due to herpes simplex virus and gram-negative bacteria, consistent with a role in stromal keratitis associated with infection by distinctly different pathogens. Proinflammatory gene expression by keratocytes may greatly exceed that of corneal epithelial cells. In this regard, keratocytes are not unlike fibroblasts elsewhere in the body, in that they capably contribute to innate immune responses in tissue substantia propria.

When liberated from the cornea and grown in the absence of serum, keratocytes express keratan sulfate proteoglycan and replicate slowly. Exposure of keratocytes to serum in culture induces a change to a fibroblast phenotype, with increased assembly of stress fibers and focal adhesion complexes, and greater expression of fibronectin, collagen, and heparan sulfate. Injury to overlying corneal epithelium, treatment with TGF-β or wounding induces a myofibroblast phenotype, associated with expression of alpha-smooth muscle actin, and enhances the capacity to contract extracellular matrix. The serum-dependent transformation from keratocyte to myofibroblast involves a phosphotyrosine signal transduction pathway and may be at least partially reversible. Owing to the effect of serum on keratocytes, they are generally referred to as corneal fibroblasts when cultured in serum-containing media. Although the keratocyte-fibroblast-myofibroblast paradigm is instructive, the normally avascular human cornea does contain serum components, albeit less so than tissues with a direct blood supply. In addition, during the acute conjunctivitis in EKC, the superficial corneal stroma is bathed in serum factors present in conjunctival exudates.

It is important to note that two separate laboratories—that of Hendricks and Dana—recently identified at least one and possibly several nonkeratocyte cell populations in the corneal stroma, including CD45+ CD11b+ leukocytes thought to be of the monocye lineage. These cells can be isolated in culture by capture of the nonadherent cells that are typically discarded from keratocyte cultures. How these resident leukocytes might contribute to innate immune mechanisms of inflammation in EKC remains unknown.

**Intracellular Signaling Determines Adenovirus Entry Into Host Cells**

Adenovirus infection of a susceptible cell begins with attachment of the penton fiber knob, the most distal component of the adenovirus capsid, to a specific receptor on the cell surface (Figure 1). Known cellular binding sites for adenovirus fiber knob include the coxsackie-adenovirus receptor (CAR), the α2 domain of MHC class I, sialic acid, and CD46 (membrane cofactor protein). Recent evidence suggests that ocular pathogenic subgroup D adenoviruses bind to sialic acid or CD46. This primary interaction facilitates an essential secondary interaction between Arg-Gly-Asp (RGD) motifs within the viral penton base protein on the proximal surface of the viral capsid and the cellular integrins αvβ3, αvβ5, and αvβ1. This requirement for integrin binding appears to be absolute for all adenoviruses except subgroup F.

Integrins are heterodimeric transmembrane glycoproteins that orchestrate interactions between individual cells and the extracellular matrix and nearest cell neighbors. Formed by the noncovalent association of one of 16 α subunits with one of eight β subunits, integrins rely on interactions with secondary cytoplasmic proteins for the generation of downstream intracellular signaling.

Integrin receptor occupation and clustering by a multivalent ligand lead to rapid formation of focal adhesions within cells (the linking of integrins to intracellular cytoskeletal complexes and associated signaling proteins) and phosphorylation of intracellular tyrosine and serine-threonine kinases.

Binding of integrins by five RGD motifs on the adenovirus penton base optimally promotes integrin clustering in a five-sided ring and initiates a signaling cascade necessary and sufficient for efficient virus internalization (Figure 1). In immortalized epithelial colon carcinoma (SW480) cells infected with HAdV2 (subgroup C), adenovirus internalization required the activation of phosphoinositide 3-kinase (PI3K), and Rho family GTPases. PI3K is a lipid kinase that catalyzes phosphorylation of phospholipid second messengers, that in turn activate the small GTP-binding proteins Ras, Rho, Rac, and CDC42. Activated Rac and CDC42 induce actin polymerization, and the end result is dynamin-dependent viral internalization into endocytic vesicles via clathrin-coated pits. Viral internalization, transport, and uncoating into the nucleus may occur within 2 hours of attachment to the cell. The adenovirus replicative cycle that follows is divided into early (E) and late (L) phases with the late phase commencing with onset of protein.
viral DNA replication. The early phase of adenovirus gene transcription has been further subdivided into “immediate early” (E1A), “delayed early” (E1B, E2A, E2B, E3, and E4), and “intermediate” (IVa2, IX) transcripts. In summary, adenovirus infection of target cells is an active cell-mediated process that requires intracellular signaling.

**FIGURE 1**

General schematic of adenovirus internalization cascade, based on studies performed with HAdV2 and SW480 cells. After binding of the HAdV2 penton fiber knob to one of several possible primary receptors, a secondary interaction between the viral penton base and cellular integrins mediates activation of phosphoinositide 3-kinase (PI3K) and Rho GTPases. Subsequent dynamin-dependent actin polymerization induces clathrin-mediated endocytosis of viral particles.

**Interleukin-8 and Monocyte Chemoattractant Protein-1: Paradigm Chemokines and Their Regulation**

Chemokines are 8 to 10 kilodalton (k), basic, heparin-binding peptides with a four-cysteine motif that cause leukocyte chemotaxis with a high degree of specificity for leukocyte cell type. Prior studies have suggested that the production of chemokines by adenovirus-infected tissues may be critical to the subsequent immunopathology associated with infection. The α chemokines, known as CXC chemokines, contain one amino acid between the first and second cysteine, whereas the β or CC chemokines have adjacent cysteines. Interleukin-8 (IL-8, also known as CXCL8) is an α chemokine that strongly and selectively induces chemotaxis and granulocytes with an exceptionally long duration of action, and to a lesser degree instigates chemotaxis of T lymphocytes. Monocyte chemoattractant protein-1 (MCP-1, also known as CCL2), is a β chemokine that induces chemotaxis of monocytes, basophils, CD4+ and CD8+ lymphocytes, and T lymphocytes of the activated memory subset.

Lipopolysaccharide treatment of keratocytes induces expression of both IL-8 and MCP-1. Molecular regulation of IL-8 expression occurs largely at the transcriptional level via activation of latent transcription factors that bind to two distinct promoter regions in the 5′-flanking region of the IL-8 gene: a proximal region with adjacent binding sites for NF-κB and NF-IL6, and a distal binding site for AP-1. The MCP-1 promoter contains two binding sites for NF-κB, and sites for AP-1 and Sp1 but not NF-IL6, suggesting the potential for the differential regulation of these two chemokines. In viral infections, IL-8 expression may occur due to secondary oxidative stress, early viral gene product activation of host gene transcription, or protein kinase signaling cascades initiated by viral binding. Early viral gene products can and do influence host gene expression during infection, but once host cell machinery shuts down, viral gene products cannot possibly influence transcription or translation of host genes.

Recent efforts toward understanding the pathogenesis of adenovirus keratitis have addressed the effects of cell signaling in HAdV19-infected human corneal fibroblasts and are summarized in Figure 2. Increased IL-8 expression upon HAdV19 infection of human corneal fibroblasts was shown to require host intracellular signaling but not viral gene or TNFα/IL-1β expression. Increased MCP-1 mRNA and protein expression in HAdV19-infected human corneal fibroblasts, like IL-8, was shown to be dependent upon intracellular signaling. These published findings using HAdV19 in human corneal fibroblasts were similar to those of Bruder and Kovess-Minszki, who showed that attachment of replication-defective HAdV5 (subgroup C) to HeLa cells led to phosphorylation of Raf-1 and ERK2 and downstream activation of IL-8 gene expression, suggesting a potential mechanism for the inflammation associated with adenoviral gene therapy. Antiviral treatments targeted to viral replication and not viral binding to target cells, would not be expected to block chemokine expression due to viral attachment, suggesting that classic antiviral drugs would not prevent leukocyte infiltration into the cornea once epithelial infection is underway.
Chemokines can be extraordinarily resistant to degradation once bound by negatively charged proteins within extracellular matrix.\textsuperscript{101} For example, IL-8 is extremely stable in tissues due to its propensity to bind to glycosaminoglycans.\textsuperscript{102-105} When expressed into the cornea, IL-8 and MCP-1 likely bind to extracellular matrix components such as heparan sulfate at the corneal epithelial basement membrane to provide a degradation-resistant chemotactic signal that persists to some degree regardless of later events in infection.

**FIGURE 2**
Model of cell signaling and downstream effects in HAdV19-infected human corneal fibroblasts.\textsuperscript{2-5,98,99} Following primary and secondary binding to CD46 or sialic acid, and then \( \alpha_v\beta_3 \) integrin, respectively, the subgroup D adenovirus HAdV19 is internalized by the activity of Src kinase, leading to multiple downstream signaling events, and culminating in enhanced cell survival and proinflammatory gene expression.

**Animal Models of Ocular Adenovirus Infection**
Adenoviruses classically demonstrate a restricted host range; with few exceptions, human adenoviruses replicate poorly or not at all in nonhuman animals. Limited ocular infections with human isolates have been induced experimentally in the cotton rat\textsuperscript{106} and the rabbit.\textsuperscript{9,14,17,107-109} The cotton rat model was successfully utilized for the study of topical antivirals,\textsuperscript{110} but was abandoned as a pathogenesis model for technical reasons. The rabbit model of ocular adenovirus infection has been used extensively for studies of antiviral therapy.\textsuperscript{111-116} However, group D adenoviruses do not replicate in rabbits, and infection with group D adenoviruses does not induce keratitis.\textsuperscript{108} Furthermore, the relative lack of species-specific reagents for immunologic studies in the cotton rat and rabbit models has made them impractical for study of immune responses to infection.

In contrast to other potential animals, the mouse has emerged as a favored model in which to study host-pathogen interactions.\textsuperscript{117} Mouse cells are not susceptible to productive infection by human adenoviruses, but early viral gene expression was shown in human adenovirus-infected mouse 3T3 fibroblasts.\textsuperscript{118} In these studies, the E1B 21k gene product, necessary for efficient viral DNA synthesis and protection of newly synthesized viral DNA against cell nucleases,\textsuperscript{119-121} was not expressed, nor did viral replication occur. Despite such limitations, mouse models of several human adenovirus infections have been developed to study innate immune responses to the virus.\textsuperscript{122-128} For example, infection of the mouse respiratory tract with human adenoviruses results in pneumonia,\textsuperscript{129-132} despite limited viral gene expression.\textsuperscript{133} In a fashion, the absence of adenoviral replication in the mouse has allowed investigators to analyze innate immune responses without the confounding influence of viral replication. Therefore, those wishing to study host-pathogen interactions in mice with human adenoviruses are either limited to or graced with (depending on one’s perspective) experimental adenoviral pathogenesis models without the full repertoire of adenoviral gene and protein expression. This thesis will explore the consequences of infection of mouse cells by human adenoviruses and provide preliminary evidence for the feasibility of a mouse model for adenovirus keratitis.

**METHODS**

**VIRUSES AND CELLS**
Human adenovirus (HAdV) serotypes 8 and 37 were obtained from the American Type Culture Collection (ATCC, Manassas, Virginia) and grown in A549 cells (ATCC CCL 185, human lung carcinoma cells) in Minimum Essential Medium (MEM, Invitrogen, Carlsbad, California) with heat-inactivated 2% fetal bovine serum (FBS) and antibiotics. Four banked clinical EKC isolates of HAdV19 were used and cultured in similar fashion. After growth in A549 cells from the ATCC (American Type Culture Collection, Manassas, Virginia), viruses were cesium chloride gradient-purified and dialyzed against 10 mM Tris (pH 8.0) buffer with 80 mM NaCl, 2 mM MgCl\textsubscript{2}, and 10% glycerol, titered in triplicate by the Tissue Culture Infectious Dose (TCID) assay, and stored at \(-80^\circ\)C. Balb/c 3T3 cells (clone A31- mouse embryo) were obtained from ATCC and cultured and maintained in Dulbecco’s Modified Eagle’s Minimum Essential Medium (DMEM) supplemented with heat-inactivated 10% FBS and antibiotics.
REAGENTS
SYBR green master mix for real-time PCR was purchased from Applied Biosystems, (Foster City, California). Other PCR reagents, including Taq polymerase, oligo dT (15mer) primers, and recombinant RNasin inhibitor, were obtained from Promega (Madison, Wisconsin). The ApoTag peroxidase in situ apoptosis detection kit was obtained from Serological (Norcross, Georgia). Horseradish peroxidase-conjugated donkey anti-rabbit IgG and chemiluminescent reagents were obtained from Amersham (Piscataway, New Jersey). The polyclonal anti-phospho-ERK 1/2 antibody was obtained from Cell Signaling (Beverly, Massachusetts) and the polyclonal anti-total ERK 1/2 antibody was obtained from Promega.

IN VITRO VIRAL INFECTION
Balb/c 3T3 monolayer cultures were grown to ~ 95% confluence in six well culture plates, washed thrice with MEM 2% FBS, and infected with adenoviruses at multiplicities of infection (MOI) of 10 and 50 TCID/cell or mock infected with virus-free dialysis buffer. After 1 hour incubation at 37°C, the cells were washed gently, fed with MEM 2% FBS, and reincubated. Cell cultures were then observed for up to 28 days for the development of adenoviral cytopathic effect.

For growth titer experiments, 3T3 cells were grown in 48 well plates and infected or mock infected in triplicate for each time point. The cells and supernatants were harvested each day for 6 subsequent days postinfection, and titered. For viral and host gene expression experiments, RNA was harvested from mock and virus-infected cells as described below, at the indicated time points.

CORNEAL INJECTIONS
Balb/cJ mice aged 6 to 8 weeks were purchased from Jackson Labs (Bar Harbor, Maine) and maintained in strict accordance with institutional animal care facility guidelines and the ARVO Statement on the Use of Laboratory Animals in Ophthalmic Research. The animal protocol was approved by the University of Oklahoma Health Sciences Center Institutional Animal Care and Use Committee. Anesthesia was provided by intraperitoneal ketamine (80 mg/kg, Phoenix Scientific, St Joseph, Missouri) and xylazine (5 mg/kg, Bayer, Shawnee Mission, Kansas), followed by ocular topical application of proparacaine (0.5%, Allergan, Hormigueros, Puerto Rico).

The right eye was gently prolapsed with curved forceps, and corneal injections were administered under a Zeiss OPMI VISU 140 Surgical Microscope (Thornwood, New York) with digital imaging system, using sterile glass needles heat-pulled from micropipettes, (Needle/Pipette Puller, Model 730, David Kopf Instruments, Tujunga, California, and KT Brown Type Micro-Pipette Beveler, Model BV-10, Sutter Instrument, Novato, California), and a CO2-powered microinjection system (PM2000 Cell Microinjector, MicroData Instruments, Plainfield, New Jersey). Either 1 μL of the dialysis buffer used to dialyze the purified virus (mock-infected control) or an equal volume of cesium chloride gradient-purified HAdV37 was injected into the right cornea, with the left eye left untouched in all mice. The injection needle was placed in the paracentral stroma of each cornea, and transient whitening of the corneal stroma was used to indicate a successful injection. Confirmation of the injection site was obtained by analysis of injected corneas with light and confocal microscopy (data not shown). After euthanasia of mice at selected time points post-infection, corneas were removed using Vannas scissors (Katena, Denville, New Jersey), and placed into RNAlater (Ambion, Austin, Texas) for PCR analysis, or chilled PBS (pH 7.4) for protein studies.

RNA ISOLATION
For RNA isolation, the corneas were lysed in 1 mL of TRIZol reagent (Invitrogen, Carlsbad, California) using RNase free, disposable, Pellet Pestle (Kimble/Kontes, Vineland, New Jersey), and RNA was isolated as per the manufacturer’s protocol. Proteins were removed by a chloroform extraction of the lysate. RNA was precipitated from the supernatant with ethanol, and the pellet was resuspended in Tris-EDTA (pH 8.0). RNasin (Promega) was added to the RNA solution to prevent RNase action. Contaminating DNA was removed by DNase I (Promega) treatment followed by a phenol/chloroform extraction and subsequent ethanol precipitation of the RNA. The RNA was resuspended at a concentration of 1 mg/mL in DEPC-treated water. A spectrophotometric reading at a wavelength of 260 nm was used to determine the concentration of RNA. The quality of each RNA sample was determined by calculating the ratio of optical density of each RNA sample at 260:280 nm; a ratio of approximately 1.8 indicated that samples contained only nondegraded RNA.

REVERSE TRANSCRIPTASE POLYMERASE CHAIN REACTION
For synthesis of cDNAs, 5 μg of total RNA was reverse transcribed with Moloney murine leukemia virus reverse transcriptase (Promega) using an oligo-dT 15mer (Promega) as the primer. The reaction mixture for the reverse transcription reaction was composed of 1.5 U/μL of RNasin, 50 mM Tris-HCl (pH 8.3), 75 mM KCl, 3 mM MgCl2, 10 mM DTT, 500 μM dNTPs and 10 U/μL of reverse transcriptase. A reaction without reverse transcription, which was composed of all of the above reactants except for the reverse transcriptase, was run with each experiment to rule out the possibility of amplification of contaminating genomic DNA in the PCR reaction step.

For reverse transcriptase PCR (RT-PCR) amplification of adenoviral mRNAs, optimal primers (Tm about 45°C to 55°C; GC content range 55% to 60%; primer length 14 to 24mer) were designed using the Primer Macintosh software and verified for interprimer and intraprimer interactions and self-dimerization using the Primer3 and Integrated DNA Technology (Coralville, Iowa) calculator software. Primers for amplification of viral genes were designed using known sequences of human adenoviruses, as shown in Table 1. Two μL of the cDNA obtained by reverse transcription were used with the PCR reaction mixture composed of 50 mM Tris-HCl (pH 9.0), 50 mM NaCl, 10 mM MgCl2, 200 μM dNTPs, 20 μg/μL primers and 1 U of Taq polymerase (Promega). Thin-
walled PCR reaction tubes were used for the reactions, and the assay was performed on a programmable thermo-minicycler (MJ Research, Waltham, Massachusetts) using one cycle that comprised a denaturation step at 96°C for 2 minutes followed by 30 cycles of 96°C for 1 minute and 68°C for 2 minutes. The final extension step was carried out at 72°C for 5 minutes. The amplification products were analyzed by gel electrophoresis in 1% agarose gels, and the sizes of the amplicons were verified by comparing them to the 100 bp DNA marker (GIBCO, Carlsbad, California).

### TABLE 1. PRIMER PAIRS FOR RT-PCR DETECTION OF HADV37 MRNAS

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RT-PCR = reverse transcriptase polymerase chain reaction.

*Genbank accession numbers: a. AF099665; b. AF086569; c. SB2508; d. AF099665; e. NC 001454; f. NC 001405.

Quantitative real-time PCR analysis of viral and host gene expression in HAdV37 and buffer-injected corneas was performed using the RNA pooled from three corneas in each treatment group, using the ABI Prism 7700 Sequence detection System (PE Applied Biosystems, Foster City, California) according to the manufacturer’s instructions. Primers for real-time PCR of viral and mouse mRNAs were designed using Primer Express Software (PE Applied Biosystems) and are shown in Tables 2 and 3, respectively. RNA concentrations of samples were normalized using quantification of GAPDH mRNA. Three μL of cDNA were subjected to real-time PCR amplification in a final volume of 50 μL containing 25 μL of 2X SYBR green master mix and 250 nM of specific forward and reverse primers. Amplification curves were generated by monitoring the fluorescence of SYBR Green I as a measure of incorporation into the amplified product. Samples were then analyzed by comparison of the number of PCR cycles required to reach the midpoint of
each amplification curve, or threshold cycle (\(C_T\)). Comparison of gene expression between two samples was performed after GAPDH normalization by calculating the \(n\)-fold difference in mRNA abundance using the formula \(y = 2^{-x}\), where \(x = (C_T\) of sample 1 – \(C_T\) of sample 2) and \(y\) = (\(n\)-fold difference in mRNA abundance). For each gene, a range of concentrations for both the forward and reverse primers allowed us to determine the combination with optimum amplification. Reactions lacking template were used to control for primer-dimer formation. To control for contamination by residual genomic DNA, reverse transcriptase-negative and nontemplate controls were run in parallel with each experiment.

<table>
<thead>
<tr>
<th>VIRAL GENE*</th>
<th>SOURCE (HADV)</th>
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<th>SEQUENCES</th>
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</tbody>
</table>

PCR = polymerase chain reaction.

*Genbank accession numbers: E1A 10S and E1B 19K: AF099665; IIIa: AF108105.

**CYTOKINE QUANTIFICATION**

For the quantification of cytokine proteins, corneas harvested at the specified times postinfection were dissected into 150 \(\mu\)L of chilled 1% SDS lysis buffer with protease inhibitors (10 \(\mu\)g/mL leupeptin, 1 mM phenylmethylsulphonylfluoride, and 10 \(\mu\)g/mL aprotinin), lysed using pellet pestle, and further homogenized by cold sonication (Sonic Dismembrator, Fisher, Pittsburgh, Pennsylvania). Supernatants were collected after centrifugation and the pooled proteins from three corneas for each condition and time point were analyzed by the Bio-Plex cytokine assay system (BioRad, Hercules, California) per the manufacturer’s instructions.

**DETECTION OF APOPTOSIS**

Balb/c 3T3 cells were infected with purified HAdV37 or mock infected with virus free buffer, and then fixed at 12, 24, 48, or 72 hours postinfection with 1% paraformaldehyde. Terminal deoxynucleotide transferase dUTP nick end labeling (TUNEL) analysis for DNA fragmentation was carried out using the TUNEL Apoptosis Detection Kit (Upstate Biotechnology, Lake Placid, New York), following the manufacturer’s instructions. Each experimental condition was duplicated in three wells, and the number of apoptotic cells in 10 high-power fields in each well was counted in masked fashion and averaged. The means for each experimental condition were compared by ANOVA with Scheffe’s multiple comparison test.

**SDS-PAGE AND IMMUNOBLOT ANALYSIS**

For preparation of protein lysates, untouched, buffer-, or HAdV37-injected corneas were homogenized in 250 \(\mu\)L of chilled lysis buffer consisting of phosphate buffered saline (PBS), 1% Triton X-100 and 2 mM EDTA, 0.2 mM sodium orthovanadate, along with protease inhibitors including phenylmethylsulphonyl fluoride (1 mM), leupeptin (1 \(\mu\)g/mL) and aprotinin (10 \(\mu\)g/mL), and freeze thawed once in liquid nitrogen. Samples were sonicated on ice for 5 to 10 seconds, followed by centrifugation at 13000 \(\times\) g for 10 minutes at 4°C. Thirty micrograms of total protein from each supernatant was boiled in 2X sample buffer (BioRad 2X Laemmli buffer: 62.5 mM Tris HCl [pH 6.8], 2% SDS, 25% glycerol, 0.01% Bromophenol Blue, and 5% B-Mercaptoethanol) and immediately loaded on 10% SDS-PAGE. The resulting gels were transferred to nitrocellulose membranes using a BioRad Mini-Protein II transblot apparatus.

Nitrocellulose membranes were blocked overnight at 4°C in 4% bovine serum albumin. Incubation with primary antiserum was performed for 2 hours at room temperature. Immunoblots were washed thrice with Tris-buffered saline after both the primary and secondary incubations. Antibody reactivity was determined with enhanced chemiluminescent reagents (Amersham Biosciences, Piscataway, New Jersey), using the appropriate peroxidase conjugated secondary antibody.

**REPETITION OF EXPERIMENTS AND STATISTICAL ANALYSIS**

All experiments were performed at least three times. For statistical analysis when appropriate, the means from each experimental condition were compared by ANOVA with Scheffe’s multiple comparison test, using SAS statistical software (Cary, North Carolina). \(P < .01\) was considered significant.
HADV37 INDUCES CYTOPATHIC EFFECT IN MOUSE CELLS

At the onset of these studies, we sought to determine if a human adenovirus from subgroup D could be identified that would infect mouse cells in culture with the idea that such a virus would be the most likely to establish infection in the living mouse cornea. Cultured Balb/c 3T3 fibroblasts were seeded with HAdV8, 19 (four clinical isolates), or 37, at two different MOI (10 and 50 TCID/cell). Only HAdV37 infection at an MOI of 50 induced cytopathic effect, evident after 2 days and extensive at 3 days postinfection (Figure 3). Similar findings were seen in CRL cells, a Balb/c epithelioid cell line (ATCC) and Du17 cells, a C57BL/6J fibroblast cell line (provided courtesy of Dr Dusko Illic, data not shown). None of the other viruses tested induced cytopathic effect at these MOIs.

In light of the induction of apparent viral cytopathic effect in mouse cells upon infection with HAdV37, we then sought to determine whether the observed cell death was due to viral replication. Balb/c 3T3 cells were infected in triplicate with HAdV37 or buffer, and the cells and supernatants harvested each day for 6 subsequent days, and titered. The titer of HAdV37 fell steadily each day, from 57.0 TCID/cell immediately after infection to 1.5 TCID/cell on the sixth day postinfection (Figure 4), consistent with a lack of viral replication. Adenoviral capsids devoid of viral DNA can be produced in some adenovirus infections, and the empty capsids are toxic and induce cytopathic effect. To test this alternate explanation for the viral cell death induced by HAdV37 in Balb/c 3T3 cells, we performed Western blot analysis comparing Balb/c 3T3 cultures after the development of cytopathic effect with those immediately after viral adsorption, but could not detect any increase in capsid protein (data not shown).

It has been suggested that the inability of human adenoviruses to replicate in mouse cells may be due to a failure to synthesize specific viral gene products. We infected (human) A549 cells and Balb/c 3T3 fibroblasts in parallel, and performed RT-PCR at various times postinfection to detect expression of specific adenoviral genes. Although only a few regions of HAdV37 have been sequenced, we were able to use other published adenovirus sequences to develop primer pairs that would detect mRNA for HAdV37 genes in A549 cells (Table 1). Surprisingly, we found that HAdV37 gene expression did indeed take place in Balb/c 3T3 fibroblasts, although it typically occurred days rather than hours post-infection (Figure 5). For example, PCR products for the three E1A transcript sedimentation products examined (10s, 12s, and 13s) were evident at 2, 2, and 4 hours postinfection in A549 cells, but not until 48, 96, and 168 hours, respectively, in Balb/c cells. The same gene products are typically evident within 4 hours postinfection in human corneal fibroblasts (unpublished data). Furthermore, we could not detect some viral mRNAs at any time point in the mouse cells, including E3B 14.7k, E4, L1 52-55k, and L2 III. Interestingly, the 14.7k product of the E3B gene, with mRNA evident at 4 hours postinfection in A549 cells, protects adenovirus-infected cells against cytolyis by tumor necrosis factor (TNF). Similarly, the failure of L1 52-55k gene to express in mouse cells—the transcript was noted in A549 cells at 6 hours postinfection—was also
noteworthy, because this gene product is essential to viral DNA encapsidation. In contrast to our results with HAdV37, after infection with HAdV19 we could not detect any of the E1A sedimentation products in Balb/c 3T3 cells (data not shown), consistent with the observed failure to develop cytopathic effect after HAdV19 infection of mouse cells. Our data are consistent with internalization of HAdV37 into Balb/c 3T3 fibroblasts, and subsequent viral gene expression, albeit abortive. Whether HAdV19 or other noncytopathic HAdVs can be internalized in mouse cells but then fail to express early viral genes is not known.

FIGURE 4
Growth curve for HAdV37 in Balb/c 3T3 fibroblasts. Cell monolayers at 95% confluence were infected in triplicate with HAdV37 at a multiplicity of infection of 50 TCID/cell, and cells and supernatants harvested daily for 6 days postinfection for titering. Error bars show the standard deviation of the mean titer at each time point.

FIGURE 5
Expression of specific HAdV37 mRNAs by reverse transcriptase polymerase chain reaction (RT-PCR) in Balb/c 3T3 cells (○) vs A549 cells (■). Cell monolayers at 95% confluence were infected in triplicate with HAdV37 at a multiplicity of infection (MOI) of 50 TCID/cell, and RNA harvested as described in the “Methods” section, and subjected to RT-PCR with primers listed in Table 1. The time of earliest detection of each adenoviral transcript in at least three independent experiments is shown for each cell type. Those genes with an asterisk (*) were found to be expressed only in the human A549 cells, not in the mouse cells.

HADV37 INDUCES APOPTOTIC CELL DEATH IN MOUSE CELLS
To test the hypothesis that the cytopathic effect seen in mouse cells upon HAdV37 infection was due to apoptosis, Balb/c 3T3 fibroblasts were infected with HAdV37 or mock-infected with buffer in triplicate and fixed for TUNEL staining at 12 to 72 hours postinfection. As seen in Figure 6, at 48 and 72 hours postinfection, there was a significantly greater number of apoptotic cells in HAdV37-infected cell cultures than in mock-infected cultures (number of apoptotic cells/high-powered field for virus vs mock infection: 61.0 ± 7.3 vs 2.7 ± 0.9, and 44.7 ± 3.3 vs 8.7 ± 5.8, at 48 and 72 hours postinfection, respectively; \( P < .01 \) for 48- and 72-hour comparisons, ANOVA with Scheffe’s multiple comparison test). We quantified the proportion of apoptotic cells per total cells, comparing mock- vs HAdV37-infected cells at these two time points. At 48 and 72 hours postinfection, the relative proportions were 1.3% vs 44.8%, and 11.8% vs 54.2%, respectively; similar findings were obtained in Du17 cells (data not shown).
FIGURE 6
TUNEL-positive Balb/c 3T3 cells per high-powered field (hpf) at indicated times postinfection. Balb/c 3T3 fibroblasts were infected with buffer or HAdV37 at a multiplicity of infection of 50 TCID/cell. Significantly increased numbers of apoptotic cells were noted in HAdV37-infected cell cultures at 48 and 72 hours after infection, as compared to cells mock-infected with buffer (* P < .01). The experiment shown is representative of three independent experiments. Error bars show the standard deviation of the mean number of apoptotic cells at each time point.

HADV37 INDUCES PROINFLAMMATORY CYTOKINE EXPRESSION IN MOUSE CELLS

To determine whether HAdV37 infection of mouse cells was associated with induction of cytokine gene expression, real-time PCR was performed on HAdV37- and mock-infected Balb/c 3T3 cells. Cytokine mRNA levels were normalized by GAPDH, and the fold-difference between virus and mock-infected cells was determined as described in the “Methods” section, using the primers shown in Table 3. We chose four cytokines to examine, based on their possible importance in adenovirus keratitis. Interleukin-6 (IL-6) was studied because of its consistent induction in adenovirus infection in varied models.130,137,138 The neutrophil chemokine and human IL-8 homolog, KC, was chosen because of evidence for IL-8 induction by adenovirus infection of human corneal cells,2,4 and its similarity in function and regulation to human IL-8.138-141 MCP-1 was selected because of studies that show it is upregulated in human corneal fibroblasts after HAdV19 infection.53 IP-10, a CXC chemokine that attracts CD4+ T lymphocytes and has been shown to mediate a broad array of functions in viral infections,142,143 was studied because of existing evidence for its role in inflammation associated with experimental adenovirus infections.144-149 By real-time PCR, HAdV37 induced increased mRNA levels for all four cytokines at 30 minutes postinfection. Figure 7 shows one representative experiment of a total of three performed. While levels of KC and MCP-1 appear to have already peaked at or before 30 minutes postinfection with maximal fold-increases of mock-infected cells of 10.5 and 8.4, respectively, at 30 minutes, IL-6 and IP-10 mRNA levels appear to be increasing through the first 90 minutes, with fold-increases of 9.6 and 20.0, respectively, at 90 minutes postinfection.

<p>| TABLE 3. PRIMER PAIRS FOR REAL-TIME PCR DETECTION OF HOST MRNAS |
|------------------|------------------|------------------|------------------|</p>
<table>
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PCR = polymerase chain reaction.
*Genbank accession numbers: GAPDH: BC083080; KC: U20527; IL-6: NM_031168; MCP-1: AF065931; IP-10: M33266.
Real-time polymerase chain reaction (PCR) analysis of cytokine mRNA change induced in vitro by HAdV37 infection of Balb/c 3T3 fibroblasts. Cell monolayers were infected at a multiplicity of infection of 50 TCID/cell with HAdV37 or mock-infected with buffer and the RNA harvested at 30, 60, and 90 minutes postinfection. The height of each bar indicates the fold-increase in host cell cytokine gene expression induced at the indicated time after HAdV37 infection, as compared to mock-infected cells. Fold-increases in KC and MCP-1 mRNA expression appear maximal at 30 minutes postinfection, whereas fold-increases in IL-6 and IP-10 mRNA expression appear to be highest at 90 minutes postinfection. This experiment was repeated twice with similar results.

HADV37 INDUCES PROINFLAMMATORY CYTOKINE EXPRESSION IN THE BALB/C MOUSE CORNEA

Our in vitro data performed with cultured mouse cells strongly suggested the possibility that HAdV37, if properly applied to the mouse cornea, might induce inflammatory cytokines and thus allow the study of inflammatory gene expression in a mouse model of infection, even in the absence of viral replication. Our in vitro data was also consistent with other published studies showing that adenoviruses induce inflammation in living mice without viral replication.\(^{126-133}\) We subsequently found that topical corneal application of HAdV37 to Balb/c mice without corneal scarification failed to induce changes in host gene expression, whereas infection with corneal scarification created too much of a local inflammatory response in controls to allow meaningful comparisons between groups at early time points (data not shown). When HAdV37 was injected into the cornea with a 33-gauge needle and Hamilton syringe, cytokine mRNA levels rose in both mock and HAdV37 injected mouse corneas in the first 4 to 8 hours, presumably due to the trauma of injection (data not shown). Finally, a less traumatic injection technique using sterile glass needles heat-pulled from micropipettes (see “Methods” section), and a CO\(_2\)-powered microinjection system was developed that induced only minimal alterations in cytokine gene expression in controls, thus allowing us to perform our analyses. The right corneas of anesthetized mice were injected in the midcorneal stroma with either 1 \(\mu\)L of the dialysis buffer used to dialyze the purified virus (mock-infected control) or with an equal volume of 10\(^8\) TCID/mL, cesium chloride gradient-purified, HAdV37, and kept the left corneas as untouched controls. A successful corneal stomal injection was evident by transient whitening of the corneal stroma and confirmed by subsequent analysis of injected corneas with light and confocal microscopy (data not shown). Mice were euthanized at 4 hours postinjection, and three corneas in each treatment group were harvested and pooled for RNA extraction and subsequent real-time PCR for IL-6, KC, and IP-10. Fold-change in cytokine mRNA expression was determined in comparison to the quantity of mRNA for the same cytokine in untouched control corneas. At 4 hours postinfection, marked increases were noted in cytokine expression for all three cytokines (Figure 8), consistent with similar elevations at earlier time points in vitro for a panel of mediators similar to those elevated in HAdV37-infected mouse cells in vitro. Levels of fold-increase were 13.7, 25.6, and 12.9 for IL-6, KC, and IP-10, respectively, with a similar pattern of increase seen in two additional experiments. The small amount of RNA present in each mouse cornea allowed us to test only three cytokines at any one time, and MCP-1 was omitted from these analyses.

Because of the importance attributed to neutrophils in the acute inflammatory response to adenovirus infection,\(^2\) we sought in particular to confirm the real-time PCR data for the neutrophil chemoattractant KC. For the quantification of KC protein, three corneas each were harvested at 0, 4, and 8 hours postinfection, and protein was extracted and pooled for analysis by the Bio-Plex cytokine assay (see “Methods” section). We found that changes in KC mRNA were mirrored by increases in KC protein expression (Figure 9) and that the level of KC in HAdV37-injected corneas appeared greatest at 8 hours postinjection. The amount of KC detected was 235.0 pg per virus-infected cornea vs. 8.0 pg per mock-infected cornea at 8 hours postinfection, an approximately 30-fold increase. A smaller elevation of IL-6 (13-fold) protein was detected after 8 hours (data not shown); MCP-1 and IP-10 proteins were not evaluated. The KC experiment was performed three times with similar results.
Real-time PCR analysis of change in host cytokine gene expression in Balb/c mice corneas at 4 hours after HAdV37 injection as compared to untouched and buffer-injected control corneas. The RNA from three corneas was pooled for analysis of each cytokine at each time point. The level of mRNA for each cytokine in buffer and HAdV37-injected corneas was expressed as the fold-increase over that of untouched corneas, set arbitrarily at a value of 1.0. mRNA levels for IL-6, KC, and IP-10 were all increased at 4 hours postinfection. A similar pattern of increased expression for all three cytokines was seen in two replicates of this experiment.

Analysis of KC protein expression in corneas of Balb/c mice at 0, 4, and 8 hours postinfection. Right corneas were injected with $10^8$ TCID/ml of HAdV37 or buffer in 1 μL, and the left corneas used as untouched controls. Corneas were harvested at the specified times postinfection and dissected into chilled 1% SDS lysis buffer with protease inhibitors, lysed, and homogenized. Supernatants were collected after centrifugation, and KC protein quantity in each sample was analyzed by the Bio-Plex cytokine assay system and reported as the quantity of KC protein per cornea. KC levels were highest in HAdV37-injected corneas at 8 hours postinfection. This experiment was performed three times with equivalent results.

By real-time PCR performed on corneas at 4 hours postinjection, we also identified mRNA expression for E1A and E1B 19k HAdV37 genes but not IIIa (Figure 10), a late gene encoding a viral capsid protein, consistent with only early viral gene expression at a time when host proinflammatory gene expression is already upregulated. Because prior studies had demonstrated that host cytokine gene expression in HAdV19-infected human corneal fibroblasts was controlled by intracellular kinase activation events following viral binding to the cells, we sought to determine whether cell signaling could be identified in the HAdV37-injected mouse cornea. Protein was harvested from corneas at 4 and 24 hours after injection of buffer or HAdV37 and subjected to Western blot for tyrosine phosphorylated and total ERK1/2, a tyrosine kinase shown previously to control induction of IL-8 gene expression in HAdV19-
infected corneal fibroblasts. As shown in Figure 11, increased phosphorylation of ERK1/2 was seen at both time points in HAdV37-injected corneas as compared to buffer-injected corneas, consistent with enhanced tyrosine phosphorylation in virus-injected Balb/c corneas.

DISCUSSION

Ocular infection by the subgroup D HAdV8, 19, or 37 has long been associated with EKC. Since Jones first suggested almost 50 years ago that subepithelial stromal infiltrates in EKC might follow adenovirus infection of corneal epithelium, secondary viral antigen deposition in the superficial corneal stroma, and lymphocyte infiltration in a multifocal pattern at sites of antigen-antibody complexes, remarkably little progress has been made with regard to understanding the molecular pathogenesis of adenovirus keratitis. Experimental evidence for Jones’ theory is still lacking, and we now know that antigen-antibody reactions in the cornea typically present as Wessely rings.

Trousdale and coworkers’ more recent finding that a nonreplicating adenovirus could induce subepithelial infiltrates when injected into the rabbit corneal stroma showed that viral replication in the stroma was unnecessary for the development of stromal keratitis in EKC, but did not elucidate a specific molecular mechanism of inflammation. In an attempt to explain corneal stromal inflammation in

FIGURE 10
Real-time polymerase chain reaction (PCR) analysis for expression of adenoviral gene transcripts in corneas of Balb/c mice 4 hours after intrastromal injection with HAdV37. An increase in threshold cycle (C_T) units reflects an increase in the level of viral gene expression as compared to C_T levels obtained immediately after injection (before any viral gene expression has occurred). Viral mRNAs for E1A and E1B 19k transcripts were present in HAdV37-injected mouse corneas at 4 hours postinfection, but the IIIa viral capsid gene was not detected (n.d.) in the virus-infected cornea in any of three separate experiments at this time postinfection.

FIGURE 11
Immunoblot analysis for phosphorylation of ERK1/2 at 4 and 24 hours after HAdV37 injection into the corneas of Balb/c mice. HAdV37 or buffer was injected intrastromally at the indicated time points, and proteins from three corneas for each condition and time point were harvested and pooled for Western blot analysis. Each phospho-ERK blot (upper of each paired blot) was stripped and reprobed for total ERK (lower of each paired blot). Phosphorylation of ERK1/2 was increased at both 4 and 24 hours after infection with HAdV37 as compared to buffer-injected controls. No differences in total ERK1/2 levels were seen at either time point. This experiment was repeated twice with similar results.
EKC as the outcome of the interaction between adenoviruses and resident cells of the corneal stroma, Chodosh and coworkers\textsuperscript{2-5,99} demonstrated IL-8 and MCP-1 expression by HAdV19-infected keratocytes, and in a series of manuscripts showed that the production of these chemokines was induced by viral binding to the cells, followed by an intracellular signaling cascade leading ultimately to chemokine gene expression. In these studies, ultraviolet light inactivation of HAdV19 did not significantly reduce chemokine expression, whereas heat inactivation of the virus abrogated IL-8 expression, consistent with others’ observations that binding of the virus to its target cell rather than viral replication might be the requisite step preceding corneal stromal inflammation in EKC. The observations of these latter two groups are consistent with one aspect of Jones’ theory of subepithelial infiltrate formation: the presence of nonreplicating viral components in the corneal stroma may be sufficient to induce the necessary inflammatory signal. In this context, the obvious limitations of in vitro studies using human cells in monolayer culture and the limited availability of immunologic reagents for the rabbit model speak to the need for a mouse model of adenovirus keratitis with which to test such observations. The studies presented herein were designed to evaluate the possibility of development of just such a model.

In light of prior evidence that binding of adenoviruses to host cell target receptors was sufficient to initiate proinflammatory cytokine expression,\textsuperscript{2,91} we set out to find a human adenovirus capable of binding and entering mouse cells. We chose to use BALB/c 3T3 fibroblasts, because of their availability and easy growth and maintenance, but confirmed our results in two other cell lines and a total of two mouse strains. Six different subgroup D adenoviruses were used, representing three different adenoviral serotypes, but only HAdV37 caused cytopathic effect in the mouse cells, evident as early as 3 days postinfection (Figure 3), but requiring a relatively high MOI. The infection of mouse cells by a human adenovirus is not completely without precedent. Eggerding and Pierce\textsuperscript{118} demonstrated limited viral gene expression but not cytopathic effect after HAdV2 (subgroup C) infection of Balb/c 3T3 cells. Our growth curve (Figure 4) and viral gene expression studies (Figure 5) confirmed that HAdV37 entered the Balb/c cells and effected early viral gene expression, whereas the TUNEL assay (Figure 6) suggested that mouse cell death due to HAdV37 infection was secondary to apoptosis rather than productive infection and secondary cell lysis. Subsequent evaluation of proinflammatory gene expression in HAdV37-infected Balb/c 3T3 cells by real-time PCR showed that infection was associated with increased transcription of IL-6, KC, MCP-1, and IP-10 at 30 to 90 minutes postinfection (Figure 7), well before even early viral gene expression (Figure 5), suggesting that the transcription of cytokine genes in infected cells occurs despite delayed viral gene expression and in the absence of productive viral replication. These results are consistent with in vitro data in human corneal fibroblasts from Chodosh and coworkers,\textsuperscript{2} in which host chemokine expression preceded HAdV19 gene expression. The apoptosis induced in three different mouse cell lines by HAdV37 was curious in light of the recent observation that HAdV19 infection of human corneal fibroblasts induces activation of the anti-apoptotic PI3K/Akt signaling pathway. Blockade of this pathway with genetic or chemical inhibitors results in increased apoptosis and reduced viral gene expression, perhaps owing to a lack of viable cells in which to replicate.\textsuperscript{99} Why adenovirus infection of human cells results in prolongation of cell viability and enhancement of viral replication by the prevention of apoptosis, whereas infection of mouse cells leads to early apoptosis and a failure of viral replication, remains unknown, but may have to do with the receptors utilized for entry into the cell or with differences in cell signaling after entry. It is tempting to speculate that chemical or molecular inhibition of apoptosis in HAdV37-infected mouse cells might prolong cell viability and make them permissive for viral replication.

In light of the experimental evidence that viral binding rather than viral replication might be responsible for the stromal keratitis in EKC,\textsuperscript{2} our principal goal was to develop a mouse model that would allow for the analysis of early molecular events in the cornea due to viral binding to resident stromal cells, without inducing secondary inflammation from the means of inoculation itself. When scarification and injection of virus-free buffer into the cornea with a metal needle each were shown to result in high background levels of proinflammatory gene expression, we turned to a novel injection method utilizing a fine glass micropipette needle and gas-powered microinjection system that allowed a relatively atraumatic injection of a small volume of virus or buffer directly into the mouse corneal stroma. The atraumatic nature of this system was confirmed by the relatively low level of cytokines expressed in buffer-injected corneas (Figures 8 and 9).

For these studies, we chose to evaluate a limited array of proinflammatory mediators in the HAdV37-infected Balb/c mouse cornea. Mice do not produce IL-8, but because of in vitro evidence that IL-8 might be important to the pathogenesis of EKC,\textsuperscript{2} we focused on the neutrophil chemokine, KC, which is expressed by resident tissue fibroblasts after injury,\textsuperscript{102} and is a close homolog for IL-8. Elevations of KC mRNA (Figure 8) and protein (Figure 9) were found at 4 to 8 hours after infection; these findings may allow the study of mechanisms for increased cytokine gene expression due to adenovirus infection. KC and other cytokines were upregulated despite the absence of viral capsid gene IIIa expression (Figure 10), consistent with a model of adenoviral inflammation in which viral replication is not an essential component.\textsuperscript{100} At the same time, real-time PCR confirmation of early adenoviral gene expression in the living mouse cornea means that HAdV37 must have entered and infected resident cells in the cornea, albeit abortively. Our demonstration of ERK1/2 phosphorylation in virus- but not buffer-injected corneas (Figure 11) was consistent with the hypothesis previously put forth by Chodosh and associates in which inflammation in the corneal stroma after adenovirus infection occurs after activation of a keratocyte signal transduction cascade leading to host proinflammatory gene expression.\textsuperscript{3-5,99} Our HAdV37 mouse model should allow testing of this hypothesis, assuming a way can be found to block specific signaling events in the mouse cornea. Systemic and local administration of protein kinase inhibitors has been shown to reduce neutrophil infiltration in other mouse models of acute inflammation,\textsuperscript{153-157} suggesting that similar approaches might be successful in the cornea.

Our goal was to develop a model in which we could analyze early gene expression events resulting from the interactions between human adenoviruses and resident cells in the mouse cornea. The development of corneal infiltration by leukocytes after corneal infection would represent an unanticipated largesse, but at no time in the limited observation time (first 24 hours) did we recognize
corneal infiltrates or other clinical signs of inflammation in HAdV37-infected Balb/c corneas. Longer observation times after injection combined with immunohistochemical evaluation will be necessary to determine if increases in proinflammatory gene expression are accompanied by the eventual infiltration of leukocytes and the development of clinically recognizable keratitis. Possibly, as in other mouse infection models, strains of mice other than Balb/c might be more susceptible to infection, and experiments are under way to determine if there is a better mouse strain in which to perform these experiments.

**SUMMARY**

This study focused on the development of a mouse model in which to study pathogenic mechanisms in adenovirus keratitis. Data was presented to show that HAdV37, an etiologic agent of EKC, infects mouse cells in vitro, where the virus undergoes limited viral gene expression, induces proinflammatory gene expression by host cells, and eventually kills the cells by apoptosis in the absence of viral replication. A novel means of mouse corneal infection was developed that allows for theatraumatic delivery of virus to the corneal stroma, where host proinflammatory gene and protein expression occurs along with tyrosine phosphorylation of a key protein kinase, despite delayed and ultimately incomplete viral gene expression.

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