The mucosal expression pattern of interferon-ε in rhesus macaques

Andrew Demers  
*University of Nebraska-Lincoln, andrew.demers@huskers.unl.edu*

Guobin Kang  
*University of Nebraska-Lincoln, gkang2@unl.edu*

Fangrui Ma  
*University of Nebraska - Lincoln, fma2@unl.edu*

Wuxun Lu  
*University of Nebraska-Lincoln, wlu2@unl.edu*

Zhe Yuan  
*University of Nebraska-Lincoln, s-zyuan1@unl.edu*

Follow this and additional works at: [http://digitalcommons.unl.edu/biosciqingshengli](http://digitalcommons.unl.edu/biosciqingshengli)

Part of the Allergy and Immunology Commons, Biological Phenomena, Cell Phenomena, and Immunity Commons, Female Urogenital Diseases and Pregnancy Complications Commons, Infectious Disease Commons, Medical Immunology Commons, Medical Pathology Commons, Reproductive and Urinary Physiology Commons, Veterinary Infectious Diseases Commons, Veterinary Microbiology and Immunobiology Commons, and the Virus Diseases Commons

Demers, Andrew; Kang, Guobin; Ma, Fangrui; Lu, Wuxun; Yuan, Zhe; Li, Yue; Lewis, Mark; Kraiselburd, Edmundo N.; Montaner, Luis; and Li, Qingsheng, "The mucosal expression pattern of interferon-ε in rhesus macaques" (2014). *Qingsheng Li Publications*. Paper 18.  
[http://digitalcommons.unl.edu/biosciqingshengli/18](http://digitalcommons.unl.edu/biosciqingshengli/18)
The mucosal expression pattern of interferon-ε in rhesus macaques

Andrew Demers,* Guobin Kang,* Fungrui Ma,* Wuxun Lu,* Zhe Yuan,* Yue Li,* †
Mark Lewis, ‡ Edmund N. Kraiselburd, § Luis Montaner,* and Qingsheng Li* †

*Nebraska Center for Virology, School of Biological Sciences, University of Nebraska-Lincoln, Lincoln, Nebraska, USA; ‡College of Life Sciences, Nankai University, Tianjin, China; †Biosqual, Rockville, Maryland, USA; §Department of Microbiology and Zoology, University of Puerto Rico-School of Medicine, San Juan, Puerto Rico; and †The Wistar Institute, Philadelphia, Pennsylvania, USA

RECEIVED FEBRUARY 11, 2014; REVISED JULY 28, 2014; ACCEPTED AUGUST 2, 2014. DOI: 10.1189/jlb.3A0214-088RRR

ABSTRACT

Type I IFNs play an important role in innate and adaptive immunity against viral infections. A novel type I IFN, namely IFN-ε, which can protect against vaginal transmission of HSV2 and Chlamydia muridarum bacterial infection, has been described in mice and humans. Nevertheless, the principle cell type and the expression pattern of IFN-ε in tissues remain uncertain. In addition, the expression of IFN-ε in Indian rhesus macaques (Macaca mulatta) has not been reported. Here, we analyzed IFN-ε expression in multiple mucosal sites of uninfected or SIV-infected Indian rhesus macaques using IHCs. We report for the first time the detection of IFN-ε expression in situ in the lung, foreskin, vaginal, cervical, and small and large intestinal mucosa of rhesus macaques. We found that the expression of IFN-ε was exclusive to the epithelial cells in all of the aforementioned mucosal tissues. Furthermore, the macaque IFN-ε sequence in this study revealed that macaque IFN-ε is highly conserved among human and other nonhuman primates. Lastly, SIV rectal infection did not significantly alter the expression of IFN-ε in rectal mucosae. Together, these findings indicate that IFN-ε may function as the first line of defense against the invasion of mucosal pathogens. Further studies should be conducted to examine IFN-ε protection against gastrointestinal as well as respiratory infections. J. Leukoc. Biol. 96: 000–000; 2014.

Introduction

Type I IFNs are fundamentally important in innate immunity against viral infections, cellular proliferation, regulation, and effector cell activation of the adaptive immune system [1, 2]. Viral infection can induce the expression of type I IFNs by activating TLRs through their pathogen-associated molecular patterns [3–6].

In recent years, a novel type I IFN, IFN-ε, has been described [7]. The IFN-ε gene is located within the type I IFN gene locus and is conserved across many mammalian species [8–12]. Although the IFN-ε protein has only 30% aa homology with IFN-α and -β [7], IFN-ε was found to use the type I IFN-R chains IFN-α-R1 and IFN-α-R2 for signaling; therefore, IFN-ε is classified as a type I IFN [11, 13]. However, compared with type I IFN-α and -β, IFN-ε has significantly lower antiviral and NK enhancement activity in vitro [9]. In addition, the expression of IFN-ε cannot be induced by pattern recognition receptors or viral infection [8–12]. In contrast, TNF-α stimulation of HeLa cells and seminal plasma in cervico-vaginal tissues could increase the expression of IFN-ε [14, 15]. Furthermore, the constitutive expression of IFN-ε in mouse lung, brain, small intestine, and male reproductive tissues has been reported [7, 9, 12, 16]. In addition, the expression of IFN-ε in epithelial cells in the genital tracts of women and female mice was found to be regulated by estrogen [11]. Very recently, it was also shown that IFN-ε-deficient mice were more susceptible to vaginal transmission of HSV2 and C. muridarum bacteria, the etiologic agent of chlamydiosis, suggesting that IFN-ε may serve as the first line of defense against sexually transmitted pathogens [11]. Despite the clear evidence of IFN-ε expression in mucosal tissues, the spatial distribution pattern in the mucosa, the cell type responsible for IFN-ε expression and the expression level change in response to SIV infection remain poorly understood.

In this study, we sought to determine the following: 1) whether the mucosae of the Indian rhesus macaque, a commonly used, nonhuman primate model of human infectious diseases, including HIV-1, express IFN-ε; 2) which mucosal sites and what type of cells express IFN-ε; and 3) whether

Abbreviations: DAB=diaminobenzidine, ENV=envelope glycoprotein, HNF-1=hepatocyte NF-1, HSV2=herpes simplex 2 viruses, HTLV=human T cell leukemia virus, IHCs=immunohistochemical staining, LP=lamina propria, Mk67=cytokeratin marker, MUSCLE=multiple sequence comparison by log-likelihood, NCBI=National Center for Biotechnology Information, qPCR=quantitative PCR, qRT-PCR=quantitative RT-PCR, Rh=recombinant human, UTR=untranslated region, vRNA= viral RNA

The online version of this paper, found at www.jleukbio.org, includes supplemental information.

0741-5400/14/0096-0001 © Society for Leukocyte Biology

Volume 96, December 2014 Journal of Leukocyte Biology 1

Copyright 2014 by The Society for Leukocyte Biology.
IFN-ε expression is altered in the rectal mucosa after early SIV infection of rhesus macaques. Here, we show for the first time that IFN-ε is expressed in the lung, male and female reproductive tracts, and the gastrointestinal tract of the Indian rhesus macaques. We also show that the distribution of IFN-ε in all of the aforementioned mucosae was exclusive to epithelial cells, and the expression of IFN-ε in the rectal mucosa was unaffected by SIV rectal infection. Finally, we determined the full-length sequence of rhesus macaque IFN-ε mRNA, of which only a partial sequence was available previously. This macaque sequence reveals a high level of conservation across human and other nonhuman primate species. Together, these findings will aid future studies examining the role of IFN-ε in combating mucosal pathogens, not just in the genital tract but also in respiratory and gastrointestinal tract tissues, which have yet to be explored.

MATERIALS AND METHODS

Rhesus macaques and viral inoculation

The mucosal tissues of Indian rhesus macaques (M. mulatta) came from two studies, in which the macaques were housed and maintained in animal housing facilities at Bioqual (Rockville, MD, USA) and the Caribbean Primate Research Center (Puerto Rico), in accordance with the Guide for the Care and Use of Laboratory Animals. All animals were free of simian retrovirus type D, simian T-lymphotropic virus type 1, and herpes B virus. Animals were sedated with ketamine or telazol for all technical procedures and were fully anesthetized for SIV inoculation with SIVmac251 at the dose of 3.4 × 10^6 50% tissue culture infective dose intrarectally. Animals were euthanized by exsanguinations under deep (surgical plane) anesthesia using a 5:1-M ratio. The competition assay was conducted by the coincubation of the peptide antigen with the anti-IFN-ε antibody (1:400 or 1:800) at a 5:1- or 10:1-M ratio at room temperature for 2 h. Then, IHCS was performed as described above.

Immunofluorescence staining to colocalize IFN-ε and Mak6

The experiment was conducted as described above for IHCS, except that the tissue sections were coincubated with a rabbit anti-IFN-ε antibody (1:400) and a mouse Mak6 antibody (1:200 dilution; Invitrogen, Carlsbad, CA, USA), a pan-cytokeratin marker for epithelial cells, overnight at 4°C. After washing in PBS, the slides were incubated at room temperature for 2 h with anti-mouse IgG conjugated to AlexaFluor 594 (1:200 dilution; Life Technologies, Carlsbad, CA, USA) and anti-rabbit IgG conjugated to AlexaFluor 488 (1:200 dilution; Life Technologies). After washing, the slides were coverslipped and examined using an inverted confocal microscope (Olympus IX 81; Olympus, Center Valley, PA, USA).

Detection of SIV RNA in rectal tissue using in situ hybridization

SIV RNA in rectal tissues was detected using in situ hybridization as described previously [18]. Briefly, 6 µm sections were cut and adhered to slides. After deparaffinization in xylene; rehydration in PBS; and permeabilization with HCl, digitonin, and proteinase K, the sections were acetylated and hybridized to 32P-labeled, SIV-specific antisense riboprobes or sense riboprobes as a negative control. After washing and digesting with RNases, the sections were coated with a nuclear track emulsion and exposed, developed, and counterstained with H&E.

IFN-ε quantification in the mucosal tissues of uninfected and infected macaques

Tissue sections stained using IHCS were digitized using Scanscope, and the IFN-ε signal was quantified using the Spectrum Plus analysis program (Version 9.1; Aperio ePathology Solutions, Vista, CA, USA). Briefly, a scanned digital slide was opened in ImageScope, and areas morphologically representative of each tissue were selected for analysis using the ImageScope drawing tools; the signal was quantified using a positive pixel count algorithm in the Spectrum Plus analysis program. The algorithm parameters were manually tuned to match the positive markup image accurately over the DAB stain. Once the parameters were set, the algorithm was applied automatically to all of the digital images to measure the IFN-ε expression in the tissues.

Statistics

A statistical analysis of rectal IFN-ε image signal quantification and qPCR data was conducted using an unpaired Student’s t test, as well as a Wilcoxon rank sum test with R statistical software (http://www.r-project.org). P < 0.05 was considered significant.

Total RNA extraction and PCR amplification of IFN-ε mRNA

Total RNA was extracted from rhesus rectal tissues using a previously published protocol [19]. Briefly, rectal tissues were homogenized with a power homogenizer in TRIzol solution (Life Technologies), followed by purification with an RNeasy Mini Kit (Qiagen, Hilden, Germany). Five micrograms of total RNA was used for RT-PCR with Superscript III RT (Life Technologies) and the IFN-ε-R1 primer 5’-TCATGTGCTTCAAGCTTCTG-3’. The resulting cDNA was amplified via nested PCR using High Fidelity Platinum Taq Polymerase (Life Technologies), the first-round IFN-ε-R1 antisense primer, and the IFN-ε-FORWARD sense primer 5’-ATG ATT ATC ATC AAG CAC TCT TTT GAA-3’. Second-round nested PCR was performed using the IFN-ε-F2 sense primer 5’-ACT CTT GTA TAA GTT GCA AAC C3’ and the IFN-ε-R2 antisense 5’-CTGTAAGACCTGAAACAAAGG-3’ primer.
amplicons were sequenced, and the resulting sequence was used to design primers for amplifying the 5’ and 3’ UTR to characterize the IFN-γ mRNA regulatory elements. The amplification of the IFN-γ mRNA 3’ UTR was performed using the IFN-γ-RACE1 sense primer 5’-CTTGGCCGCTTGGATGTA-3’ and the antisense primer 5’-TTTGGAAATGACCCATATTTAG-3’. For the amplification of the IFN-γ mRNA, the 5’ UTR AD02135 sense primer 5’-CTTAGATATGAACTGATAGGATA-3’ and the antisense IFN-γ-RACE2 primer 5’-GCCAGCAGCAGACACATATTT-3’ were used. The final PCR products were run on a 1% agarose gel, purified using a QIAquick Gel Extraction Kit (Qiagen), and sequenced directly.

The quantification of IFN-γ expression in rectal tissues using qRT-PCR

qRT-PCR was conducted in a final volume of 20 μl with 800 ng cDNA, 0.2 μM of each primer, and Platinum Taq High Fidelity Polymerase (Invitrogen) using the CFX96 Real-Time detection system (Bio-Rad Laboratories, Hercules, CA, USA), using a hot start (95°C for 3 min) and 40 amplification cycles (95°C for 15 s, 57°C for 30 s). The cDNA was synthesized using an Oligo (DT) primer and Superscript III RT (Life Technologies). The following primers and probes were used for amplification and detection: Rh-IFN-γ forward CTG TGG CAA ACC TCA and Rh-IFN-γ reverse 5’-CCT GCT GAA GCA TCT CAT GG-3’; GAPDH forward 5’-ACA TGA TCC CCT GCT CTA CT-3’, Rh-IFN-γ probe 5’-G/MGB/TAG-3’; GAPDH probe 3’-TCT CCT CCT CCT TAT GCG-3’. The full-length Indian rhesus macaque IFN-γ mRNA sequence derived from this study was used for MUSCLE multiple alignment and phylogenetic analysis against the following sequences: chimpanzee (Pan troglodytes), accession #GABE01011555; northern white-cheeked gibbon (Nomascus leucogenys), accession #XM_004092857.1; gorilla (Gorilla gorilla), accession #XM_004047874; and human (Homo sapiens), accession #XM_176891, and the full length was determined experimentally from this study for the rhesus macaque (M. mulatta), accession #KP955535.

Multiple sequence alignment and phylogenetic analysis of IFN-γ mRNA

The full-length Indian rhesus macaque IFN-γ mRNA sequence derived from this study was aligned with other mammalian IFN-γ mRNA sequences obtained from NCBI using the MUSCLE multiple alignment tool [20] with default settings and a maximum iteration of 16 times. The resulting multiple alignments were verified and edited manually in BioEdit. A phylogenetic analysis was conducted using the maximum likelihood method, and the tree was generated using the PhyML program [21]. The HKY85 nucleotide substitution model and the nearest neighbor interchange algorithm were used for the tree topological search. A BioNJ tree was built for the starting tree. The branch lengths and substitution model parameters were optimized for the best tree output. The phylogenetic accuracy and reliability were tested by bootstrapping with 1000 repeat calculations. The tree was viewed and edited using FigTree (by Andre Rambaut; http://tree.bio.e-d.ac.uk/software/figtree/).

Nucleotide sequence accession numbers

The full-length Indian rhesus macaque IFN-γ mRNA sequence derived from this study was used for MUSCLE multiple alignment and phylogenetic analysis against the following sequences: chimpanzee (Pan troglodytes), accession #GABE01011555; northern white-cheeked gibbon (Nomascus leucogenys), accession #XM_004092857.1; gorilla (Gorilla gorilla), accession #XM_004047874; and human (Homo sapiens), accession #XM_176891, and the full length was determined experimentally from this study for the rhesus macaque (M. mulatta), accession #KP955535.

RESULTS

IFN-γ is expressed in the cervix, vagina, and foreskin of the Indian rhesus macaque

Previous studies have demonstrated that IFN-γ could be detected in epithelial cells in the female reproductive tracts of mice and humans [11, 12]. In addition, mouse testes were found to express IFN-γ, demonstrating that both the male and female reproductive organs can express IFN-γ [12]. However, whether IFN-γ is expressed in nonhuman primates is unknown. We examined IFN-γ expression in the vagina and cervix tissues of four SIV-uninfected female Indian rhesus macaques and the foreskin tissue of seven SIV-uninfected male Indian rhesus macaques using IHCs. In the vagina and ectocervix tissues, IFN-γ was expressed exclusively in epithelial cells (Fig. 1A and B and Supplemental Fig. 1) but not in the LP, follicular aggregates, or any other cells in these tissues.
This result was confirmed by the colocalization of IFN-\(\epsilon\) and Mak\(\delta\), a marker for pan-cytokeratin (epithelial-specific proteins), using immunofluorescence staining (Supplemental Fig. 1). Of note, the highest IFN-\(\epsilon\)-expressing cells in the vagina and ectocervix were the basal epithelial cells. Similar to the ectocervix and vagina, in the endocervix, IFN-\(\epsilon\) expression was localized solely in the single layer of columnar epithelial cells (Fig. 1C and Supplemental Fig. 1).

Similar to the female reproductive tract tissues, IFN-\(\epsilon\) expression in the foreskin (Fig. 1D) was localized in unkeratinized epithelial cells. Again, the highest IFN-\(\epsilon\) expression was in the basal epithelial cells. However, the signal intensity in the foreskin relative to the female reproductive tissues was significantly lower. Together, these findings demonstrate for the first time that the epithelial cells lining the cervix, vagina, and foreskin of rhesus macaques constitutively express IFN-\(\epsilon\).

The antibody used in this study is against human IFN-\(\epsilon\) and its specificity for rhesus macaques IFN-\(\epsilon\) was unknown. To confirm the specificity of the antibody, we conducted IHCS using an isotype control antibody, in which no staining was observed (Fig. 1E and Supplemental Fig. 2). As the human IFN-\(\epsilon\) antigen amino acid sequence contains five mismatches with the same region of the rhesus macaque sequence, to confirm further the specificity of the antibody and ensure that the human IFN-\(\epsilon\) antibody was recognizing rhesus macaque IFN-\(\epsilon\), we performed a peptide antigen and antibody competition assay. The preincubation of the peptide antigen with the IFN-\(\epsilon\) antibody at a 5:1-M ratio resulted in reduced staining compared with the nonpeptide control (Supplemental Fig. 3), and a 10:1 ratio resulted in complete elimination of IFN-\(\epsilon\) staining of the tissue. This reduction in staining could be completely rescued when a nonspecific peptide (HTLV) of similar length was used. Together, these data demonstrate the specificity of this antibody in recognizing rhesus macaque IFN-\(\epsilon\) protein (Supplemental Fig. 3).

**IFN-\(\epsilon\) is expressed in lung mucosal tissue**

IFN-\(\epsilon\) mRNA has been detected previously in mouse lung tissue through qPCR [7]. However, it had not been reported in any other animals. Here, we sought to determine if IFN-\(\epsilon\) is expressed in the lungs of Indian rhesus macaques. We identified IFN-\(\epsilon\) expression in the lungs of all three animals examined. IFN-\(\epsilon\) was expressed in the epithelial cells lining the bronchioles (Fig. 2 and Supplemental Fig. 1) but not in other cells, including epithelial cells of the alveoli. This finding revealed that the respiratory mucosa of Indian rhesus macaques express IFN-\(\epsilon\), with expression exclusive to bronchial epithelial cells, supporting the notion that IFN-\(\epsilon\) expression is limited to epithelial cells in the mucosa.

**IFN-\(\epsilon\) is expressed in the mucosa of small and large intestines**

Next, we sought to determine whether IFN-\(\epsilon\) expression could be detected in intestinal mucosa, especially the rectum, which is an important portal of entry for many pathogens, including HIV-1 [22]. In the jejunum (Fig. 3A and B) and the rectum (Fig. 3D–G, top, and Supplemental Fig. 1), robust expression of IFN-\(\epsilon\) was detected in the epithelial cells but not in the LP or any other cells using IHCS. These data demonstrate that epithelial cells of the small and large intestinal mucosae of the rhesus macaque express IFN-\(\epsilon\).

**IFN-\(\epsilon\) expression in rectal mucosa was not altered in the early rectal transmission of SIV**

Receptive rectal intercourse is a common mode of HIV-1 transmission in humans. As we found that IFN-\(\epsilon\) is constitutively expressed in the rectal epithelial cells of SIV-uninfected macaques, we next wanted to determine whether IFN-\(\epsilon\) expression in the rectal mucosae would be altered during early rectal transmission of SIV. Rectal tissues collected at 10, 14, or 28 days post-SIVmac251 inoculation, confirmed to be SIV vRNA positive through in situ hybridization (Fig. 3D), and qRT-PCR (data not shown) were used for this purpose. Quantitative image analysis of rectal tissues stained using IHCS (Fig. 3E and G) showed no significant difference in IFN-\(\epsilon\) levels between the uninfectected and acutely infected rectal tissues or between the different time-points postinfection, and qRT-PCR confirmed that IFN-\(\epsilon\) mRNA expression was not different between the uninfected and infected animals (Supplemental Fig. 4). Furthermore, the spatial distribution of IFN-\(\epsilon\) (Fig. 3E and F) within these tissues was similar. These data indicate that SIV infection of the rectum did not alter the expression of IFN-\(\epsilon\), at least in acute infection, which supports the previous notion that IFN-\(\epsilon\) expression is not induced directly by viral infection.
Figure 3. IFN-γ expression in the gut mucosae of uninfected and SIV-infected macaques. Representative micrographs of IFN-γ expression (stained as brown) in the epithelial cells lining the jejunum (A and B) and rectums (E–G, top) of seven SIV-uninfected Indian rhesus macaques, detected using IHCS. (C) Quantitative analysis of IFN-γ expression in the rectums of uninfected or 12 SIV-infected macaques; the box plot shows the ratio of IFN-γ-positive pixels versus the epithelial area in pixels. The differences were not significant (horizontal black line denotes the median, boxes denote 25 and 75 percentiles, and whiskers denote the sd). (D) SIV vRNA+ cells in rectums detected using in situ hybridization with an S35-labeled ribo-probe. After radioautographic development, the clusters of discrete black silver grains (white arrows) overlay the vRNA+ cells at 6, 10, 14, and 28 days postinoculation (dpi). (E) Representative images of IFN-γ staining in infected rectal tissues from three male animals at each time-point postinfection. (F) The images at high magnification from the boxes in image E. (G) Markup images, in which red, pink, and yellow represent strong, medium, and weak positive signals, respectively, used for quantification of IFN-γ in SIV-uninfected and infected rectums at different time-points postinfection. FA, Follicular aggregate.
conclude definitely that IFN-ε expression is not altered in SIV infection. Therefore, the role of IFN-ε in protecting against rectal transmission of SIV remains unknown and is beyond the scope of this study.

IFN-ε mRNA is highly conserved among nonhuman primates

Mouse and human IFN-ε have been sequenced previously; however, for rhesus macaques, only a putative sequence was available before this study. To confirm the specificity of our IFN-ε IHCS and to obtain the full-length IFN-ε sequence of the Indian rhesus macaque, IFN-ε mRNA, isolated from rectal tissues, was amplified using RT-PCR and sequenced. The macaque full-length coding sequence of IFN-ε (positions 622–1203 in the full-length sequence), obtained from this study, was aligned and compared with the available IFN-ε sequences derived from the mRNA of human and other nonhuman primates in the NCBI database using MUSCLE multiple sequence alignment [20, 23]. Overall, there was a 94% nucleotide identity in the IFN-ε coding sequences of rhesus macaques, humans, and other nonhuman primate species. The 5′ UTR region of the rhesus macaque had not been predicted before this study. The alignment of the 5′ UTR region derived from this study (positions 1–621 in the full-length sequence) revealed that the rhesus macaque IFN-ε 5′ UTR region has 96% identity to the human and other nonhuman primate 5′ UTR sequences, including several known transcription factor-binding sites (Supplemental Fig. 5). The most notable difference in the 5′ UTR region was at positions 570–574, in which the rhesus macaque and olive baboon (Papio anubis) have a 4-nt insertion/disruption of the HNF-1 transcription factor-binding site (Supplemental Fig. 5). This observation suggests that the function of the IFN-ε protein and the expression regulation of the IFN-ε gene are likely to be highly similar among other nonhuman primates and humans.

Maximum likelihood phylogenetic analysis [21] (Fig. 4) resulted in two distinct clusters or groups. The full-length rhesus macaque IFN-ε mRNA clustered with olive baboon in one group, while human, chimpanzee, and other nonhuman primates clustered in the other group. Overall, the sequence conservation observed in the IFN-ε mRNA Indian rhesus macaque, relative to the other IFN-ε mRNA sequences, demonstrates that IFN-ε is highly conserved among human and nonhuman primates.

DISCUSSION

Here, we report for the first time the expression of IFN-ε in epithelial cells lining the jejunal, rectum, bronchioles (lung), and both female and male genital tract tissues of rhesus macaques. Previous studies in mice demonstrated that IFN-ε was expressed by mucosal epithelial cells only in the female reproductive tract and could protect against HSV2 and C. muridarum infections [11]. Furthermore, before this study, there was only a predicted rhesus macaque IFN-ε mRNA sequence, which did not include the 5′ UTR region. In this study, we show that the rhesus macaque IFN-ε 5′ UTR region is 96% identical to that of humans and other nonhuman primates. In addition, with the exception of only the HNF transcription factor-binding site, the transcription factor-binding sites are conserved among the rhesus macaque, human, and nonhuman primate IFN-ε 5′ UTRs, including in the progesterone transcription factor-binding site. Previous studies have shown that mouse and human IFN-ε transcription was increased in response to higher estrogen levels within female genital tract tissues [11]. Our finding of IFN-ε expression in the lungs and gastrointestinal tract and the conservation of several transcription factor-binding sites may suggest a tissue-specific mechanism for regulating the expression of IFN-ε, as sex hormones, such as estrogen or others, are likely to be largely absent in these tissues [11]. The dichotomy of the absence of IFN-ε transcriptional up-regulation in response to viral infection and the fact that it can provide protection against viral infection [11, 12] remain poorly understood, although one plausible explanation is that the IFN-ε-protective effect may be dependent on other cofactors expressed in cells or specific tissues that are inducible in response to pathogens.

Of note, the full-length macaque IFN-ε mRNA sequenced in this study revealed high homology (94% identity) in the coding region with that of humans and other nonhuman primates. Therefore, it is plausible that IFN-ε would have similar function across these species. Further investigation would be needed to address this question, as well as the function and...
protective effects of IFN-ε in the gut and lung tissues and between female and male reproductive tissues, as these mucosal sites have drastic anatomic, physiological, and microbiological differences. The examination of the breadth and potency of IFN-ε in mediating protection against different pathogens in vivo in different mucosal tissues is also needed.

Previous studies have demonstrated that IFN-ε expression could not be induced by viral infection in vitro or in vivo. Consistent with previous in vitro and in vivo studies in mice [11, 12], IFN-ε expression was not altered significantly throughout the early course of infection, suggesting that rhesus macaque IFN-ε expression in the rectum is not influenced directly by SIV infection. One caveat of this study is the small sample size; therefore, future studies using larger numbers of rhesus macaques may be better suited to fully address whether SIV infection has any influence on rhesus macaque IFN-ε expression.

In conclusion, we report here, for the first time, the expression of IFN-ε in epithelial cells of multiple mucosal tissues in Indian rhesus macaques. In addition, we show that Indian rhesus macaque IFN-ε expression in the rectum is not altered after SIV infection.

Finally, rhesus macaque IFN-ε mRNA, both in the coding region and the 5′ UTR, is highly conserved compared with humans and other nonhuman primates. The findings reported here may aid future studies to address the role of IFN-ε in protecting against mucosal infections.

AUTHORSHIP
A.D. and Q.L. conceived of and designed the experiments and wrote the manuscript. G.K., Y.L., and F.M. provided aid in data analysis. W.L. and Z.Y. performed tissue collection. Z.Y. also provided aid for the qPCR assay design. M.L. and E.N.K. housed and cared for the animals. L.M. provided some of the samples used in this study and contributed to the discussion.

ACKNOWLEDGMENTS
This work was supported by U.S. National Institutes of Health Grants DK087625 (to Q.L.), PO4 OD012217 (to M. I. Martinez, University of Puerto Rico), and R01AI094603 and R01AI084142 (to L.M.). We thank members of the Q.L. lab for their insightful technical advice on this project. The authors thank Mark Lewis at Bioqual and for the support of T. A. Santiago, P. P. Maldonado, C. A. Sariol, and M. I. Martinez at the University of Puerto Rico for all of the macaque care and of S. Abdulhaqq and B. Ross at Wistar. We also thank Dr. Dong Wong for his assistance with statistical analysis of the data and Dr. Todd Wical for editing of this manuscript.

DISCLOSURES
The authors declare no conflicts of interest.

REFERENCES


SUPPLEMENTARY FIGURE LEGENDS

Supplemental Figure 1: Confocal images showing co-localization of IFN-ε and pan-cytokeratin confirming IFN-ε is expressed in epithelial cells. IFN-ε (green) and pan-cytokeratins (red) in the vagina, endocervix, rectum or lung tissues from uninfected rhesus macaques were detected using immunofluorescence staining. Rabbit anti-IFN-ε and mouse anti-pan-cytokeratin antibodies (Mak6) were used as primary antibodies, and anti-rabbit-Alexa 488 and anti-mouse Alexa 594 were used as secondary antibodies. The merged panel shows the co-localization (yellow) of IFN-ε and Mak6 over the differential interference contrast (DIC) images of the designated tissues.

Supplemental Figure 2: Rabbit IgG isotype control for IHCS of rhesus macaque tissues. Rabbit IgG was incubated in the same concentration as IFNE antibody on respective tissues under the same IHCS conditions as used for IFNE staining. Each image is representative of at least 3 sections from at least 3 animals for each tissue examined. Scale bars are drawn to 100 µm.

Supplemental Figure 3: An anti-human IFN-ε antibody specifically stains IFN-ε proteins, as demonstrated by peptide antigen and antibody competition assays. IFN-ε peptide antigen or HTLV envelope peptide and antibody competition staining assays on rectal tissues of an SIV-uninfected rhesus macaque. Human anti-IFN-ε antibody at a working dilution of 1:400 or 1:800 was incubated at room temperature for 2 h without IFNE or HTLV peptide antigen (no peptide) or with peptide antigen at a 5:1 (5X), 10:1 (10X) molar ratio, respectively, prior to a standard IHCS procedure.
Supplemental Figure 4: IFN-ε mRNA expression is unaltered in SIV-infected rhesus macaques. Quantitative RT-PCR of IFN-ε mRNA expression was normalized to GAPDH expression from the rectal tissues of SIV-uninfected rhesus macaques or at 28 days post SIV infection. Data are shown as the mean ± SEM of 3 individual rhesus macaques. There was no significant difference using an unpaired Student’s t test.

Supplemental Figure 5: The IFN-ε 5’ UTR of the rhesus macaque has 96% identity with human and non-human primates. MUSCLE alignment of the IFN-ε 5’UTR nucleotide sequence from non-human primates, humans and mice. Transcription factor binding sites are shaded in yellow, and black boxes highlight nucleotide mismatches from mouse and non-human primates. Transcription factor binding sites were determined based on conservation of human IFN-ε 5’ UTR and Hardy et. al. as a reference [7].
Supplemental Figure 5

[Image of a figure showing sequence alignments for different species, including Mouse, Macaca, Papio Anubis, Pongo Abelii, Gorilla, Human, and Chimpanzee, with highlighted regions for SATB1 and ISL1]