First Phase 1 Double-Blind, Placebo-Controlled, Randomized Rectal Microbicide Trial Using UC781 Gel with a Novel Index of Ex Vivo Efficacy

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Abstract

Objectives: Successful control of the HIV/AIDS pandemic requires reduction of HIV-1 transmission at sexually-exposed mucosae. No prevention studies of the higher-risk rectal compartment exist. We report the first-in-field Phase 1 trial of a rectally-applied, vaginally-formulated microbicide gel with the RT-inhibitor UC781 measuring clinical and mucosal safety, acceptability and plasma drug levels. A first-in-Phase 1 assessment of preliminary pharmacodynamics was included by measuring changes in ex vivo HIV-1 suppression in rectal biopsy tissue after exposure to product in vivo.

Methods: HIV-1 seronegative, sexually-abstinent men and women (N = 36) were randomized in a double-blind, placebo-controlled trial comparing UC781 gel at two concentrations (0.1%, 0.25%) with placebo gel (1:1:1). Baseline, single-dose exposure and a separate, 7-day at-home dosing were assessed. Safety and acceptability were primary endpoints. Changes in colorectal mucosal markers and UC781 plasma drug levels were secondary endpoints; ex vivo biopsy infectibility was an ancillary endpoint.

Results: All 36 subjects enrolled completed the 7–14 week trial (100% retention) including 3 flexible sigmoidoscopies, each with 28 biopsies (14 at 10 cm; 14 at 30 cm). There were 81 Grade 1 adverse events (AEs) and 8 Grade 2; no Grade 3, 4 or procedure-related AEs were reported. Acceptability was high, including likelihood of future use. No changes in mucosal immunoinflammatory markers were identified. Plasma levels of UC781 were not detected. Ex vivo infection of biopsies using two titers of HIV-1Lai showed marked suppression of p24 in tissues exposed in vivo to 0.25% UC781; strong trends of suppression were seen with the lower 0.1% UC781 concentration.

Conclusions: Single and 7-day topical rectal exposure to both concentrations of UC781 were safe with no significant AEs, high acceptability, no detected plasma drug levels and no significant mucosal changes. Ex vivo biopsy infections demonstrated marked suppression of HIV infectibility, identifying a potential early biomarker of efficacy. (Registered at ClinicalTrials.gov; #NCT00408538)

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Introduction

Efforts to reduce the sexual transmission of HIV-1 are pivotal to controlling the AIDS pandemic. Sustained plasma suppression reduces transmission but trials of HIV-specific vaccines and topical microbicides have been challenging in heterosexual couples and men who have sex with men (MSM) populations, especially given the still-poorly understood immune responses at the sexually-exposed mucosal portals of virus entry [1–9]. The recent results from both the Phase IIb CAPRISA 004 Trial of vaginally-applied 1% tenofovir gel and the Phase III iPrEx Trial of oral Truvada tablets (a co-formulation of tenofovir disoproxil fumarate and emtricitabine) have been exciting, first-time achievements in HIV prevention [10,11].

Microbicides have been advanced as a topical mode of reducing HIV-1 transmission “per sexual act.” While discussed as a topical version of PrEP [12], use of topical microbicides is intended to provide a safe, acceptable, affordable form of protection from HIV-1 transmission, providing receptive partners (women and men) with options, especially when condom use is non-negotiable [13]. The spermicidal and contraceptive vaginal agent, nonoxynol-9 [N9] was demonstrated, post-approval, to create an increased risk for HIV-1 acquisition with frequent vaginal use. Significant epithelial sloughing was seen when applied rectally. This experience identified newer safety parameters to consider when evaluating microbicidal agents [14–16]. Until recently, clinical trial efforts have focused on vaginal transmission with mostly disappointing results [17–21]. A first-in-field success, CAPRISA 004 utilized a reverse-transcriptase inhibitor (1% tenofovir) gel applied 12 hours before and after vaginal intercourse. The study demonstrated a >50% reduction in HIV-1 transmission in those women using the gel for >80% of episodes [10,11]. Equally exciting, in different risk groups, was the recent iPrEx trial demonstration of 44% reduction of HIV-1 transmission in ~2500 higher-risk MSM at 11 study sites worldwide [11]. As in the CAPRISA trial, when the inherently difficult issue of adherence is teased apart, sub-analyses suggest the prevention rate may be 50% or higher. Both studies successfully demonstrated proof-of-concept for topical microbicides.

Rectal transmission of HIV-1 is thought to be 20–200-times more likely per sexual act than vaginal transmission, perhaps related to the single-cell epithelial lining and extensive, activated resident immune populations [1–7,22,23]. Receptive anal intercourse (RAI) is highly prevalent among MSM and also in heterosexual sexual partnerships [24–30]. It is anticipated that when the mucosa is co-infected (such as with HSV) or significant trauma, the rate of rectal transmission per sex act would markedly increase [31–34].

This report describes the first IND-supported Phase 1 safety trial of two concentrations of UC781 (0.25% and 0.1%) as a rectal microbicide. UC781 is a potent non-nucleoside reverse transcriptase inhibitor (NNRTI) which binds tightly to HIV-1 RT [35–40], has activity against a wide range of subtype HIV-1 isolates and is poorly absorbed from mucosal surfaces with systemic limited bioavailability. UC781 displays in vitro nanomolar range EC_{50} activity against wild type HIV-1 virus and little to no cytotoxic effect on cell lines and primary cells. In pre-clinical studies of human cervical and colorectal explants pre-incubated with UC781, R5 HIV_{BaL} was markedly suppressed, decreasing the infection in migrating lymphoid cells [41,42]. UC781 added in vitro showed 100% inhibition of HIV_{BaL} at 3.3 μg/ml and 90% inhibition at 0.33 μg/ml. These infectious doses are thought to be far in excess of ejaculate concentrations [43–45]. For comparison, the delivered doses (empirically assuming a 10× dilution by rectal fluids) in this trial for the 0.1% gel was a dose of 3.5 mg in 3.5 ml (1000 μg/ml) and for the 0.25% gel, a dose of 0.75 mg in 3.5 ml (2500 mg/ml).

Two concentrations of UC781 gel (0.10% and 0.25%) formulated for topical vaginal application and demonstrating safety in early dose-ranging vaginal safety studies [46–40] were applied rectally in this study. The product was delivered using the same applicator design as used in vaginal microbicide trials. The novelty of this “first in field” study was facilitated by the HPTN-056 study which established normative ranges and inter-subject variabilities in a host of recently developed mucosal indices to assess potential mucosal injury [49].

An innovative 2-stage trial design was used consisting of an initial, single rectal application of either of the two concentrations of UC781 or hydroxyethyl cellulose (HEC) placebo gel followed several weeks later by 7 once-daily, self-administered rectal applications. We have previously reported the acceptability evaluations carried out in this trial [50]. Here we report the clinical safety of vaginally-formulated UC781, plasma drug concentrations, mucosal safety and the use of novel ex vivo biopsy HIV-1 infection challenges of in vivo exposed tissue samples. Both concentrations of UC781 were shown to be clinically safe by all indices used with few Grade 2 and no Grade 3, 4 or procedure-related AEs. Plasma levels of UC781 were not seen and no significant mucosal abnormalities were detected. Remarkably, the ex vivo HIV-1 infection of tissue biopsies showed marked suppression with UC781 0.25%.

Methods

The study was designed by the investigators with collaborative input from CONRAD and the NIAID/DAIDS/Prevention Sciences Integrated Preclinical-Clinical Program (IPCP), as stipulated in the award notice and reviewed by the U.S. Food and Drug Administration (FDA). The study was approved by the UCLA Office of the Human Research Protection Program Institutional Review Board (UCLA IRB) and all subjects provided written informed consent. The trial is registered at ClinicalTrials.gov, number NCT00408538 and is in compliance with the CONSORT 2010 recommendations for reporting of trial results (www.consort-statement.org) [51,52].

Study population

HIV-1 seronegative men and women between the ages of 18 and 64 (inclusive) with a history of consensual receptive anal intercourse (RAI) at least once in lifetime were eligible for screening. This was included to identify individuals with some familiarity of sexual or product insertion per rectum. Lower risk of HIV infection was supported by HIV screening tests for inclusion, the additional inclusion criteria of willingness to be sexually active regarding rectal sex or any other rectal insertions one week prior to treatment and one week before and following each flexible sigmoidoscopy. Ongoing monitoring of plasma HIV RNA levels is further supported this. Exclusion criteria included HIV infection, known inflammatory bowel disease or any other chronic gastrointestinal disorder and/or history of significant gastrointestinal bleeding as well as any allergies to methylparaben, propylparaben or sorbic acid.

Study Products

Two concentrations of UC781 (0.1% or 0.25% w/w), Carbomer 974P, methylcellulose, glycerin, purified water and common preservatives (methylparaben and propylparaben), adjusted to pH 5.2, were prepared in vaginally-formulated, aqueous

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gel formulations. The single dose, prefilled vaginal applicators (the same type of applicators as used in UC781 vaginal microbicide trials) were packaged, overwrapped and shipped directly to the study site pharmacy. Each applicator contained either a dose of 3.5 mg in 3.5 ml (1000 μg/ml) for the 0.1% gel, a dose of 3.75 mg in 3.5 ml (2500 μg/ml) for the 0.25% gel, or 3.5 g in 3.5 ml of aqueous HEC (hydroxyethyl cellulose) gel, adjusted to pH 4.4, as the placebo gel [53].

Study Design
This was a single site, blinded, multi-arm, two-treatment stage Phase 1 trial using two established UC781 drug concentrations and HEC placebo in 36 subjects in a 1:1:1 randomization (Figure 1). The protocol for this trial and supporting CONSORT checklist are available as supporting information; see Checklist S1 and Protocol S1. The first stage was a randomized single exposure, directly applied by clinical personnel, with the subject remaining supine for the 30 minute period until sample collection. The second stage, assessed separately, was a randomized (same groups), 7-day cumulative exposure delivered by participants at home. Participants were given a Product Use Log for the 7-day use and were called daily to encourage use and recording of symptoms/time of use.

The primary objectives of the study were to evaluate the safety and acceptability of 0.1% and 0.25% UC781 vaginal microbicide gel versus placebo when applied rectally.

The primary endpoints were (i) the frequency of ≥Grade 2 adverse events and (ii) extensive acceptability evaluations (reported elsewhere [50]. The primary tool for safety assessments and grading adverse events was the DAIDS AE Grading Table Version 1.0, December 2004, Addenda 1 (female genital) and 3 (rectal-specific) Grading Tables for Use in Microbicide Studies (http://rsc.tech-res.com/safetyandpharmacovigilance). AE gradings are defined as symptoms causing no or minimal (Grade1), causing greater than minimal (Grade 2) interference with or causing inability to perform (Grade 3) usual social and functional activities. Physical signs and laboratory values are graded similarly to the ones listed in the main DAIDS AE Grading Table. Details of demographics and adverse events are provided in Table S1 and Table S2.

One secondary objective of the study was to determine whether use of study products was associated with rectal mucosal damage. Secondary endpoints included rectal epithelial sloughing, histopathology, rectal microflora, fecal calprotectin, and a group of “mucosal immunotoxicity indices,” which included flow cytometric characterization of isolated mucosal mononuclear cells (MMCs), secreted mucosal immunoglobulins, secreted rectal fluid cytokines, mucosal tissue cytokine mRNA and susceptibility to HIV infection of rectal tissue biopsies.

An additional secondary objective was to determine the pharmacokinetics (PK) of UC781 vaginal microbicide gel administered rectally in a subset of participants. The endpoint was UC781 blood levels as an indicator of absorption of drug from the GI tract. Enrollment was protocol-defined as having met initial, screening criteria at Visit 1 (eligibility requirements, consent signing, STI evaluations and baseline acceptability questionnaires answered via CASI) and having been randomized post Visit 2. During Visit 2, baseline sample collections and flexible sigmoidoscopy for each subject’s baseline mucosal biopsy infectibility, PK as well as additional acceptability assessments occurred. PK assessments for UC781 were measured in a subset of enrolled, randomized participants. All subject visits were at the UCLA Digestive Diseases Clinic and/or UCLA Outpatient Endoscopy Suite.

Following enrollment, 36 subjects were randomized, in a double-blind fashion, to one of the three treatment arms. Randomization was performed in blocks of six, with each block containing two assignment codes for each of the three treatment groups. A subset of 9 subjects (3 from each group) also participated in a PK sub-study, which included 6 timepoints (pre-dosing, 0.25 hr, 2 hr, 4 hr, and 24 hr after the single dose, and 1 day post-daily dosing for 9 of the 36 study participants (3 participants in each study group: placebo, 0.25%, UC781 and 0.1% UC781) were randomized for UC781 plasma levels for 24-hour PK (baseline/pre-dosing, 30 minute, 2, 4 and 24 hours following single exposure) and also single cumulative level following 7-day dosing (total of 54 samples). Randomization codes were generated by the study biostatistician using computer-generated random numbers. Once generated, the UCLA investigational pharmacy held primary responsibility for dispensing drug, randomizing subjects and maintaining the blind.

Each participant had three flexible sigmoidoscopies for mucosal biopsies and other sample collections: baseline (Visit 2), 30 minutes post single topical exposure (Visit 3) and the morning following completion of 7 daily at-home topical exposures (Visit 5). Rectosigmoid colon biopsies (referred to hereafter as “colonic biopsies”) were collected endoscopically using large cup biopsy forceps via a flexible sigmoidoscope as previously described [49,54]. The first set of colorectal biopsies were then acquired at 10 cm from the anal verge and then at 30 cm (14 biopsies acquired at each site; 28 total biopsies per visit).

Each study visit to the endoscopy suite for flexible sigmoidoscopy had an established sequence to minimize confounding of endpoint results. Phlebotomy for safety laboratories as well as UC781 PK plasma samples occurred first (pre-exposure at Visit3 and then at 30 minutes, 2 hr, 4 hr and 24 hr); rectal sponges (ULTRACELL® Aspen Surgical #40415, Caledonia, Michigan) then were applied rectally in a subset of participants. The endpoint was UC781 pharmacokinetics (PK) of UC781 vaginal microbicide gel administered rectally in a subset of participants. The endpoint was UC781 blood levels as an indicator of absorption of drug from the GI tract.
collected unit for stool sample collection for calprotectin (same-day shipment to Genova Diagnostics, Asheville, NC). Microflora samples were shipped same day for characterization to Dr. Sharon Hillier’s Laboratory at Magee-Womens Research Institute, Pittsburgh, PA. Participants were then taken to the endoscopy room. Following digital examination, the flexible sigmoidoscope was introduced to a distance of 5 cm in the rectal vault where 50 cc of Normosol® solution was slowly infused. Following 30 seconds, at least 25–30 ml of lavage fluid was aspirated through the endoscope to a collection trap vial to assess epithelial sloughing. The endoscope was then advanced to collect biopsies as described above. Endoscopic pictures/grading were not recorded due to the known variability of the colorectal lining in health.

Behavioral and acceptability measures (primary endpoint)

All participants completed the Baseline Behavioral Questionnaire (BBQ) prior to Visit 3. Product Acceptability Questionnaires (PAQs) were completed after the 7-day exposure. The PAQ included both close-ended and open-ended questions, assessed using a password-protected Web-based survey, enabling direct data entry. An extensive, final 1-hour, in-depth acceptability interview was conducted. Detailed methods and results of this portion of this trial have been published [50].

Indicators of rectal mucosal damage

A diverse panel of mucosal assessments to evaluate potential injury or significant alteration from baseline immune parameters was used to detect changes post-gel exposure compared to baseline findings within individual subjects and between study groups [49]. The broad profile of mucosal markers of inflammation/injury is included based on several years of development, optimization and determination of in vitro stability, reproducibility and variability. The clinical relevance of two-fold or more changes in any of these parameters is not known but the absence of any changes provides a degree of confidence of non-injury [49]. The assays used to monitor mucosal injury, activation and/or inflammation are described below. Following the methods is a listing of which samples were collected at which visits.

Fecal calprotectin

During enhanced mucosal inflammation, intestinal granulocytes, containing large amounts of the cytoplasmic protein calprotectin, migrate through the mucosal epithelia and granulocyte-derived calprotectin can be found in feces, providing a useful indirect index of mucosal inflammation [56,57]. Fecal calprotectin levels correlate well with disease activity in Crohn’s disease and ulcerative colitis [58,59] and were utilized here to indirectly detect acute inflammation. Stool was collected using a kit with mailing materials (Genova Diagnostics, Asheville, NC). A 20 ml stool sample was collected from participant’s enema evacuations and mailed same day. Reported results from the company have a range of <30 μg/ml = normal; greater than 50 μg/ml to 500 μg/ml equating to varying levels of inflammation. Fecal calprotectin assays have a sensitivity of 96% in discriminating between healthy controls (2 mg/l, 95% CI 2–3 mg/l) and subjects with active inflammatory bowel disease (91 mg/l, 95% CI 59–105 mg/l) [56,57].

Rectal secretions of immunoglobulins and cytokines collected by sponges

Rectal secretions were collected using 4 cellulose sponges (ULTRACELL®, Aspen Surgical Caledonia, Michigan) for secreted immunoglobulins (2 dedicated sponges) and secreted cytokines (2 dedicated sponges) per collection event, as previously reported [49,60]. Sterile sponges were pre-moistened with 50 μl of phosphate buffered saline (Gibco BRL, Gaithersburg, MD) and attached to an adapted 2 ml plastic transfer pipette (Fisher Scientific, Pittsburgh, PA). Two pipettes with attached sponges were introduced into the rectum via the anoscope and held against the rectal mucosa under direct vision for 5 minutes, then placed on ice until freezing at −80°C for batch processing. Samples with visible blood were discarded. Sponge tips were transferred to a 2 ml Spin-X column (Corning, Corning, NY), from which the acetate membrane had been removed. This was repeated for the second set of two sponges. Absorbed rectal secretions were eluted twice with a total volume of 250 μl of cold elution buffer (PBS containing 0.25% BSA (Sigma Chemicals, St Louis, MO), 1% Igepal (Sigma Chemicals, St Louis, MO) and 1× protease inhibitor cocktail (Sigma Chemicals, St Louis, MO) from the sponges by centrifugation (10,000 rpm, 30 minutes at 4 degrees). The recovered eluate was transferred to a pre-weighted 1.5 ml Eppendorf tube (Fisher Scientific, Pittsburgh, PA) and re-weighted. The recovered volume of secretion was calculated by subtracting the recovered volume from that recovered from control sponges that were run in parallel. Duplicate samples were pooled, (usually yielding 400 μl), frozen and retrieved in batches for further analysis.

Total IgG and total IgA were quantified (in duplicate) in the eluted rectal secretions by ELISA [54] and results expressed as ng/ml. Values in ng/ml were extrapolated from the relevant standard curves and means calculated for each sample. The Coomassie DryTM Blue Protein Assay (Pierce, Rockford, IL) was used to extrapolate the quantity of total protein expressed as ng/ml [49]. Cytokine samples were quantified (pg/ml) using 50 μl in a Lincoplex® Human cytokine/chemokine multiplex immunoassay kit (used in BioRad Luminex® 100 System multiplexing array instrument). Seven secreted cytokine proteins were quantified in pg/ml (RANTES, MIP-1α, TNF-α, IFNγ, IL-12 (p40), IL-6, IL-1β).

Microflora

Rectal specimens obtained per rectum with a sterile cotton swab were placed in an anaerobic transport tube (Port-a-Cul; Becton-Dickinson Corp., Cockeysville, MD) and shipped by overnight mail to Dr. Sharon Hillier’s Laboratory at Magee-Womens Research Institute, Pittsburgh, PA. The swab was removed, placed into 0.9 ml of buffered salt solution and vortexed to release fluid and processed as previously reported [7]. Plates were incubated in an anaerobic chamber for 72–96 hours for detection of anaerobes; agar plates for aerobic bacteria were evaluated after 48 hours of incubation at 37°C in 6% CO2. Given the enormous variety of normally-present bacteria in the colon and the unknown variations related to diet and time, the bacterial groups selected for monitoring before and after product exposure was based on a FDA defined and accepted panel. These microflora are identified in Table 1. Changes in bacterial frequencies and concentrations within exposed individuals as well as between study groups were examined between baseline visit and following the 7-day exposure (which included any changes occurring after the single dose exposure). Results are quantified by colonization growth on a scale from 0–4 as follows: 0 = no growth; 1 = 103 cfu/ml; 2 = 105 cfu/ml; 3 = 107 cfu/ml; 4 = 109 cfu/ml. Raw means and SDs were computed by time (before or/after gel use), using McNemar’s test for evaluating paired changes in colonization status [61,62].

Epithelial sloughing

Collected lavage fluid (>25 ml) was transported on ice to the laboratory. Using a protocol adapted from Patton and Phillips [14,63], the fresh lavage samples were placed in a 100×15 mm
Petri dish, on the platform of a Leica EZ4 inverted microscope with attached CANON Powershot A630 camera, to count epithelial sheets. The sheets were measured in quadrants using a permanent ruler, affixed to the platform, under the Petri dish, as previously reported [14,63]. Positive findings were defined as a clear, spongy, elastic piece of cellular material (with nucleus) at least 2 mm in size. As epithelial sloughing does not have an absolute, quantifiable threshold, the previously used scoring system of 0–4 was used. Each of four Petri-dish quadrants was scored as either 0 or 1, (absence or presence of epithelial sheets) for a total of 0–4. As epithelial sloughing does not have an absolute, quantifiable threshold, the previously used scoring system of 0–4 was used. Each of four Petri-dish quadrants was scored as either 0 or 1, (absence or presence of epithelial sheets) for a total score from 0–4.

Histology
Histopathological scoring of inflammation was carried out on oriented, formalin-fixed paraffin-embedded, hematoxylin- and eosin-stained tissue biopsies. Colonic mucosal biopsy samples were obtained at both 10 cm and 30 cm (measured by endoscopic markings) from the dentate line. A pathologist with specialty training in gastrointestinal pathology, blinded to sample group, performed the batched, qualitative (scale: 0–5) histological assessments. A validated, qualitative scoring scale of chronic active inflammation used in ulcerative colitis trials previously adapted and reported for baseline mucosal indices in microbicidal training in gastrointestinal pathology, blinded to sample group, performed the batched, qualitative (scale: 0–5) histological assessments. A validated, qualitative scoring scale of chronic active inflammation used in ulcerative colitis trials previously adapted and reported for baseline mucosal indices in microbicidal training in gastrointestinal pathology, blinded to sample group, performed the batched, qualitative (scale: 0–5) histological assessments.

Isolation of mucosal mononuclear cells and flow cytometry
MMCAs were isolated from intestinal biopsies obtained at both 10 cm and 30 cm (measured by endoscopic biopsies) [49]. Percentage values for the stained subsets were recorded.

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### Table 1. Changes in pre-defined rectal bacteria populations following cumulative exposure to 0.1% or 0.25% UC781 topical gel or HEC placebo compared to baseline.

<table>
<thead>
<tr>
<th>Bacteria</th>
<th>0.1% UC781 (n = 12)</th>
<th>0.25% UC781 (n = 12)</th>
<th>HEC (n = 12)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Baseline</td>
<td>Post-exposure</td>
<td>p value</td>
</tr>
<tr>
<td>Lactobacillus (H₂O₂-producing)</td>
<td>33%</td>
<td>50%</td>
<td>0.68</td>
</tr>
<tr>
<td>Lactobacillus (H₂O₂-nonproducing)</td>
<td>17%</td>
<td>17%</td>
<td>1</td>
</tr>
<tr>
<td>Gardnerella vaginalis</td>
<td>0</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Escherichia coli</td>
<td>83%</td>
<td>83%</td>
<td>1</td>
</tr>
<tr>
<td>Other gram-negative rods</td>
<td>17%</td>
<td>8%</td>
<td>1</td>
</tr>
<tr>
<td>Anaerobic gram-positive cocci</td>
<td>92%</td>
<td>92%</td>
<td>1</td>
</tr>
<tr>
<td>Anaerobic gram-positive rods (Clostridium)</td>
<td>67%</td>
<td>83%</td>
<td>0.64</td>
</tr>
<tr>
<td>Anaerobic gram-positive rods (other)</td>
<td>83%</td>
<td>67%</td>
<td>0.64</td>
</tr>
<tr>
<td>Anaerobic gram-negative rods</td>
<td>100%</td>
<td>100%</td>
<td>1</td>
</tr>
<tr>
<td>Black-pigmented anaerobic gram-negative rods</td>
<td>58%</td>
<td>83%</td>
<td>0.37</td>
</tr>
</tbody>
</table>

NB: No differences between the HEC and the UC781 exposed groups were seen. No differences in female to male cultures were seen.

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Changes in pre-defined rectal bacteria populations following cumulative exposure to 0.1% or 0.25% UC781 topical gel or HEC placebo compared to baseline.
HIV	extsubscript{BaL}, strain (10^4 \text{TCID}_{50} \text{ or } 10^2 \text{TCID}_{50}) \text{ TCID}_{50} \text{ was determined by titration using pooled PBMCs. The same viral stock used throughout the study. Explants were incubated with virus for 2 hours and then thoroughly washed and six explants from 10 cm and six explants from 30 cm were placed (one biopsy/raft) on gelfoam rafts in individual wells of a 24 well plate (Costar #3524, Corning, Inc., Corning, NY). Explants were followed for 14 days with supernatants for ELISA quantification of p24 (ng/ml) collected every 3–4 days and replaced with media (100%). Results are reported here as cumulative p24 detected at day 14 with ‘uninfecible’ defined as less than 100 ng p24/ml (p24 kits from AIDS & Cancer Virus Program, NCI, Bethesda, MD; detection to 78 ng/ml). At baseline (Visit 2), all participants’ HIV-exposed explant biopsies had a positive control of added UC781 to demonstrate suppressive potential.

The following samples were collected and assessed at Visit 2, 3 & 5:

- rectal microflora to assess changes with swabs collected at baseline and post 7-day exposure (only Visits 2 and 5)
- sloughing of rectal epithelial cells using endoscope-collected lavage at Visits 2, 3 & 5
- histopathology at 10 cm and 30 cm at Visits 2, 3, & 5
- fecal calprotectin to assess mucosal inflammation from post-encephal evacuation at Visits 2, 3 & 5
- mucosal mononuclear cell [MMC] phenotype by flow cytometry at Visits 2, 3 & 5
- secreted mucosal immunoglobulins collected by sponge (total IgG, total IgA) at Visits 2, 3-pre and 3-post & 5
- secreted mucosal cytokine proteins collected by sponge at Visits 2, 3-pre and 3-post product exposure & 5
- mucosal tissue mRNA cytokine profile at Visits 2, 3 & 5
- biopsies for ex vivo HIV-1 explant infection Visits 2, 3 & 5.

**Plasma levels of UC781**

Bridge Laboratories (Gaithersburg, MD) conducted FDA GCP compliant quantifications of plasma levels of UC781 within concentration ranges of 0.25 ng/ml to 200 ng/ml, using a validated LC-MS/MS system (lower limit of quantitation (LLOQ): 0.25 ng/ml). Samples evaluated included all 9 subjects (placebo and two UC781 concentration exposure groups) randomized to undergo the PK plasma sampling (baseline, 30 minutes, 2 hr, 4 hr, 24 hr and post 7-day exposure). Mucosal biopsies were obtained to quantify tissue levels of UC781 once the method had been optimized. The assay was recently validated and results obtained. These are being analyzed and will be reported in a subsequent paper.

**Statistical Analysis**

Evaluations of the different data collections were done as follow: Analysis of baseline variability among the three treatment groups was performed for continuous measure using analysis of variance and Kruskal-Wallis test. This exercise assured investigators that baseline ranges here were consistent with those previously published by this group in HPTN-056 [49]. Paired t-test and other appropriate tests were used to compare results from tissue samples collected at the 10 cm and 30 cm biopsy locations (histopathology, tissue cytokine mRNA, flow cytometry and ex vivo explant infections). To analyze the treatment effect of the UC781 gel, changes in mucosal indices from baseline to post-single exposure (V3) or post-7-day exposure (V5) were compared between each of the UC781 treatment groups and the HEC placebo group. A two sample t-test was used for continuous variables (tissue mRNA for cytokine, fecal calprotectin, flow cytometry). For continuous variables with two baseline measurements (rectal fluid cytokine profile, immunoglobulins), a mixed linear model with person-level random effect was used. For categorical variables (epithelial sloughing, histopathology), 95% confidence intervals for the difference in correlated proportions were calculated.

**Statistical Analysis of Ex Vivo Biopsy Infectibility**

**Suppressive impact of in vivo UC781 on ex vivo explant infectibility assays.** The aim of this explant study was to determine whether a topically administered topical product could suppress ex vivo HIV-1 infection in biopsies based on the ancillary study endpoint of cumulative p24 antigen at day 14 (already shown to tightly correlate with qRT-PCR for HIV) [64]. Comparisons evaluated the difference in tissue infectibility between baseline (Visit 2) and post-single dose exposure (Visit 3) or post-7-day exposure (Visit 5) in each study group using paired t-tests. Assessments of the treatment effect on altered infectibility from baseline (Visit 2) to post exposure single (Visit 3) or post-repeated (Visit 5) were conducted at each combination of biopsy locations (10 cm and 30 cm) and viral titers (10^2, 10^4) using two sample t-tests. Evaluation of potential baseline differences in biopsy infectibility (10 cm versus 30 cm) for each viral titer used was conducted by using paired t-tests. To determine whether treatment effect differs between the two biopsy locations, a mixed linear model evaluating differences in cumulative day-14 p24 antigen levels was fitted. This model incorporated person-level random effects to account for correlations among biopsies at 10 cm and 30 cm taken from the same individual. Four separate analyses were done, one for each combination of UC781 product (0.1%, 0.25%) and viral titer (10^2, 10^4). The mixed model afforded a number of comparisons, among which are a test for treatment effect (product vs. placebo), a test for biopsy location difference (10 cm vs. 30 cm), and a test for location x treatment interaction (addressing whether treatment varies by biopsy location). All biopsy infection results were independently confirmed by a secondary analysis using a comparison of 4 alternative analyses of explant readouts (day 14 p24, AUC, slope, and soft-endpoint analyses [65]).

**Results**

**Participant Demographics, High Study Retention and Sample Acquisition**

The 36 subjects completed the study in 16 months, each requiring a 7–14 week participation including 5 clinic visits, 3 visits for flexible sigmoidoscopies and sample collection, 2 ACASIs and 1 in-depth phone interview. Of 155 potential participants telephonically pre-screened, 55 were eligible for the screening visit (V1) with 36 participants meeting “enrollment” criteria for baseline sampling at Visit 2. All 36 participants initially enrolled completed the study (100% retention). The median age was 41 (range: 24–64) with 26 males (72%) and 10 females (28%). Participants were 41% African American, 39% Caucasian, 14% Hispanic, 3% American Indian and 3% Pacific Islander. Few differences between the participants randomized to each of the 3 study groups were appreciated (see Demographics Table S1).

**No Serious (Grade 3 or 4) Adverse Events (AEs) Reported**

Thirty-six AEs were reported of which 28 (78%) were Grade 1 and 8 (22%) were Grade 2. There were no Grade 3 or 4 AEs nor any procedure-related AEs (108 flexible sigmoidoscopies with just
over 3000 biopsies were performed in the 36 participants). Only AEs occurring following product exposure (Visit 3 forward) are reported; all are described in Adverse Events Table S2 by number/percentage of subjects having AEs, whether they were gastrointestinal (GI) related or not and which AEs occurred during which trial stage. Table S2 shows that the number of subjects with AEs of any grade are similar between groups after the single or 7-day exposure.

Nearly all the Grade 1 abnormalities (19/28; 68%) were recorded as not related to product. Of the 9/28 (32%) Grade 1 AEs reported as related to product, 6 were in the placebo group, 2 in the 0.1% UC781 group and 1 the 0.25% UC781 group. Eight Grade 2 AEs were reported among 5 subjects. Four of these 8 reports were from 1 individual (#401 - placebo group) who reported fever, cramps, flatulence, and diarrhea, considered possibly related, after the single exposure (Visit 3). Of note, no Grade 2 AEs were seen in this individual following the 7-day exposure. The other 4 Grade 2 AEs included the following limited diarrhea in 2 subjects following the 7-day exposure and considered possibly related (414 in the 0.25% UC781 group and 418 in the 0.1% UC781 group); transient thrombocytopenia in 1 subject following the 7-day exposure and considered not related (420 in the 0.25% UC781 group); and a spider bite in 1 subject after single dose and considered not related (4432 in the placebo group).

No Epithelial Sloughing from Rectal Lavage
No significant differences at baseline (Visit 2) were identified among enrolled subjects. No differences in the degree of epithelial sloughing between single exposure (Visit 3) and baseline or 7-day exposure (Visit 5) and baseline were appreciated. No UC781 treatment effect was identified for either low-concentration gel or high-concentration gel relative to placebo.

No Changes in Histology at 10 cm and 30 cm
Neither the placebo nor either of the two UC781 gel concentrations showed significant differences between either single or 7-day exposure and baseline. No differences were seen when each subject’s baseline samples at 10 cm and 30 cm were compared using this index of inflammation.

No Changes in Rectal Microflora
No significant changes in any of the rectal microflora profiles assessed were seen within subjects exposed to UC781 (either low or high dose) or the HEC placebo compared to their baseline culture results. As well, no significant differences were seen between study groups after cumulative (single and 7-day) exposures (Table 1). Although the study participant subpopulations were too small for statistical evaluation, no trends toward differences were seen between microflora samples from men compared to women (overall) or with between-group analyses of men compared to women, including Lactobacillus and Gardnerella vaginalis.

No Changes in Fecal Calprotectin
No significant differences in stool calprotectin, an indirect indicator of mucosal inflammation, were seen among study participants at baseline. No significant differences in calprotectin within subjects or between groups were appreciated following single-dose or after 7-day exposure (Table 2).

Minimal Changes in Mucosal Immunotoxicity Indices
The following data were studied at baseline and compared within and between groups after single-dose and after 7-day exposure: (i) phenotypic, HIV-1 co-receptor and activation profiles of mucosal CD4+ and CD8+ T lymphocytes at 10 and 30 cm; (ii) rectal fluid IgG and IgA immunoglobulins; (iii) rectal fluid cytokine protein yields; and (iv) tissue cytokine mRNA at 10 cm and 30 cm. These unique, baseline data from mucosal samples of healthy HIV-1 seronegative subjects will be enormously helpful to help power future study designs. Consequently, the detailed resultant means and standard deviations for each set of indices are added in Table S3.

Baseline variability was examined for all indices above. For flow cytometric analyses, no significant result was identified among 18 baseline comparisons performed. Importantly, the findings are consistent with those seen in the earlier HPTN-056 study [49], aimed at defining normative mucosal ranges for immunoinflammatory indices. However, using paired t-tests, baseline differences in lymphocyte surface phenotype and activation markers were seen between 10 cm and 30 cm, as we’ve reported previously [56,57]. Significant between-site differences at baseline (Visit 2) were identified at a level of \( \alpha = 0.05 \) for % CD4+ T cell lymphocytes, CD38 RFI on CD4+ T cell lymphocytes, CD38% on CD4+ T cell lymphocytes, HLA-DR% on CD4+ T cell lymphocytes, CCR5% on CD4+ T cell lymphocytes and % CCR5/CXCR4 double-positives on CD4+ T cell lymphocytes. These may reflect real differences since the total number of comparisons here was 9 [41].

Following intervention, before-after comparisons (baseline to single exposure or baseline to 7-day exposure) were conducted to evaluate changes in each of the above mucosal immune indices in each study group. From the 180 comparisons performed, 26 were significant at \( \alpha = 0.1 \). However, at \( \alpha = 0.05 \), 11 were significant (closer to the expected number of 9). There were no consistent patterns seen in the distribution of these 26 results. No evidence of consistent mucosal reaction to UC781 treatments or vehicle was identified following either single or 7-day exposure.

Table 2 presents the group analysis results of the clinician-delivered single-dose and self-administered 7-day exposure UC781 treatment effects compared to placebo on changes in all mucosal immune indices. Given the relatively small sample size and given the safety consideration of these first Phase 1 studies, we felt it important not to overlook any possible altered findings associated with exposure. Hence, those comparisons with p-values .10 and lower (corresponding to an alpha of 10\%), are highlighted in Table 2, which gives a higher power for identifying significant differences, at the cost of increased Type I error probabilities. For single-dose UC781 treatment effect compared to placebo, on changes in all mucosal immune indices (Visit 3-Visit 2) where 60 comparisons were performed, the single significant result (RANTES in the 0.1% UC781 vs. placebo comparison) is not unusual. The results support the conclusion that neither of the UC781 gel concentrations had any significant effect on mucosal immune indices compared to placebo. Importantly, when the HEC placebo responses were compared to baseline after single dose exposure (Visit 3 vs. Visit 2), no changes were identified. This is the first such detailed assessment of the HEC placebo on rectal mucosa (data not shown).

Following 7-day exposure (Visit 5-Visit 2), an additional 60 comparisons were compared to baseline values (Table 2). Seven significant results were identified at \( \alpha = 0.1 \), again, not an unusual number given the number of comparisons. The 4 findings that may have clinical relevance were the following but need to be kept in context with published trends toward differential expression at baseline [41]:

(i-ii) in the higher concentration UC781 group compared to placebo, a difference in the change of CCR5 expression was
seen on mucosal CD4+ T lymphocytes at both the 10 cm and 30 cm level (reduced; P = 0.025 for both)

(iii) for co-expressed CCR5/CXCR4 on CD4+ T lymphocytes at only 30 cm (reduced; p = 0.020)

(iv) for CXCR4 alone on CD4+ T lymphocytes only at 10 cm (reduced; p = 0.095).

It should remain clear that the selection of this broad profile of immunoinflammatory indices of mucosal injury represents a best effort to assess any changes of significance related to product (or placebo) in this vulnerable compartment with few standardized measures predicative of clinical toxicity. The overarching absence of changes or differences is relevant.

**Undetectable Plasma Levels of UC781**

No detectable levels of UC781 were seen by LC-MS/MS analysis in any samples (LLOQ in plasma: 0.25 ng/ml). As no product was identified in the 7-day exposed participants, the only confirmation of adherence to home use of the product was by participant-completed diary.

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**Table 2. Summary of p-values for comparing changes in mucosal immune parameters compared to baseline following single or 7-Day exposure.**

<table>
<thead>
<tr>
<th></th>
<th>Changes after single exposure (Visit 3-Visit 2) (p-value)</th>
<th>Changes after 7-day exposure (Visit 5 – Visit 2) (p-value)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>UC781 0.1% vs. Placebo</td>
<td>UC781 0.25% vs. Placebo</td>
</tr>
<tr>
<td>Fecal Calprotectin</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>0.417</td>
<td>0.587</td>
</tr>
<tr>
<td>Mucosal Immunoglobulins*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>IgG</td>
<td>0.297</td>
<td>0.777</td>
</tr>
<tr>
<td>IgA</td>
<td>0.754</td>
<td>0.458</td>
</tr>
<tr>
<td>Cytokine*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>RANTES</td>
<td>0.033***</td>
<td>0.569</td>
</tr>
<tr>
<td>MIP-1α</td>
<td>0.696</td>
<td>0.406</td>
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<td>TNF-α</td>
<td>0.436</td>
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<tr>
<td>IFN-γ</td>
<td>0.296</td>
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<td>IL-12 (p40)</td>
<td>0.488</td>
<td>0.428</td>
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<tr>
<td>IL-6</td>
<td>0.410</td>
<td>0.761</td>
</tr>
<tr>
<td>IL-1β**</td>
<td>0.347</td>
<td>0.896</td>
</tr>
<tr>
<td>Tissue mRNA for cytokine</td>
<td></td>
<td></td>
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<tr>
<td>IFN-γ</td>
<td>10 cm 0.407</td>
<td>0.546</td>
</tr>
<tr>
<td>IFN-γ</td>
<td>30 cm 0.999</td>
<td>0.303</td>
</tr>
<tr>
<td>Flow Cytometry: Mucosal Mononuclear Cells (MMCcs)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD4 lymphocytes</td>
<td>10 cm 0.561</td>
<td>0.806</td>
</tr>
<tr>
<td></td>
<td>30 cm 0.258</td>
<td>0.747</td>
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<tr>
<td>CD38+/HLA-DR+ on CD4</td>
<td>10 cm 0.467</td>
<td>0.404</td>
</tr>
<tr>
<td></td>
<td>30 cm 0.572</td>
<td>0.971</td>
</tr>
<tr>
<td>CD38 RFI on CD4</td>
<td>10 cm 0.877</td>
<td>0.815</td>
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<tr>
<td></td>
<td>30 cm 0.269</td>
<td>0.299</td>
</tr>
<tr>
<td>CD38% on CD4</td>
<td>10 cm 0.311</td>
<td>0.397</td>
</tr>
<tr>
<td></td>
<td>30 cm 0.698</td>
<td>0.369</td>
</tr>
<tr>
<td>HLA-DR1% on CD4</td>
<td>10 cm 0.307</td>
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</tr>
<tr>
<td></td>
<td>30 cm 0.419</td>
<td>0.749</td>
</tr>
<tr>
<td>CCR5 RFI on CD4</td>
<td>10 cm 0.401</td>
<td>0.103</td>
</tr>
<tr>
<td></td>
<td>30 cm 0.837</td>
<td>0.151</td>
</tr>
<tr>
<td>CCR5% on CD4</td>
<td>10 cm 0.702</td>
<td>0.668</td>
</tr>
<tr>
<td></td>
<td>30 cm 0.659</td>
<td>0.232</td>
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<tr>
<td>CXCR4% on CD4</td>
<td>10 cm 0.342</td>
<td>0.116</td>
</tr>
<tr>
<td></td>
<td>30 cm 0.511</td>
<td>0.211</td>
</tr>
<tr>
<td>CCR5%/CXCR4% on CD4</td>
<td>10 cm 0.546</td>
<td>0.372</td>
</tr>
<tr>
<td></td>
<td>30 cm 0.516</td>
<td>0.186</td>
</tr>
</tbody>
</table>

*For Mucosal Immunoglobulins and Cytokine, p-values were calculated based on two baseline measurements (Visit 2 and pre-Visit 3).

**One subject’s (ID = 417) baseline measurements were extreme outliers and excluded from analysis.

***Statistically significant at alpha level of 0.10.

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Biopsy Infection studies: Successful Suppression of ex vivo HIV-1 Infection of Colorectal Biopsies Exposed in vivo to UC781

Characteristics of biopsy infections at baseline (Visit 2). All 36 participants had baseline colonic biopsies obtained from 10 cm and 30 cm immediately set up for ex vivo HIV-1 infection with R5 HIV_HBXL at two titers (10^4 TCID_50 and 10^5 TCID_50). All but one participants’ biopsies were infectible at baseline with the higher 10^5 TCID_50 viral titer. Only ~60% of participants’ biopsies were infectible at baseline with the lower 10^4 TCID_50 titer. The single individual with biopsies not infectible at baseline with either titer (subject #439) demonstrated normal ranges of surface CCR5 expression on mucosal CD4 T lymphocytes by flow cytometry.

The ~40% of participants with uninfected biopsies at baseline with the lower TCID_50 10^5 titer were not the same ~40% with uninfected biopsies at subsequent visits. Over the course of the trial, evaluating only the placebo group (n = 12), as at baseline, all were infectible at all visits with the higher 10^5 TCID_50 titer. With the lower 10^5 viral titer, the 33.3% (subjects #401, 407, 432, 445) not infectible at baseline (Visit 2) was a different 33.3% subset than those not infectible at Visit 3 (subjects #426, 432, 442, 455). At Visit 5, 66.7% of placebo-treated explants were not infectible with the lower viral titer (subjects #407, 416, 417, 426, 432, 442, 445, 455). With the lower viral titer, a participant’s biopsies being uninfected at one visit did not predict uninfectibility at later visits.

To determine whether biopsy infectibility ex vivo was impacted by location (10 cm versus 30 cm), baseline p24 infection levels for each titer of virus was compared, using paired t-tests. No evidence for differences in explant infectibility ex vivo was seen between two biopsy locations of 10 and 30 cm. These data support using a single site in future similarly designed trials.

Single-Dose Exposure: Ex vivo infections were suppressible following single-dose in vivo product exposure. Data are shown in Figure 2 for the within-group suppression of cumulative p24 following ex vivo infection of participants’ biopsies when exposed in vivo to the single dose of the topical product (Visit 3 compared to Visit 2). These products were directly applied by study clinicians. In the 0.25% UC781 treatment group, the sharp decreases in viral infections from Visit 2 to Visit 3 are striking. This decrease is consistent across both virus titer levels and both biopsy locations.

The viral suppression shown in Figure 2 (right column) using the 0.25% UC781 formulation is significant at p < 0.05, based on two-sided paired t-tests for the four separate before-after comparisons (10^5 virus at 10 cm, 10^5 virus at 30 cm, 10^4 virus at 10 cm, 10^4 virus at 30 cm) (p < 0.02). The same paired before-after comparisons for the 0.1% UC781 formulation demonstrate a trend toward suppression but with marginally significant or non-significant differences (10^5 virus: p = 0.59 at 10 cm, p = 0.038 at 30 cm; for 10^4 virus: p = 0.051 at 10 cm, p = 0.035 at 30 cm). None of the placebo comparisons showed a significant change between Visit 2 and Visit 3.

Single exposure: Between group differences in biopsy viral suppression. The effect of 0.25% UC781 formulation relative to placebo was significant for 10^4 TCID_50 at 10 cm (p = 0.002) and close to significant for 10^5 TCID_50 at 30 cm (p = 0.059) at the significance level of p = 0.05. For 0.1% UC781, placebo demonstrated a significant change between Visit 2 and Visit 5 under any combination of virus titers and biopsy locations. 7-day exposure: Between group differences in explant viral suppression. Between-group reductions were also not seen following the 7-day exposure group comparisons.

Mixed model analyses showed “treatment,” not biopsy location, was responsible for the observed significant effects. This approach addressed whether differences in the cumulative p24 between baseline (Visit 2) and post-single exposure (Visit 3) were primarily related to treatment and/or biopsy location. The treatment effect was highly significant for the 0.25% UC781 in vivo treated biopsies exposed ex vivo to 10^4 TCID_50 virus (p < 0.001). The tests for biopsy location as well as location×treatment interaction were not significant (p = 0.65 and p = 0.40, respectively). The other three analyses (0.1% formulation with 10^5 TCID_50, 0.1% formulation with 10^4 TCID_50, 0.25% formulation with 10^2 TCID_50) also did not show any significant treatment, location or treatment×location effects.

Discussion

There are five main findings in this first-in-field, Phase 1, randomized, double-blind, placebo-controlled trial of a vaginally formulated HIV-1 microbicide used rectally in 36 men and women: (i) The results showed both doses of the vaginally-formulated UC781 gel used rectally are safe by every index used and are highly acceptable to participants [50]; (ii) The design of the trial was innovative by including two separate exposures (single and 7-day) as independent but linked components to expedite safety assessments; (iii) A novel profile of diverse mucosal “immunotoxicity” indices was included to evaluate potential changes in colorectal mucosa following product exposure, some of which might later point to increased risk of HIV infection; (iv) The immunotoxicity results here correlated well with previously published data from HPTN 056 [49], designed specifically to identify mucosal markers of interest and their stability over 6 weeks; (v) The novel Phase 1 inclusion of ex vivo biopsy infectibility studies based on in vivo exposed tissue samples was demonstrated as a robust, clinically-relevant, potential early bio-indicator of efficacy. After this trial was completed, it was decided that further development of UC781 would not be pursued because of problems with solubility, stability and other reasons unrelated to this trial. However, findings (ii) through (v) above represent significant advances in the field of microbicide development independent of the product studied.

Colorectal mucosa is composed of a single layer of columnar epithelial cells, extremely receptive to injury but capable of rapid repair. It is highly vulnerable to HIV-1 infection [7,22], related to the subjacent lamina propria which, in health, is densely populated with activated memory T-cells expressing both CD4 and both HIV-1 co-receptors CCR5 and CXCR4, as well as dendritic cells (DCs) and macrophages [6,7,22,66,67]. The physiologically inflamed tissue is more infectible per sex act (possibly by 20 to 2000-fold depending on co-STI infections, inflammation, etc) than is the vaginal mucosa [68–70]. These differences may also increase rectal compared to vaginal susceptibility to microbicide-induced toxicity, potentially favoring HIV infection [16] as seen with other sexually transmitted infections [31,33,34]. The rapidly reactive rectal tissue responses make selecting sample study points difficult, as evidenced by the differing reports and clinical implications seen with N9. Tabet et al. in evaluating an approved vaginal formulation of N9 in MSM [16], described mild to no rectal histological changes in participants receiving up to 6 weeks of daily N9 (or placebo gel) with samples collected up to 12 hours after N9.
exposure [15]. In contrast, Phillips et al. using rectal lavage and histology endpoints, saw marked rectal epithelial exfoliation 15 minutes after a single N9 topical exposure in men which disappeared after 2 hours [14].

In this Phase 1 trial, the rectally-applied, vaginally-formulated UC781 0.1% and 0.25% gel and the HEC placebo were safe. There were no Grade 3 or higher AEs reported nor any procedure-related AEs despite over 3000 biopsies having been performed. Retention was 100% post-enrollment, emphasizing participant’s understanding and willingness to undergo these procedures repeatedly.

Acceptability (using web-based surveys and an in-depth interview at study’s end) was high, especially when focusing on “likelihood” to use the product in the future [50]. The importance of determining acceptability in the early phases of microbicide development is critical, enabling earlier, less costly product changes and raising the likelihood of real-life use.

The “immunotoxicity assays” as a whole, showed no significant changes from baseline for each product or any significant differences between study groups. This was true for both the 10 cm and 30 cm assessments following the single-dose and the 7-day dose exposures. It is reassuring that the baseline results here were generally within the normative ranges reported in HPTN 056 [49]. This confirmation is important to the field.

One of the most impactful endpoints in this Phase 1 trial was the first-time attempt to infect ex vivo tissue biopsies that had been exposed to drug only in vivo. Given limits on the number of biopsies obtainable, only the laboratory strain of R5 HIV-1_RbaL was used at two titers (TCID50 of $10^7$ and $10^9$ using the same lot of virus). These titers are thought to be far in excess of ejaculate viral titer (closer to clinically relevant titers) could be a statistically significant, even with these small subject numbers. This was seen only with the higher viral titer. The data are compelling when evaluating the 30 minute topical exposure of the higher UC781 concentration (0.25% gel) at 10 cm (area most likely drug exposed) using the higher viral titer (assuring baseline infectibility). UC781 is a potent inhibitor of viral RT. For this to be the main mediator of suppression in these heavily washed biopsies, other mechanisms may need to be invoked and tested, including an intracellular activity and/or retained drug in tissue/membranes due to the lipophilic nature of UC781 inhibiting further replication. This latter observation is the known (but poorly understood) “memory” effect of UC781. A clinically relevant advantage may be prolonged resistance to transmission due to this feature of UC781, possibly allowing for coitally-dissociated administration.

These results showing suppression of ex vivo biopsy infection were consistent and reproducible, within groups and between groups. Given that these are clinical samples with expected inter-subject variability and washed frequently by shaking in media and during transport from the endoscopic unit to the laboratory, it is remarkable that any ex vivo impact was seen with only 12 participants in each study group. Additional confounders (that might have taken years to segregate out) include in vivo variables such as the biological activity of the colorectal area (continuous mucus secretion, peristaltic activity, frequent stool passage) and the absence of an independent, visual identifier to guide biopsy acquisition relative to where the investigator-administered topical drug might be located 30 minutes after exposure.

Evaluating the 7-day (uncontrolled) exposure

As no independent, reliable biological indicator of home delivery/compliance was used (a great need in this field), there are many contributory explanations for the apparent absence of ex vivo suppression of biopsy infections. Maintained suppression would be dependent on the subjects using the product daily, as instructed, and that there be some tissue retention, given the active GI elimination processes. In addition, some of the biochemical, lipophilic features of UC781 as well as its tissue half-life in vivo remain undefined. When coupled with complex mucosal factors of mucus, microflora and dynamic epithelial turnover, the absence of 7-day exposed ex vivo suppression is more understandable and points out challenges to address in future trials.

At this point, the ex vivo biopsy infection suppression assays may be viewed as a possible bioindicator of pre-clinical efficacy. Importantly for future trials: (i) no differences were seen using the 30 cm or the 10 cm biopsies, supporting the selection of a single anatomical site in similarly designed trials (benefiting participant safety as well) and (ii) using a sufficiently high titer of
standardized virus minimizes data loss due to uninfectible baseline samples.

Conclusions
The microbicide field has met with significant difficulties in attempting to move a product from design through development to demonstrating efficacy. The first hopeful signs were the dramatic results from CAPRISA 004 [10]. While not discussed much 10 years ago, awareness of the high prevalence of receptive anal intercourse (RAI) in heterosexual partnering [24–30,74–80] as well as MSM, combined with the enhanced infectibility per sex act of this compartment has driven the microbicide field to actively include development, assessment and acceptability of rectal microbicides into the main portfolio of scientific and public health goals.

Many advances in technology, immunotoxicity assessments, PK and pharmacodynamic (PD) study design, formulations and acceptability measures have enabled the development of more strategic and focused efforts with rectal microbicides [41,46,47,49,54,63,64,61–65], supporting a rapid advance to clinical trials. While development of a combination microbicide useable in both sexual compartments is a long-term desired goal, the current need for a safe and effective rectal microbicide is evident. The inclusion of a novel ex vivo biomarker of efficacy in a Phase 1 trial is a boon to the field and may turn out to be an exciting addition as a “nonclinical surrogate of bio-efficacy,” helping guide decisions about which agents at which concentrations should be advanced in human trials. These intensive efforts and mucosal indices benefit the microbicide field as well as the growing focus on mucosal vaccines, enabling a more rational and cost-effective product development pipeline.

Supporting Information
Table S1 Demographics.
Table S2 Adverse Events.

References


