

**Impact of Cryotherapy versus Loop
Electrosurgical Excision Procedure (LEEP)
on Recurrence of Cervical Intraepithelial
Neoplasia and HIV-1 Cervical Shedding
among HIV-positive Women**

Study Protocol

Version 8.5

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University of Washington

November 20, 2015

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Study Investigators

Principal investigator

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Dr. Chung is the PI and will directly lead the planning, implementation, and analysis of the study. He will meet weekly with the study team, direct the study, and guide the study coordinator. He will check progress on enrollment and follow-up with the study coordinator and study doctor. In addition, Dr. Chung will serve as the point person to explain the study and share data results with others and report adverse effects associated with the study. The University of Washington will provide administrative, laboratory and data support to this project.

Co-investigators

Nelly Mugo, MBChB, MMed, MPH, Gynaecologist, Department of Obstetrics and Gynecology, Kenyatta National Hospital

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Dr. Mugo is a Co-investigator of the study and will assist Dr. Chung in the study's planning and implementation. As a gynaecologist, she will ensure that the study medical staff will provide excellent care and maintain high clinical standards. She will oversee the gynecological care and proper medical procedures by meeting regularly with the study doctor

and nurses. Dr. Mugo will be involved in any gynecological complications related to the study.

Samah Rafie Sakir, MBChB, Medical Director, Coptic Hospital of Kenya

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As Medical Director of the Coptic Hospital, Dr. Sakir will work with Drs. Chung and Mugo to implement the study at the Coptic Hope Center for Infectious Diseases. He and the Coptic Hospital will provide the clinical infrastructure where HIV patients will be enrolled and followed in the study. Dr. Sakir will manage the health care workers at the Hope Center and ensure that clinical data that is collected from the Hope Center and shared with the study is accurate and timely.

Grace John-Stewart, MD, PhD, Professor, Department of Medicine, University of Washington

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Dr. John-Stewart will lend her epidemiology expertise to the analysis of the study. She has significant experience in conducting and examining randomized clinical trials in Kenya. Dr. John-Stewart will help analyze the data, prepare any manuscripts, and give feedback on implementation of the trial.

Barbra Richardson, PhD, Research Professor, Department of Biostatistics, University of Washington

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Dr. Richardson will contribute statistical support to the study and will be deeply involved in statistical analysis of its findings. She will prepare the method to randomize subjects and will analyze results of the study.

Dr. Hugo De Vuyst, MD, PhD, epidemiologist, Infections and Cancer Epidemiology, International Agency for Research on Cancer (IARC-WHO).

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Dr. De Vuyst will contribute his expertise and epidemiological skills in issues of cervical cancer screening, HPV and HIV in developing countries. He will help analyze the data and its association with HPV results.

Silvia Franceschi, MD, Epidemiologist, Head of Infections and Cancer Epidemiology Group, IARC-WHO

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Dr. Franceschi will contribute her extensive epidemiological expertise in the field of cervical cancer, HPV and HIV.

Martin Steinau, PhD, Team Lead HPV DNA, Chronic Viral Diseases Branch (CVDB), Division of High-Consequence Pathogens and Pathology (DHCPP), National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC) 1600 Clifton Road, MS G41, Atlanta, GA 30329-4027

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Dr. Steinau will coordinate the HPV-related study activities and data-analysis. He will contribute his expertise and epidemiological skills in issues of cervical cancer screening, HPV and HIV in developing countries. He will oversee the testing of HPV samples at CDC, ensure quality control, and help analyze the data and its association with HPV results.

Elizabeth R. Unger, PhD, MD, Chief CVDB, DHCPP, NCEZID, CDC

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Dr. Unger will contribute her expertise in the field of cervical cancer, HPV and HIV. She will supervise and manage the laboratory where the HPV sample testing will occur and help analyze the results of the study.

Nelly Yatich, DrPH, MPH, Clinical Assistant Professor, Department of Global Health, University of Washington

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Dr. Yatich will work closely with Drs. Chung and Mugo to implement the study at the Coptic Hope Center. She will provide mentorship to the research team at weekly meetings, provide mentorship to the Data Manager in data analysis, and guide the study coordinator. She will provide other on the ground support as needed.

Dara A. Lehman, MHS, PhD, Staff Scientist, Human Biology, Fred Hutchinson Cancer Research Center
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Dr. Lehman will lead efforts to quantify the HIV-1 RNA levels of cervical and plasma samples collected from subjects enrolled in the study comparing cervical cancer treatments in HIV-positive women.

Summary and Objectives

The recent scale-up of antiretroviral treatment programs in resource-limited settings provides an unprecedented opportunity to implement a comprehensive cervical cancer screening and treatment program for women who, by virtue of having HIV, are at significant risk for cervical disease. Unfortunately, even if screening is offered free of charge to millions of women living with HIV, it is unclear which treatment modality for pre-cancerous cervical lesions will be most effective since HIV appears to affect outcomes of treatment by increasing the recurrence and severity of cervical disease. Cervical treatment may also increase shedding of HIV from the cervix which may put discordant couples at risk and possibly spread HIV more widely. This study proposes to randomize HIV-positive women with cervical intraepithelial neoplasia grade 2 and 3 (CIN 2 and 3) to cryotherapy vs. loop electrosurgical excision procedure (LEEP) and measure the recurrence of cervical disease in each group over 2-years of follow-up as well as HIV shedding from the cervix for 3 weeks after treatment.

Our hypothesis is that compared to cryotherapy, LEEP is significantly more likely to prevent recurrence of cervical lesions over 2 years of follow-up and less likely to cause shedding of HIV-1 from the cervix over 3 weeks of follow-up.

The objectives of this study are:

1. To compare the rate of recurrence of cervical intraepithelial neoplasia among HIV-positive women receiving cryotherapy versus LEEP over 2 years of follow-up
2. To compare the shedding of HIV-1 from the cervix between HIV-positive women receiving cryotherapy versus LEEP over 3 weeks of follow-up

Background

The introduction of antiretroviral medications on a large-scale in resource-limited settings through funding from agencies such as the President’s Emergency Plan for AIDS Relief (PEPFAR) has decreased the number of HIV-positive women dying from AIDS. As a result, many HIV-positive women are leading longer, healthier lives. However, despite immune reconstitution many are still at risk for diseases related to their HIV infection including cervical cancer.¹ Cervical cancer is the leading cause of cancer death among women in resource-limited settings, and HIV-positive women are more likely to be infected with human papillomavirus (HPV), the primary cause of cervical cancer, and progress to invasive, life-threatening disease than those who are HIV-negative.²⁻⁶ Thus, although many women may be saved by antiretroviral therapy through PEPFAR support, they may later die of a disease that could have been detected and prevented at the same facilities where they received their antiretroviral treatment.⁷

The importance of adequate cervical cancer screening among HIV-positive women is being recognized by the Kenya PEPFAR program, Office of the Global AIDS Coordinator (OGAC), and other clinics around the world which are treating HIV-positive women.⁸ While there is a body of published knowledge on the screening and treatment of women in resource-limited settings, very little has been studied on the relevance of these findings on HIV-positive women.^{9, 10} For example, it has been suggested that visual inspection with acetic acid (VIA) along with cryotherapy be recommended as a “screen and treat” approach on the same day for women located in resource-limited settings.^{11, 12} The benefits are obvious; VIA is simpler to administer than a Papanicolaou test (Pap smear), does not require laboratory support, and is up to 20 times less expensive. Similarly, cryotherapy, a low technology treatment option, can be offered on the same day as VIA decreasing loss-to-

follow-up due to referral and waiting times, and is cheaper and easier to administer than LEEP. As a result, some HIV treatment programs in resource-limited settings are beginning to utilize this approach for their female patients. As PEPFAR begins to consider supporting cervical cancer screening among HIV-positive women enrolled in its programs, it will be essential that decisions are grounded in scientific evidence since any approach may have tremendous consequences on morbidity, mortality, and transmission of HIV-1.¹³

The issue is that there is no evidence that a “screen and treat” approach is as effective among HIV-positive women as it appears to be among HIV-negative women. In fact, there is data to suggest that this approach may be problematic. In HIV-negative women, VIA appears to be more sensitive but less specific compared to Pap smear.^{9, 14} HIV-infected women have a higher prevalence of aggressive cervical disease and are more likely to experience recurrent HPV and genital infections.^{5, 15} The presence of florid disease may alter the sensitivity and specificity of VIA in the presence of HIV disease, making it more sensitive but less specific than Pap smear. VIA may therefore detect more cervical abnormalities in HIV-infected women that are not truly cancerous. Coupled with cryotherapy, this may result in many HIV-positive women receiving unnecessary treatments that inflame the cervix and cause it to shed increased levels of HIV virus.¹⁶ Increasing cervical shedding of HIV after cryotherapy may increase HIV transmission and infectivity in a manner analogous to male circumcision which appears to increase the risk of female partners acquiring the disease.¹⁷

The standard of care for screening and treatment in the US, according to the American Society for Colposcopy and Cervical Pathology (ASCCP), is based on Pap smears and excisional (cold-knife conization, loop electrosurgical excision procedures, laser conization, and electrosurgical needle conization) or ablative treatments (cryotherapy, laser ablation, electrofulguration, or cold coagulation).¹⁸ Women who are found to have high-

grade lesions on cytology may either have their lesion treated right away with LEEP or have a colposcopy-directed biopsy. If the woman has a biopsy and the histology results show a CIN 2 or 3 and the colposcopy is satisfactory, treatment may be undertaken with either ablative or excisional therapy. If colposcopy is unsatisfactory or the CIN 2/3 is recurrent, treatment should be a diagnostic excisional procedure, which is an excisional procedure followed by a pathological examination of the sample tissue.

According to the Kenyan Ministry of Health, cervical cancer screening and treatment practices include VIA and cryotherapy at the district level health centers and below, and Pap smears with follow-up colposcopy with biopsy and LEEP at tertiary and provincial level hospitals.¹⁹ In our study, participants will be screened using Pap smear with confirmatory histology and treated with cryotherapy or LEEP. Our screening and treatment methods are consistent with standards of care set at Kenyan tertiary and provincial level facilities. As mentioned above, one of the reasons to study cryotherapy and LEEP is to understand how to refer HIV-positive women for cervical treatment within the Kenyan government health system. The reason we are utilizing cytology and histology as a screening method is the lack of evidence confirming the sensitivity and specificity of VIA among HIV-positive women. Given that the outcome of measurement in this study is recurrence of CIN, it is considered scientifically necessary and within Kenyan standards of care to use these accurate, evidence-based tests.

In terms of risk of serious complication, cryotherapy and LEEP are quite safe. In a study from Zambia, Pfaendler, et al. found that the overall complication rate of LEEP to be 3.7%, all of which was managed on-site in the clinic.²⁰ Likewise, in a study of cryotherapy in India, the overall complication rate was found to be 3.0%.²¹ In a large study from Peru that followed 1,398 women, who underwent cryotherapy for a mean of 12 months, no serious complications, including pelvic inflammatory disease, severe cramps or bleeding, or

anaphylactic reactions, were found.²² In a comparison study between cryotherapy and LEEP, cryotherapy was found to have a 2% complication rate and LEEP, an 8% complication rate, and the difference was not significant.²³

There has been some controversy surrounding cervical treatment and whether or not it is associated with adverse pregnancy outcomes. Sadler et al. showed in their retrospective analysis of 652 women that had undergone LEEP, laser ablation or laser conization that LEEP did not increase the incidence of preterm delivery.²⁴ However, the authors did note a significant increase in premature rupture of membranes. Acharya et al., in their matched cohort of 428 women undergoing LEEP, also found no correlation between the procedure and premature delivery or low birth weight, but they did find a significantly higher number of women with pregnancy complications, which included premature contractions, infections and cervical incompetence.²⁵

Finally, there is evidence that cryotherapy may be less effective compared to LEEP in preventing the recurrence of cervical intraepithelial neoplasia though the literature is equivocal. Overall, there have been few studies comparing the efficacy in treatment between cryotherapy and LEEP, especially in HIV-infected women. Chirenje et al. found a significant difference in the failure rate of cryotherapy versus LEEP, however his numbers were small with only 6 high-grade recurrences in the cryotherapy arm (14.3%) and 2 in the LEEP arm (4%).²⁶ Moreover, neither HIV nor HPV shedding was measured and follow-up time was only one year. In another randomized study comparing cryotherapy and LEEP, this time a larger study in non-HIV-infected women, no significant difference was found between the two arms in terms of failure, defined as either recurrence or persistence.²³ Additionally, in a 2000 Cochrane review, it was stated that “evidence suggests that there is no obviously superior surgical technique for treating cervical intra-epithelial neoplasia.”²⁷

If cryotherapy is found to result in a greater number of failures, it may require more frequent and careful follow-up screening than LEEP, and therefore may not be as cost-effective or therapeutic for the patient. As a result, the individual and public health risks of a “screen and treat” approach for cervical cancer screening and treatment among HIV-positive women may be much greater than its benefits.

Rationale

The University of Washington (UW) in collaboration with the Coptic Hope Center for Infectious Diseases has been providing cervical cancer screening to its female HIV positive patients in Kenya since 2006. The UW/Coptic Hope Center has enrolled over 8,000 HIV-positive women in its two Nairobi sites and offers a robust patient population for cervical screening. The UW/Coptic Hope Center has already screened over 2,000 HIV-positive women for cervical cancer using both Pap smear and VIA, and has worked in partnership with Kenyatta National Hospital (KNH) to provide LEEP to those with detectable lesions. Most recently, the collaboration has received a grant from the Puget Sound Partners for Global Health to compare VIA versus Pap smear among women enrolled at the Hope Center and to examine HIV-1 cervical shedding in a small subset who receive cryotherapy. Unfortunately, funding is only available for one year and will not allow any comparison with LEEP or a study of cervical disease recurrence after intervention.

Kenya is an appropriate site to conduct this study due to a high incidence of cervical cancer and lack of cervical screening coverage. The incidence of cervical cancer in Kenya is much higher compared to the West and measures between 43 and 45 cases per 100,000 compared to 8.4/100,000 in the USA.^{28, 29} Of 3,902 women who presented to KNH with reproductive tract malignancies between 1989 and 1998, 85% had invasive cervical cancer.²⁸ In a sampling of 1,353 patients at the same institution in Kenya, only 22% reported having received a previous Pap smear.³⁰ Although the government of Kenya has advocated the use of VIA as a primary method to screen for cervical cancer, no specific recommendations are made concerning HIV-positive patients. Cervical cancer screening for HIV-positive women in Kenya should be a high priority since HIV-infected women in Kenya with invasive cervical cancer are 10 years younger than HIV-negative women at initial presentation.^{6, 31}

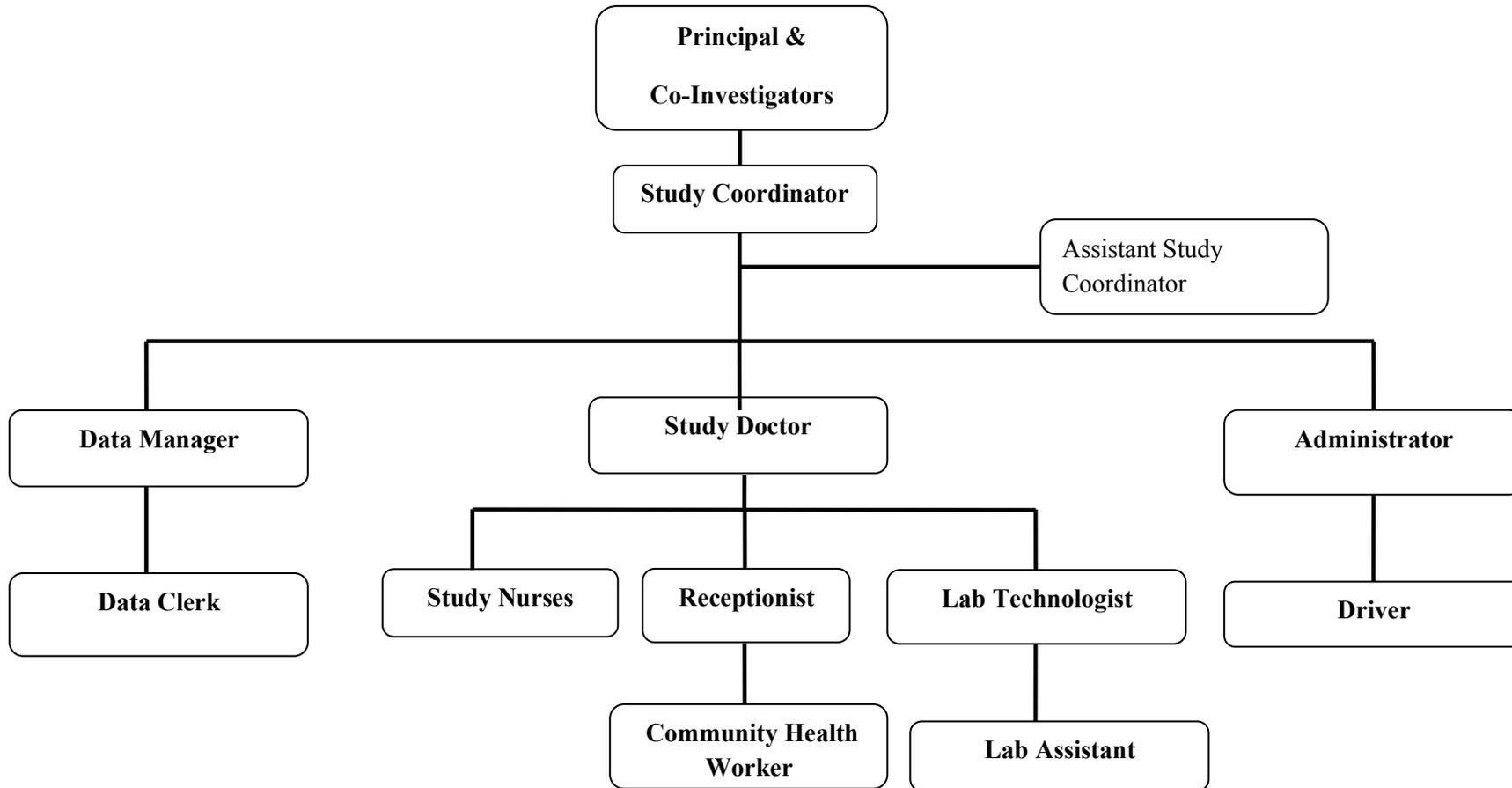
Based on the high incidence of cervical cancer in Kenya and building upon our programmatic and research work in cervical cancer screening among HIV-positive women, we propose to study the effect of LEEP versus cryotherapy on the recurrence of cervical intraepithelial neoplasia and the shedding of HIV-1 from the cervix. Our hypothesis is that compared to cryotherapy, LEEP is significantly more likely to prevent recurrence of cervical lesions over 2 years of follow-up and less likely to shed HIV-1 from the cervix over 6-weeks of follow-up. This evaluation will inform PEPFAR policies on the best method to treat pre-cancerous lesions in HIV-positive women and elucidate the importance of cervical treatment interventions according to immune status and antiretroviral therapy. Such information is directly relevant to the care of HIV-positive women in Kenya and other resource-limited countries which are significantly impacted by cervical cancer.

Timeline and Dissemination

The duration of the evaluation activity is approximately 6 years from the beginning of the funding cycle. It is estimated that it will take at least 3 years to screen at least 2,400 women and randomize approximately 400 women with high grade intraepithelial lesions to LEEP vs. cryotherapy. These randomized participants will be followed for 2 years after screening or randomization. Including preparation and analysis, it is expected that it will take approximately 6 years to complete this study.

Results of the study will be shared directly with the USG/GOK technical teams through annual reports and regular e-mail contact with designated contacts at the CDC in Atlanta and Nairobi. Reports will include analysis from regular DSMB meetings. After initiation of the study, we will confer with the CDC on whom to report to in Kenya and Atlanta. At that time, we will also determine how frequently the USG/GOK technical teams would like to be appraised of the study and its results. At a minimum, we will seek to be in phone and/or e-mail contact with USG contacts quarterly to ensure that the study and its data is relevant to USG technical working groups and policies established around cervical cancer screening in PEPFAR-supported clinics. Dissemination of study findings will also occur through public presentations and publication in internationally recognized journals. CDC and USG staff/agents will not participate in the study as co-investigators or study collaborators. They will not participate directly in the study development, analysis, or manuscript preparation.

Cervical Screening Organogram



Personnel

Study Coordinator

- Manage directly the Study Monitor, Data Manager, Study Doctor, Administrator, and Lab technologist around aspects of the PHE cervical treatment study and R01 resistance study
- Monitor the progress of research activities and ensure the smooth and efficient day-to-day operation of research and data collection activities
- Initiate and coordinate activities that improve the conduct and performance of the study
- Conduct weekly clinic meetings that are led by Study Doctor with study staff
- Conduct weekly data meetings that are led by the Data Manager to ensure data is collected in a timely fashion, is cleaned, and clearly analyzed
- Conduct and lead weekly study-related administrative meetings
- Direct and be responsible for the study budget and petty cash that is managed by the administrator
- Act as the primary administrative point of contact for research staff and as the principle operational liaison for Coptic administration and regulatory bodies
- Analyze recruitment rates, determine if rates match expectations, and implement plans that will promote recruitment
- Analyze retention rates and implement plans that will promote retention
- Supervise and coordinate the provision of support services to investigators
- Prepare periodic and ad hoc reports as required by investigators, funding agency, and/or regulatory bodies

- Be responsible for renewing, updating and modifying IRB applications at UW and KNH that are associated with this study
- Be responsible for generating Adverse Events, protocol violations and deviations, and unanticipated problems reports
- Be responsible for liaising with the DMSB as needed
- Perform any other duties and responsibilities that may be given by the PI or co-investigator

Assistant Study Coordinator

- Assist the Study Coordinator in monitoring the progress of research activities and ensuring the smooth and efficient day-to-day operation of research and data collection activities
- Implement and be responsible for renewing, updating, and modifying existing standard operating procedures (SOPs) and develop new ones as needed
- Be responsible to ensure that all staff are following SOPs
- Implement quality control procedures throughout the conduct of the study
- Review the accuracy, completeness and timeliness of completed study related records, case report forms and other documents
- Compare reported data with original source documents
- Review study related processes relative to applicable regulatory requirements, including GCP and Human Subjects Protection regulations
- Verify the following items for the study: protocol compliance (i.e. subject recruitment and eligibility criteria, informed consent and randomization procedures); that only designated

investigator(s) and/or appropriate research staff are performing study functions; that regulatory compliance is being maintained (i.e. that investigators are providing and maintaining all study related documents as required.) Be responsible for study compliance with all regulations

- Communicate any serious deficiencies noted during monitoring to the Study Coordinator
- Ensure that a record of all correspondence, monitoring reports and other written documentations are maintained by the Administrator
- Participate in all study meetings
- Organize and coordinate all training activities
- Respond to and be responsible for implementing all matters that may arise from CDC and Study Monitor visits
- Perform any other duties and responsibilities that may be given by the Study Coordinator
-

Study Doctor

- Oversee and ensure that patients in the study are receiving good medical HIV care and inform the study coordinator of any complications
- Identify subjects that require medical attention and refer them for care at the Hope Center
- Draw blood or obtain specimens from patients if the lab assistant or lab technologist is unable or unavailable
- Oversee the cervical screening clinic and ensure it is well-stocked with necessary medical supplies and equipment to perform the study

- Meet with the data clerk or data manager to correct data entry errors
- Review and confirm eligibility of each patient for study
- Administer questionnaires
- Conduct gynecological examinations, HIV and HPV swabs, colposcopy, biopsies, and cryotherapy and LEEP
- Assist the study nurses in performing their duties if they are unable or unavailable
- Confer and communicate with Hope Center clinicians and medical staff if any questions or problems arise concerning medically related issues
- Work closely with clinic staff at the Hope Center to ensure high recruitment for research study
- Work with data manager and receptionist to analyze data and prepare reports
- Present weekly summaries along with the study nurse and receptionist marking progress in enrollment and tracking of subjects in the study
- Will be responsible for reporting adverse effects to the principal investigator and co-investigators
- Perform any other duties and responsibilities that may be given by the principal investigator or the study coordinator

Study Nurses

- Conduct gynecological examinations, HPV swabs, and Pap smears

- Check age, previous cervical screenings and gynecological history before enrollment into the program
- Provide adequate knowledge and education about the study to patients so they can sign an Informed Consent
- Administer the Informed Consent and store it safely
- Administer questionnaires
- Review and confirm eligibility of each patient for research study
- Assist the study doctor in any medical procedures
- Attend weekly clinic meetings
- Maintain, and in the absence of the community health worker, clean and organize the cervical screening clinic
- Transport equipment and supplies for cleaning and autoclaving
- Meet with the data clerk or data manager to correct data entry errors
- Draw blood or obtain specimens from patients if the lab assistant or lab technologist is unable or unavailable
- Perform any other duties and responsibilities that may be given by the study coordinator or study doctor

Administrator

- Manage petty cash and study budget
- Liaise with payroll administrators to ensure salaries are paid correctly and on time

- Make purchases, photocopy data collection tools, and keep inventories of supplies
- Maintain communication between the clinic and office
- Attend and take minutes at weekly administrative and clinic meetings and present them at the next meeting
- Reconcile receipts to send to Seattle
- Communicate with Seattle when more funds are needed
- Manage the driver and arrange transportation
- Arrange for study trainings in coordination with study coordinator
- Maintain and organize files of personnel, correspondence, applications, IRB records, receipts, budget, inventories, etc...
- Coordinate staff evaluation procedures
- Oversee and record the attendance of office and clinic staff in coordination with the study nurse
- Make monthly reports of project expenses
- Facilitate in renewing personnel medical insurance and liaise between insurance and the hospital of matters of personnel appointment
- Make weekly reports of administrative issues
- Prepare IRB and government applications for the shipment of samples
- Remind study coordinator when IRB renewals are due and work with study coordinator and principal investigator to submit, modify, and renew IRB applications
- Ensure office tidiness

- Perform any other duties and responsibilities that may be given by the principal investigator or study coordinator

Receptionist

- Follow subjects enrolled in the study and ensure they are retained in the study and proper follow-up is done both at the research clinic and the Hope clinic
- Keep track of all the patients enrolled and determine if any patient has missed appointment and take action to report and bring these patients back under care and supervision
- Handle money given by the administrator and account for it by keeping the various logs (i.e. calling log, transport log and client transport reimbursement forms) and meet weekly with the administrator for reconciliation
- Track Excel spreadsheet of patient appointments, recruitment, and follow-up in the study clinic
- Develop report of clinic flow weekly for study clinic meetings
- Present weekly summaries marking progress in enrollment and tracking of subjects in the study in coordination with the study doctor
- Will work with the study doctor, lab assistant, and community health worker to follow-up subjects by phone and home visits
- Perform any other duties and responsibilities that may be given by the principal investigator or study coordinator

Data Manager

- Oversee the work of the data clerk as below and assume any of the duties of the clerk that may be required due to his absence or inability to perform
- Manage the data clerk
- Contribute to the design and modification of protocols, which define what and when data are to be collected
- Design and approve forms on which data are collected
- Be responsible for data collection forms and informed consents (both old and new) that are used in the study
- Manage data information entered by the data clerk on study patients with Hope Center
- Ensure that patient study files are properly filled, documented, and stored
- Manage data backup on weekly basis
- Coordinate the transfer of data with the Coptic Hope data manager to the research databases with the data clerk
- Design SPSS database and manage both the SPSS and Access databases for the study
- Ensure the databases meet requirements for the entry and reporting of clinical data
- Maintain daily, weekly, and monthly work schedules for the data office with the data clerk and ensure their completion
- Check for errors in the data, correct the errors, and maintain cleanliness of the data
- Check and manage the data log book of errors produced by the data clerk
- Coordinate the data-checking process and produce a monthly report on the data quality

- Thoroughly clean the data every 3 months to ensure cleansing of errors
- Sort out any data entry or error problems weekly with the study doctor and study coordinator
- Run frequencies and range checks to identify extreme values monthly
- Present weekly and monthly reports of data analysis
- Assist the study doctor and receptionist in the presentation of weekly summaries marking progress in enrollment and tracking of subjects in the study
- Assist the receptionist and study doctor in the preparation of monthly summary tables on number of women enrolled in each study arm and to consolidate the weekly reports
- Prepare laboratory shipping lists with the lab technologist
- Be responsible for maintaining the security of the data
- Generate study ID numbers
- Be responsible for linking and de-linking data
- Train clinical research staff to help improve the quality of the data being collected
- Assist in standardizing data management procedures such as documentation for study operating procedures
- Develop and maintain documentation and data management guidelines
- Perform other duties that may be given by the principle investigator or study coordinator

Data Clerk

- Enter questionnaire data and laboratory testing information into a computer database

- Scan, verify, and check data in Teleform
- Prepare new patient files and ensure all files contain the required questionnaires
- Maintain Access, Excel, and SPSS computer databases for the study
- Maintain daily, weekly, and monthly work schedules and ensure their completion
- Conduct weekly data quality checks with guidance from the data manager
- Check for errors in the data, correct the errors, and maintain cleanliness of the data
- Inform the data manager and study coordinator and of any data entry problems on a weekly basis
- Keep a data log book of data entry queries and inconsistencies
- Back-up all data weekly (Friday)
- Back-up all data to an off-site disk weekly (Friday)
- Coordinate the timely movement of questionnaires, data forms, and information between the Hope Center, the Coptic research wing, and the KNH data office
- Ensure that the computers, printers and scanner are in good order and free from viruses
- Perform any other duties that may be given by the data manager, study coordinator, and principal investigator

Lab Technologist

- Oversee the work of the lab assistant as below and assume any of the duties of the lab assistant that may be required due to his absence or inability to perform
- Manage the lab assistant

- Ensure enrolled patients have their blood samples and/or other specimens collected at all visits according to the study schedule
- Ensure equipment and supplies are available, working, and well maintained
- Ensure the lab is maintaining good laboratory practices
- Maintain and manage the inventory of laboratory supplies and equipment
- Collect laboratory specimens (including blood) from study participants if the lab assistant is absent or unable to perform
- Keep track of laboratory specimens by updating and maintaining the lab database
- Prepare for the shipment of lab specimens (HIV and HPV)
- Prepare media for collection of samples
- Be responsible for maintaining appropriate freezer temperature
- Monitor freezer temperature by keeping an accurate temperature chart if lab assistant is unavailable
- Be responsible for and enact proper emergency procedures if the freezer is not functioning
- Oversee and manage collection and storage of the following specimens: urine for pregnancy, HPV swab, HIV swab/CVL, Pap smear, and biopsy specimens
- Coordinate the delivery of lab specimens (Pap smear and biopsy) and collection of results with the pathologist
- Coordinate collection, delivery, and recording of CD4 counts

- Track specimens and results in lab book
- Perform other duties that may be given by the study coordinator or principal investigator

Laboratory Assistant

- Collect laboratory specimens (including urine and blood) from study participants
- Track CD4 results from the medical records office and update the CD4 results log in coordination with clinic lab tech, clinic data manager, and study senior data analyst
- Ensure timely transportation of questionnaires and research files between Coptic and KNH offices
- Assist the lab tech in the collection and storage of the following specimens: urine for pregnancy, HPV swab, HIV swab, Pap smear, and biopsy specimens
- Assist the lab tech in the delivery of lab specimens (Pap smear and biopsy) and collection of results with the pathologist
- Track specimens and results in lab book
- Assist the lab tech in the delivery of specimens and collection of results
- Assist the lab tech in the processing and freezing of samples
- Monitor freezer temperature by keeping an accurate temperature chart and inform the lab technologist and study coordinator if there is a failure
- Remove frost and clean the freezer
- Record and present minutes at clinic research meetings if the administrator is not available

- Clean and help organize study clinic
- Assist the receptionist in following subjects enrolled in the study and ensure they are retained in the study and proper follow-up is done both at the research clinic and the Hope clinic
- Assist the receptionist in keeping track of all the patients enrolled and determine if any patient has missed appointment and take action to report and bring these patients back under care and supervision
- Ensure completed questionnaires are delivered to the data clerk within 24 hrs
- Will perform home visits as necessary in coordination with the receptionist and study doctor
- Will help phone subjects for follow-up in coordination with the receptionist and study doctor
- Perform other duties that may be given by the laboratory technologist, study coordinator, or principal investigator

Community Health Worker

- Accompany clients for possible enrollment from the Hope Center to the cervical screening clinic or study lab for urine testing
- Accompany clients between the study lab, Coptic lab, and the cervical screening clinic
- Help clients schedule appointments at the Hope Center
- Clean and help organize study clinic

- Will perform home visits as necessary in coordination with the receptionist and study doctor
- Will help phone subjects for follow-up in coordination with the receptionist and study doctor
- Perform duties given by receptionist or study coordinator

Driver

- Transport specimens and data files
- Assist the administrator in purchase of supplies and equipment
- Maintain car and ensure it is running well and has fuel
- Perform duties given by administrator or principal investigator
- Pick supplies from the office to the clinic

Study Population & Recruitment

The study will be a prospective randomized clinical trial enrolling HIV-positive women who receive care at the Coptic Hope Center for Infectious Diseases in Nairobi, Kenya. The study clinic will screen at least 2,400 HIV-positive women from the Coptic Hope Center for cervical cancer and, of whom, approximately 400 will be enrolled and randomized to receive treatment.

It is estimated that at least 2,400 women will need to be screened in order to identify 400 women with high-grade intraepithelial lesions. This is based on a prospective analysis of 500 HIV-positive women who underwent Pap smear screening at the Hope Center between July and November 2009. In this analysis, 187 (37%) women had normal cytological results, 292 (59%) had abnormal cytological results, and 21 (4%) had results which were indeterminate due to inflammation, inadequate sample collection, or insufficient data. Abnormal cytological results included 77 women (15%) with atypical squamous cells of undetermined significance (ASCUS), 121 (24%) low-grade squamous intraepithelial lesions (LSIL), and 92 (18%) high-grade squamous intraepithelial lesions (HSIL).

Adult HIV-positive women receiving care at the Coptic Hope Center who are not pregnant by clinical examination or history, have an intact cervix, have not received prior cervical treatment, do not have a history of a bleeding disorder, do not have any known allergy to study medications OR their alternatives, have initiated sexual intercourse and are above 18 years of age will be informed of the study by non-study clinicians and health care workers. Subjects will be excluded by the study at initial cervical cancer screening if they are HIV-negative, male, below 18 years of age, pregnant by clinical examination or history, post-hysterectomy, post-cervical cancer treatment or have known allergies to study medications.

Potential participants will be identified by Hope Center clinical officers seeing patients as part of routine HIV medical care at the Hope Center. Clinical officers will use “Pre-Screening Talking Points” to inform patients about the study. If subjects are interested then they will be directed to the study clinic in a physically separate room staffed by study personnel. We will also use recruitment leaflets that will be distributed at the Hope Center Reception. The leaflets will have the contact information of the study staff. At the study clinic, a study staff member will inform potential subjects of the study using “Screening Talking Points”. If subjects remain interested and are eligible then they will be enrolled (see Study Flow I). An appointment calendar will be kept by the study receptionist on an MS Access database to track who has attended the clinic and received screening.

Interested women will be referred to the study clinic (Room A) which will be staffed by two study nurses trained to perform Pap smears on the same day after obtaining informed consent for screening. Women who enrolled in the screening portion of the study and obtained a Pap smear will be asked to return to the study clinic (Room A) for results 2 weeks later. If Pap result is positive for HSIL, the subject will be referred to a separate room (Room B) where a study doctor and a study nurse will obtain a biopsy to confirm CIN 2 and 3 by histology. From this point, the subject will return to Room B and see the study doctor and nurse for further follow-up. Four to six weeks after biopsy, the subject will return for her results and to discuss treatment options. At this point, study staff will obtain informed consent for randomization and follow-up, and the study doctor will randomize histology-confirmed subjects (approximately 400 women) to LEEP or cryotherapy. Criteria for being ineligible for cryotherapy include if a polyp or anatomic defect prevents access to the cervix and/or if the lesion size is >75% of the cervix or is larger than the cryoprobe tip, or if the lesion is not visible in its entire extent or extends more than 2 to 3 mm into the endocervical canal. If any of these subjects are diagnosed with severe cervical disease confirmed by

histology at these later screening time points, they will be offered LEEP by the study or referred for appropriate care.

Patients who require further care for cervical disease unavailable at the study clinic (eg. hysterectomy) or who decline to enroll in the treatment portion of the study will be referred to KNH. Through patient tracing the study doctor and staff connected to the KNH gynecology department will make best efforts to assure that patients receive proper care. Referral notes and copies of results for histopathology, cytopathology, CD4 count, and HIV viral load will be sent with the patient during referral. All study specific documentation including study numbers and title will be removed. Only the subject's name and age, as is relevant to the referral, will be on the documents sent for clinical care at KNH or Coptic Hospital. Subjects will receive government subsidized care at these sites but the study will not pay for these interventions or treatment.

Study clinic visits will be conducted separately from regular medical visits a patient may make at the Hope Center. Subjects will continue to follow-up with their doctors at the Hope Center, schedule their visits through the Hope receptionist, and pick up their medications at the clinic. However, study clinic visits will be conducted by staff employed by the study in physically separate locations from routine medical visits. Study appointments will not conflict with the HIV clinical care that patients receive through coordination between the study receptionist and the clinic receptionist. Study doctors will not provide HIV care or treatment, and subjects will remain enrolled as patients at the Hope Center and receive the same care as prior to enrollment. Before lab tests for the study are performed, confirmation from the subject (and if necessary, the laboratory) will be made to assure no tests, such as Pap smears ordered as standard of care or blood draws, are unnecessarily repeated.

Medical information collected by clinic staff at the Hope Center will be made available to the study team after written informed consent by the subject. This will contain all information collected at the Hope Center including laboratory data. Only clinic staff will collect medical information from subjects related to the management of their HIV disease. Study data for the PHE will be collected on separate forms and scanned, cleaned, and analyzed at a location unassociated with the clinic and its medical records. Only study staff will collect study data on forms described in this protocol. If institutional IRBs approve, we will make study data available to the clinic staff at the Hope Center after written informed consent is signed by the subject. As soon as a participant's results become available, the study staff will enter the results in a form that is stored in the participant's Hope Files as a means of communicating the results to the participant's clinicians. This will be done in coordination with the medical records department at the Hope Center. If clinical information is discovered about a subject that needs immediate attention, clinical staff will be available on site at all times to assist in care of the patient and will be informed by the study staff. In the event that a participant needs emergency care, the study clinical staff will be prepared to stabilize patients and emergency services are available at Coptic Hospital 24 hours a day. Moreover, clinical care is available at KNH at any time if needed or preferred by the patient. Participants will be given careful instruction as to which symptoms would necessitate them seeking additional medical care, especially after receiving treatment. The participant will receive contact information of study staff whom they should contact in case of an emergency or if they need guidance as to how to manage a certain symptom or condition. Study staff can share pertinent information with clinic staff in these situations by phone or in person as the study clinic is located on the hospital grounds.

Participants will be consented for screening by study staff at their enrollment visit and for randomization and follow-up at their biopsy result visit. If an

exited subject becomes eligible for re-enrollment due to re-reading of cytology results (See QA/QC section), then re-enrollment consenting will be done at the disclosure visit for the new results. All consenting study staff members will be able to describe study procedures in English, Kiswahili and possibly another local language. Other nursing and study staff are available and should be contacted in the case of a participant who presents for enrollment but is not conversant in any of the languages the consenting staff member speaks. Consent documents will be prepared in both English and Kiswahili at a basic reading level. The English version will be used by study staff to produce the Kiswahili version and several members of the staff will review the document to assure its accuracy and readability.

Participants will be encouraged to ask questions at any point during the consent process and will be given a copy of the document to review on their own. If a participant has a question that cannot be answered by the consenting staff member, other study staff will be available for consultation. The patient may also contact other study staff at her convenience. The participant may also interrupt the consent process at any time if she needs more time to consider her participation.

Subjects with initial Pap smear results that are Normal or ASC-US will be exited from the study after receiving their Pap smear results. Subjects with LSIL Pap smear results will be offered Colposcopy. According to the 2012 ASCCP guidelines, participants found with LSIL in the absence of HPV co-testing should be recommended colposcopic evaluation and biopsy. These biopsy histology results will be treated the same as other study histology, including possible treatment randomization for CIN 2/3 and CIS. This will enable researchers to monitor lesions which may escalate from low-grade to high-grade in subjects who may be eligible for subsequent randomization. Results from repeat Pap-smears will be treated the same as results from initial Pap smears: Women with Normal/ASC-US results will be exited from the study. Subjects with HSIL/ASC-H/HSIL-invasion-not-ruled-out and results

suggestive of cervical carcinoma (i.e. high grade intra-epithelial lesions) will undergo colposcopy with biopsy, and may potentially be eligible for randomization upon confirmed histology results. If a subject with high grade intraepithelial lesions detected on study-related Pap smears is later found to have normal or ASCUS on a follow up Pap smear (study related or not) that occurs prior to or in lieu of a colposcopic evaluation, the subject will still be required to have a study related colposcopy and biopsy.

Subjects who have biopsy-confirmed CIN 2, 3 or CIS will be eligible for the randomization and treatment portion of the study. They will be presented with the consent information upon receipt of biopsy result. Subjects with biopsy results suggestive of cervical carcinoma will be exited from the study and referred for further evaluation and management at KNH. Subjects with CIN I biopsy results will be asked to return to the clinic for repeat Pap smear in 6 months and will remain in the study. If a subject receives three consecutive CIN I histology results, she will be offered LEEP treatment by the study doctor. These participants will be treated and followed the same as other non-randomized LEEP participants.

Of note: Women with normal or with ASCUS Pap smear results (that are exited from the study) will be referred for subsequent cervical screening follow-up at the Hope Center Cervical Cancer Screening Program (CCSP). This program is administered by the Coptic Hospital and is supported by the Coptic Mission, PEPFAR, and other donor funding. The CCSP offers free cervical cancer screening to HIV-positive women enrolled in the Coptic Hope Clinic. Currently, a single visit ‘screen-and-treat’ strategy for cervical cancer prevention and treatment will be used. The screen-and-treat method involves visual inspection of the cervix followed by treatment of precancerous lesions by cryotherapy at the same visit. Patients who cannot be treated with cryotherapy are referred to KNH for subsidized cervical

care and treatment. As all study subjects are drawn from the Coptic Hope HIV clinic, they are all eligible to be screened, treated, and followed at the CCSP.

If a woman exited from her normal or ASCUS Pap smear result prefers to rescreen with the study, she may be re-enrolled using the re-enrollment consent form.

Study Procedures

1. SCREENING visit (at least 2,400 patients) (see Study Flow Diagram I, Enrollment)

- At screening, subjects who agree to participation will:
 - Sign a written informed consent
 - Answer the “Enrollment” questionnaire which will ask her sociodemographic background, sexual history, and cervical cancer screening history.
 - Complete “Address and Intake” questionnaire.
 - Undergo a physical examination including a pelvic examination with a Papanicolaou smear of the cervix. The results of the physical examination will be entered in the “Pap Smear” questionnaire
 - Women found to have evidence of an STI will undergo syndromic management according to the following algorithm:
 - i. Vaginal discharge without abdominal pain
 - 1. Treat for vaginitis:
 - a. Nystatin 1 pessary everyday x 5 days and
 - b. Metronidazole 2g x 1
 - 2. If no improvement after 7 days, treat for cervicitis:
 - a. Norfloxacin 800mg x 1 and
 - b. Doxycycline 100mg BD x 7 days
 - ii. Lower abdominal pain
 - 1. If due to surgical or gynecological causes, refer

2. If cervical motion tenderness, treat for pelvic inflammatory disease:
 - a. Norfloxacin 800mg x 1 and
 - b. Doxycycline 100mg BD x 7 days and
 - c. Metronidazole 400mg BD x 10 days
 - iii. Genital ulcer disease (GUD)
 1. If multiple lesions grouped together with a history of recurrence, treat for Herpes simplex genitalis:
 - a. symptomatic treatment
 2. If other GUD, treat empirically:
 - a. Erythromycin 500mg three times per day x 7 days and
 - b. Benzathine penicillin 2.4 million U IM stat
2. Two-week visit for RESULTS (at least 2,400 patients)
- Patients will return to the clinic two weeks later for Pap smear results which will be entered on the “Colposcopy” questionnaire. We expect that an estimated 1,700 women will have results (no dysplasia or ASCUS) that do not require study biopsy/colpo procedures. These patients will be ineligible for randomization.
 - Women with no dysplasia or ASCUS Pap smear results will be exited from the study. They will be followed up at the Coptic Hope Cervical Cancer Screening Program (CCSP). The Hope screening program offers free cervical screening through visual inspection with acetic acid (VIA). Women who are screened and exited from this study with normal Pap smear results will be followed up at CCSP one year after initial screening. Women who are screened and exited from this study with ASCUS Pap smear results will be followed up at CCSP 6 months after initial screening. Women who have exited the study due to a cytologic diagnosis of no dysplasia or ASCUS will

be eligible to re-enroll in the study and obtain a study related screening pap smear, approximately 12 months after their last normal/ASCUS study related pap smear.

- Women found with unsatisfactory results will have their Pap smear repeated.
- Women with LSIL will be offered Colposcopy. According to the 2012 ASCCP guidelines, participants found with LSIL in the absence of HPV co-testing should be recommended colposcopic evaluation and biopsy.
- Any patient who receives a cytological diagnosis of high-grade intraepithelial lesion (HSIL and ASC-H, and HSIL-invasion-not-ruled-out or SCC) (~18% of the patient population or approximately 400 patients), will be recommended to undergo colposcopy as per 2012 ASCCP guidelines. If a subject with high grade intraepithelial lesions detected on study-related Pap smears is later found to have normal or ASCUS on a follow up Pap smear that occurs prior to or in lieu of a colposcopic evaluation, the subject will still be required to have a study-related colposcopy and biopsy. Prior to participating in any cervical procedure, including biopsy, women will undergo a rapid pregnancy test. Pregnant subjects will be exited from the study and followed-up in the CCSP. These women will also be referred for prevention of mother-to-child transmission (PMTCT) care at Hope Center. If the pregnancy result is negative, they will be referred to the study doctor for colposcopy-directed biopsy which will clarify the extent and severity of disease. Results of the colposcopic examination will be entered in the “Colposcopy” questionnaire (see Study Flow I). For patients who have glandular epithelium abnormalities (AGC) on Pap or squamous epithelium abnormalities without visible lesions on colposcopy, endocervical sampling with curettage will be performed in order to identify possible abnormalities in the endocervical canal.

- After undergoing biopsy, women will be given free condoms and advised to abstain from sexual activities for at least 10 days.
3. 6-8-week visit for RANDOMIZATION (~400 patients) (see Study Flow II)
- Subjects who have undergone biopsy will return four to six weeks later for a follow-up visit to be given the results of the biopsy. Biopsy results will be entered in the “Treatment” questionnaire.
 - Those with biopsy results that are positive for a high grade pre-cancerous lesion (CIN 2/3 or CIS) and if the lesion is amenable to cryotherapy or LEEP as per Kenyan national guidelines, will be eligible for randomization. If they are willing to undergo informed consent for the treatment study, they will be randomized to receive either cryotherapy or LEEP (see Study Flow II). Those who decline to participate in the treatment study will be exited and referred to KNH for treatment. Randomization questionnaire will also be completed during this visit.
 - Those patients who do not have lesions which are amenable to cryotherapy due to size or access will not be randomized but will be offered free LEEP and followed up every 6 months for 2 years, the same way as randomized participants. Those who have lesions that are neither amenable to LEEP nor cryotherapy due to size or severity of disease, or anatomy does not allow proper access to the cervix, will be referred to KNH for subsidized care at this government hospital. The study will not pay for this treatment. Patients will be given follow-up appointments in the study clinic after referral to determine outcomes. They will have the option to continue to be followed at the CCSP.

- Those with CIN 1 will not be randomized or exited but will be asked to return to the study clinic after 6 months for a repeat Pap smear.
- Those who show no dysplasia on histology will be exited from the study and referred to the CCSP for further follow-up.
- It is expected that approximately 400 patients will be eligible for randomization. During randomization, a sealed envelope will be opened by the study doctor that will reveal the randomization arm that has been assigned to the study ID number. Randomization assignment will be performed on a computer by Dr. Barbra Richardson, the study statistician, in Seattle, and preparation of the envelopes will be performed by the study data manager in Nairobi in conjunction with Dr. Richardson. This is the standard used for all randomized clinical trials conducted by the University of Washington in Kenya. Study investigators and the study doctor will not have access to the randomization sequence. Study investigators and staff will not be blinded to randomization.
- Those eligible for randomization will have an HIV cervical viral level swab taken before treatment (LEEP or cryotherapy).
- After receiving LEEP or cryotherapy, women will be given free condoms and advised to abstain from sexual activities for at least 4 weeks. Women will also be offered an information sheet on the treatment procedure and abstinence information to take home for their spouse or partner. Information includes treatment given, recommended abstinence period and reasons for abstaining from sexual intercourse. These information sheets will be optional to maintain patient confidentiality and safety, especially for people who have not disclosed their status.
- Patients undergoing randomization will have the following performed:

- Blood (10mls) will be drawn to check for HIV viral load and CD4 count
 - Cervix will be swabbed to assess the presence of Human Papillomavirus (HPV) prior to LEEP or cryotherapy
 - Cervix will be swabbed for HIV viral level prior to LEEP or cryotherapy
 - LEEP or cryotherapy performed based on randomization envelope
 - Those undergoing LEEP will have the LEEP piece taken for histology
 - “Randomization” questionnaire completed
 - “Treatment” questionnaire completed
4. First, second, and third-week visits for cervical HIV-1 SHEDDING after treatment (~400 patients) (see Study Flow II)
- Patients who have been randomized and received treatment (approximately 400 patients) will be followed every 1 week for 3 weeks AFTER the treatment intervention to assess healing and measure HIV-1 viral shedding from the cervix. These events can be called the 7-9, 8-10, and 9-11-week visit time points and will probably coincide with the eighth, ninth, and tenth week after enrollment. However, for purposes of measuring HIV viral shedding, it is most important that these time points occur 1, 2, and 3 weeks after the treatment intervention. These terms are consistent and are used for descriptive purposes. They will return 3 times over a 3-week interval and have the following performed:
 - Blood (10mls) will be drawn to check for HIV viral load
 - Cervix will be swabbed to assess the presence of HIV-1
 - “Shedding” questionnaire completed

Shedding information will be considered valid if it is obtained \pm 3 days of the scheduled visit. We will collect shedding information for the first three weeks after treatment but if a patient misses those visits but comes back within six weeks, we will still collect the samples just for comparison purposes.

5. Six, twelve, eighteen, and twenty-four months after randomization visits for RECURRENCE (approximately 400 patients) (see Study Flow II)

- After the 9-11-week visit, randomized participants and those who undergo LEEP but are not randomized because their lesions are not amenable to cryotherapy will return at 6, 12, 18, and 24 months after treatment is administered. At the 24 month visit, the subject will be exited from the study. At these 6-month interval visits, the following will be performed in order:
 - Pap smear to detect recurrence of cervical intraepithelial lesions
 - Same Pap smear swab will be used to assess the presence of Human Papillomavirus (HPV)
 - Blood (5 mls) will be drawn to check for CD4 count
 - “Pap Smear” questionnaire completed
- Pap smear results that are normal, ASCUS, will be filed in the participants file and disclosed at the next scheduled follow-up visit. If pre-cancerous lesions by cytology have recurred then patient will be contacted to return to the study clinic to undergo colposcopic directed biopsy prior to treatment. Results of the colposcopic examination will be entered in the “Colposcopy” questionnaire..
 - If histology-confirmed lesions can be treated by LEEP, then the patient will receive a free LEEP and “Treatment” questionnaire will be filled.
 - If histology-confirmed lesions are too large or cancerous for LEEP then patient will be referred to KNH for subsidized care at this government

hospital. The study will not pay for this treatment. Patients will be given follow-up appointments in the study clinic after referral to determine outcomes.

- Phone calls and home visits will be performed sequentially in the case of a subject who has been lost to follow-up (LTFU) after screening or randomization (see LTFU section). LTFU will be prepared and discussed at weekly clinic meetings. Written reports will be prepared from these meetings and sent to co-investigators. Home visits will be performed by community health workers trained to be discreet in their approach to finding patients. They will not wear clothes that would identify them as health workers and will only use public transportation or unmarked private vehicles. They may ask about the participant's health or whereabouts.

6. Specimen transportation

Blood samples will be processed at the study clinic and will be sent to the US for processing at a later date when enough samples have been collected, approvals have been obtained, and the laboratory is ready to receive the specimens. The Pap smear and biopsy samples will be transported to the pathologist on a daily basis and the results will be brought back after processing. HPV collection media will be stored at room temperature at the study clinic and may be transported outside of Kenya at a later date. Biopsies will be transported to the pathologist and the results brought back within 2 weeks after processing. The paraffin-embedded tissue blocks remaining after the histology processing will be brought back and stored at the study clinic. A transport log book will be used for this purpose. A results log will be kept to account for results received from the pathologist.

7. Infection Control

Hand washing with soap, disinfecting used linen and equipment, proper waste disposal, single use needles and syringes, use of gloves and autoclaving of equipment at the Hope Center will be adhered to.

8. Transport Fees

No money for transport will be given on enrollment for this will be treated as a scheduled clinic day at the Hope Centre. All subjects screened will get Ksh 300 on return to the clinic after initial enrollment and screening. If a woman would like more time to consider entry into the study, she will be given money for travel at the time she returns with the intent to enroll in the study. Subjects will receive travel money on presentation to the study office on their scheduled visit day and a list of subjects (by study ID) who get travel money will be maintained by the receptionist on daily basis. The receptionist will be getting transport money from the administrator. A top up will be done whenever the amount fall below Ksh 3,000. Maximum amount to be collected from the administrator at any given time will be Ksh 10,000.

9. Notification of Results

Cytology results: Subjects will be notified of their screening cytology results at a study visit scheduled two weeks after their screening visit. If on routine cytology QC re-readings, consensus cytology results are discordant from the initial reading and alternative management is required, subjects will be notified and called back for appropriate follow-up. Subjects with new cytology results of LSIL will be offered colposcopy and biopsy according to 2012 ASCCP guidelines. If the new cytology results are HSIL, ASC-H, or HSIL cannot rule out invasion, patients will be presented with evaluation, treatment and follow-up options, which includes continuation in the study (even if previously exited). Subjects will be given consent information to re-enroll into the study. A Colposcopy form will be completed at this visit. Cytology results from bi-annual follow-up visits for randomized participants and non-

randomized LEEP participants will be disclosed at the next scheduled follow-up visit if the result is Normal or ASCUS. If pre-cancerous or cancerous lesions by cytology have recurred then patient will be contacted to return to the study clinic to undergo colposcopic directed biopsy prior to treatment.

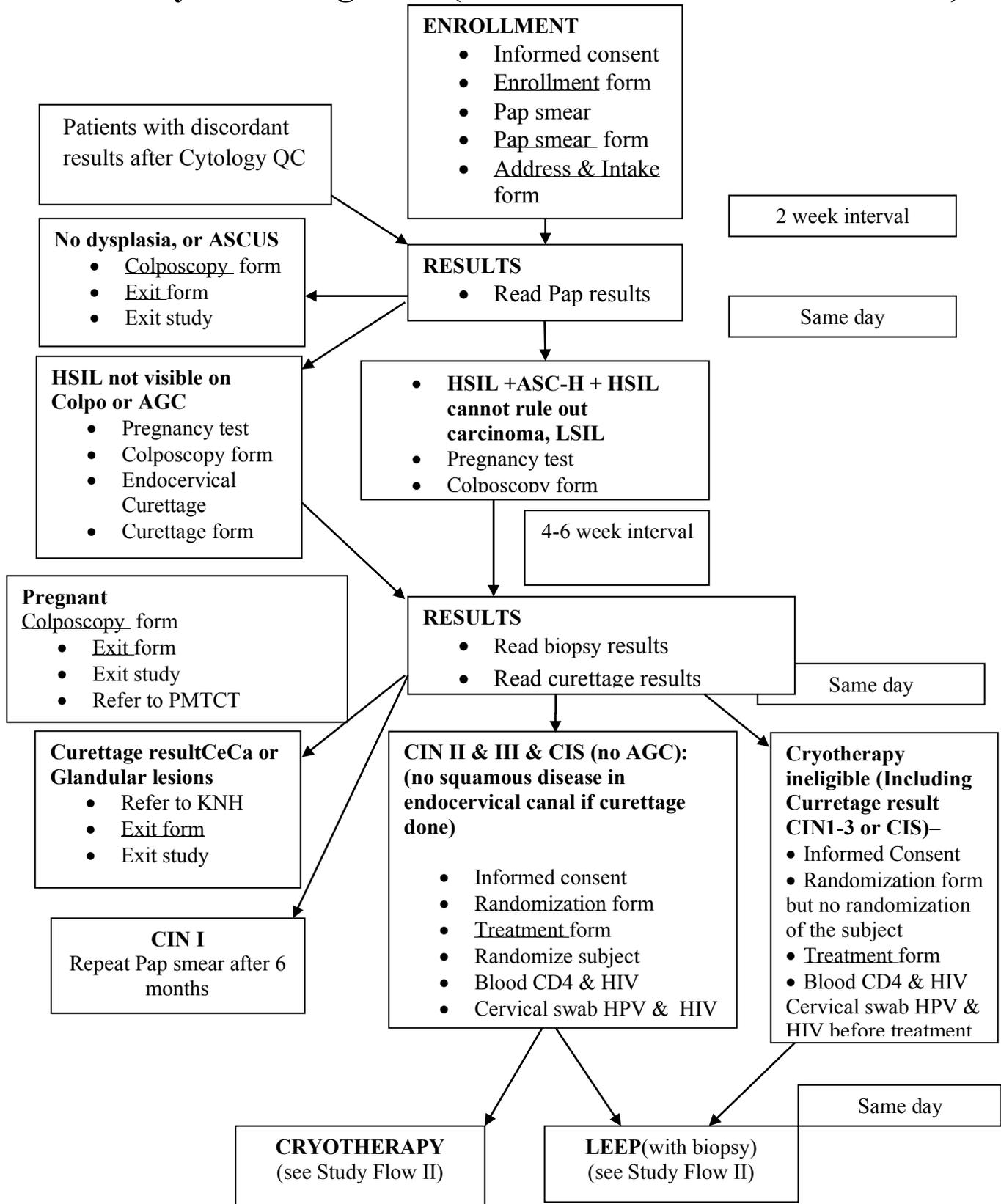
Histology results: If the participant needs a biopsy (i.e., HSIL, ASC-H, HSIL cannot rule out invasion, or ICC), they will need to return 4-6 weeks later for histology results. If study staff or a participant feel that having a counselor present at any of these visits would help facilitate the conversation, this can be arranged through utilization of Hope Center counseling staff.

Participants who do not come to their scheduled appointments will be called or visited at home using the contact information provided at enrollment (process described thoroughly below in the Loss to Follow-up & Mortality section).

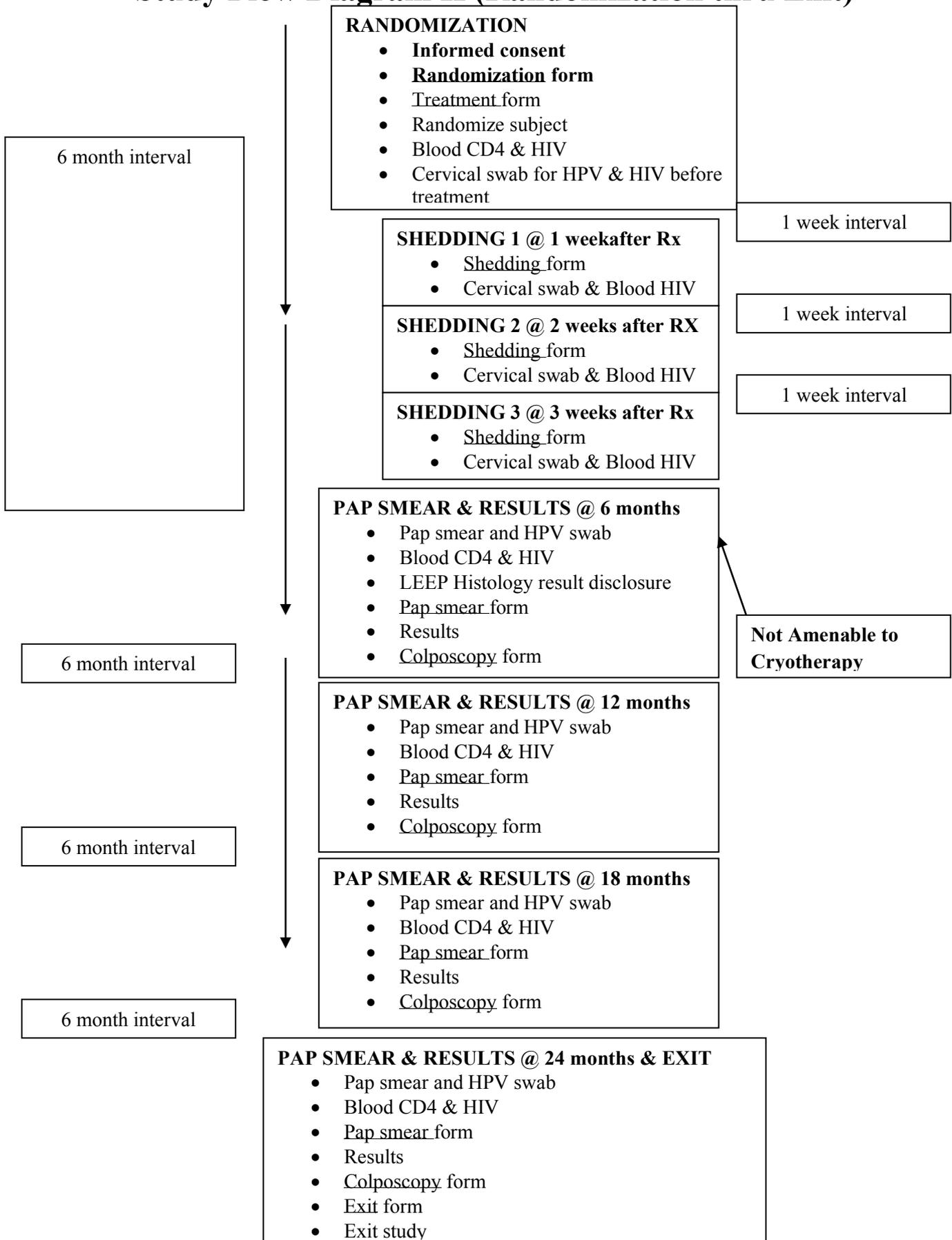
10. Exit

Subjects will exit the study after an interview with the study doctor or study nurse. An “Exit” form will be completed. Participants will be notified that if they wish to know the final results of the study, they may contact the study office 6 months or a year after their participation is complete.

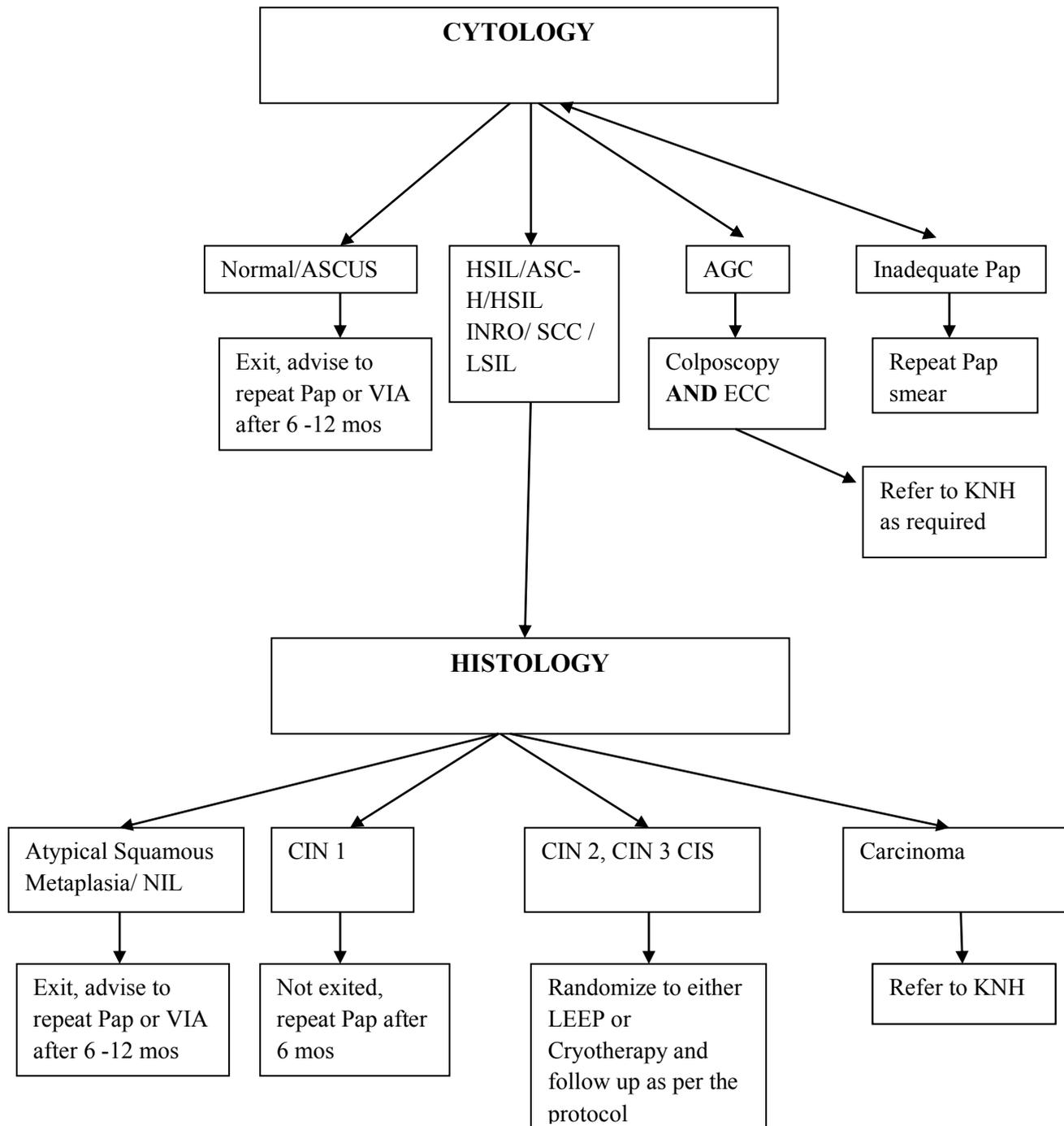
Study Flow Diagram I (Enrollment thru Randomization)



Study Flow Diagram II (Randomization thru Exit)



Study Flow Diagram III (Study Tests)



Loss to Follow-up & Mortality

Tracing of randomized participants

If a randomized subject fails to appear for a scheduled study clinic visit then she will be contacted by phone the next day by the receptionist. If the first phone call after a missed visit does not reach the patient or someone who is in contact with the patient, then calls will be repeated at least twice a week for two weeks. If the patient is reached by phone, another visit will be scheduled as soon as possible within one month. If the patient also fails to return for this rescheduled visit, the receptionist will call again the following day. If again the patient does not return for the next rescheduled visit (the third scheduled visit of its kind), or if at least 4 calls have been unsuccessful in reaching the subject, an attempt will be made to visit her at home by a community health worker. If a home visit is unsuccessful in reaching the subject, or if home-visit is successful but the subject fails to return to the study, then she will be considered lost to follow-up (LTFU). If a phone call or home visit reaches the subject, then an appointment will be made for the subject to return to the study clinic. (See Study Tracing Diagram for randomized patients).

Tracing of non-randomized participants

If a non-randomized subject fails to appear for a scheduled visit, including 6-month repeat Pap smear visit after a prior LSIL or CIN1 result she will be contacted by phone the next day by the receptionist. If the phone call does not reach the patient or someone who is in contact with the patient, then a second call will be made the following week. If both calls are unsuccessful in reaching the subject, or if the visit is rescheduled but the subject again fails to show up, then her Coptic Hope file will be flagged, and she will be considered LTFU. If a

phone call or home visit reaches the subject, then an appointment will be made for the subject to return to the study clinic.

Tracing of patients with new QA/QC Cytology results

Both enrolled participants and patients who have been exited from the study may need to be contacted to receive new cytology results based on laboratory QA/QC. Women whose final QA/QC cytology result has been downgraded from the initial result that was disclosed to them, or women whose final results do not require treatment (Normal, ASCUS) will be followed in the same manner as non-randomized subjects: two weekly phone calls and flagged Hope file. These patients will be operationally considered lost at this same point.

Women whose final results have been upgraded to ASC-H, HSIL, or SCC, and women whose final results are unsatisfactory but had at least one reading of ASC-H, HSIL, or SCC, will receive additional follow-up efforts. This includes at least 6 phone calls over 4 weeks, with at least one call over the weekend. Home visits will also be conducted for these patients when possible. These subjects will be defined as LTFU on a case-by-case basis when all efforts to contact them have been completed.

Tracing of patients with cancer

Any woman who receives an initial or final laboratory result of SCC (cytology) or ICC (histology), whether she is randomized, non-randomized, or exited, will receive the same extensive follow-up efforts as noted above. This includes at least 6 phone calls over 4 weeks, including a phone call over the weekend, and a home visit when possible. These subjects will be defined as LTFU on a case-by-case basis when all efforts to contact them have been completed.

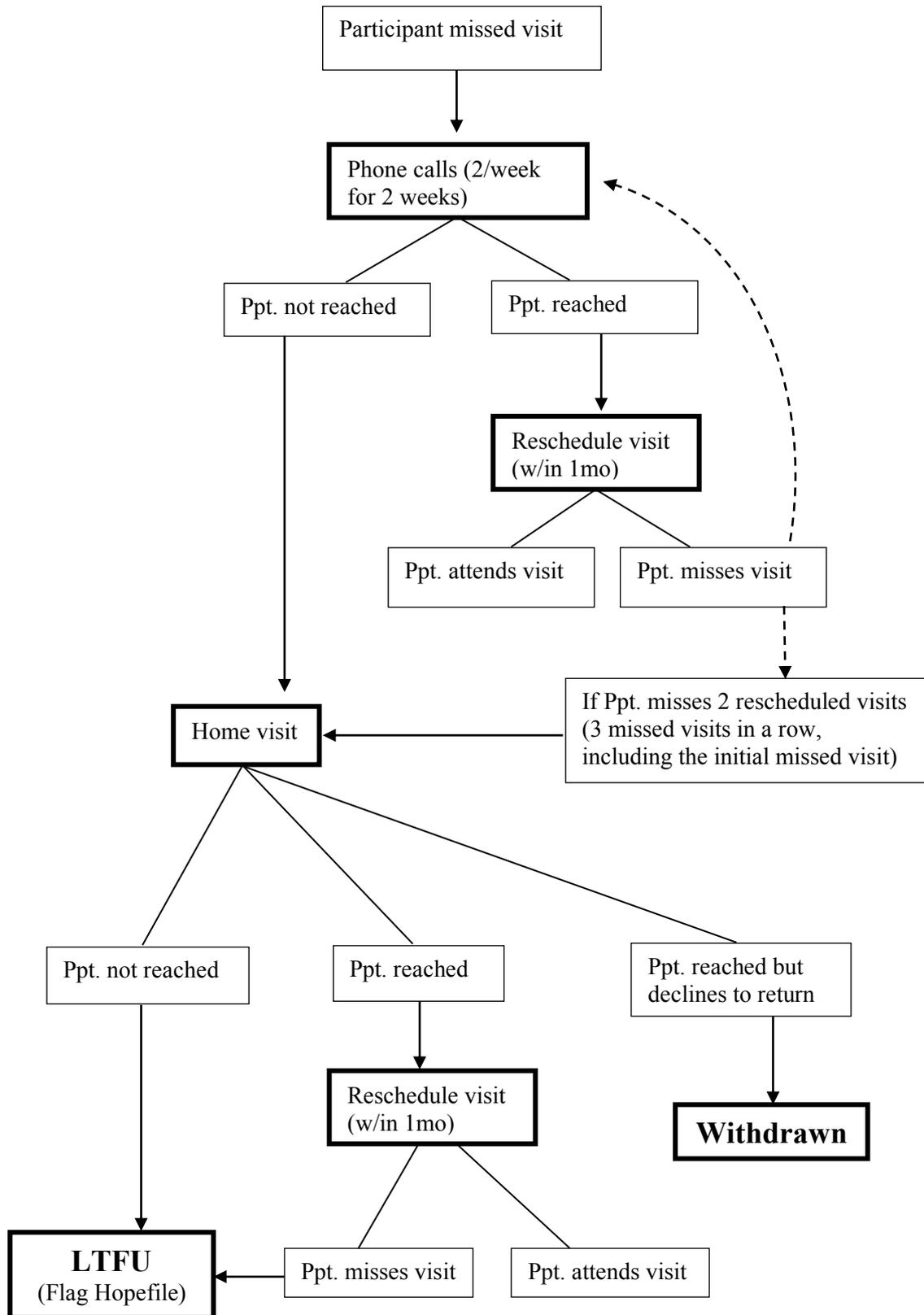
Documentation of tracing

All attempts at contacting the patient either through phone call or home visits will be documented in the participant's chart notes. All attempts at home visit will be recorded on the "Patient Contact" form. If a subject is LTFU, this will be recorded on the "Patient Contact" form, in addition to reasons for loss. If a subject withdraws or asks to dis-enroll from the study, the "Exit" form will be completed. If a subject is determined to have died through phone call or home visit, then the "Verbal Autopsy" form will be completed. Information on loss or death will be shared with the Hope clinic team and social worker so that they may update their files.

Statistical considerations for LTFU

Those who are LTFU or dropout after randomization will contribute time and data to the outcomes of measurement for as long as they have remained in the study. They will continue to remain in the arm to which they have been randomized and analyses comparing interventions will include these participants. If, for example, they reach the 6 month time point but did not appear at 12 or 24 months, then data from this subject will be included until 6 months after randomization in a survival analysis. The study coordinator and study doctor will oversee loss and mortality data in coordination with the receptionist.

Study Tracing Diagram for Randomized Participants



Statistical Methods

The sample size was calculated to detect a 10% difference in treatment outcomes between LEEP and cryotherapy (4% versus 14%)²⁶ with 80% power and a 0.05 significance level. Accounting for the possibility of a 20% loss-to-follow-up rate over two years, we estimated that 400 women with high grade lesions would need to be enrolled in the study with 200 women randomized to each arm. It is estimated that 200 women in each arm will allow detection of a 0.25 log₁₀ HIV virus level difference in cervical shedding between the two arms with greater than 90% power. Based on results from an ongoing cervical cancer screening study implemented at the Hope Center in 2009, our estimation of the prevalence of high-grade intraepithelial lesions is about 18%. Given this estimate and the number of women who are likely to be eligible for randomization, we calculate that we need to enroll 2,400 women for initial screening.

The primary analysis of the study will be to compare treatment outcomes between the LEEP and cryotherapy intervention arms in an intent-to-treat analysis. Treatment outcomes will be measured by Pap smears taken every 6 months during two-year follow-up. Pap smears demonstrating high-grade intraepithelial lesions will be considered positive and compared against negative or low-grade intraepithelial lesions. A Chi-square test will be used to compare the percentage of positive Pap smears between the intervention arms at 6, 12, 18, and 24 months. Secondary analyses will compare cervical HIV-1 RNA viral shedding between the treatment arms. A comparison of the average area under the curve (AAUCMB) of long-transformed HIV-1 RNA cervical viral loads will be performed between the two arms using an analysis of covariance (ANCOVA) model.

Our analysis will include contraceptive history as a potential confounder. While there has been some suggestion in the literature that HIV infectivity among women on oral

contraceptives is increased,^{33, 34} a recent study from the UW did not show a significant increase in levels of HIV RNA in cervical samples obtained from women on hormonal contraception.³⁵ A more recent multicenter randomized controlled trial performed in Asia and Africa also did not find an association between hormonal contraceptive use and increased HIV transmission.³⁶ However, increased viral shedding due to hormonal contraceptive use remains an important and interesting question and one that we will examine in our study.

The analysis will control for immunological status through measurement of CD4 count as well as presence of and duration on antiretroviral medications. Given that the study will recruit patients from an antiretroviral treatment clinic, it is expected that the majority of subjects will be on antiretroviral medications (~75%). Type and duration of antiretroviral drugs (ARVs) will be obtained through accessing clinical medical files and CD4 count measurements will be obtained regularly during study follow-up. The impact of ARVs on recurrence of cervical intraepithelial lesions is unclear and may have minimal effect.^{37, 38} Also, despite ARV use, HIV shedding has still been detected in the genital tract.³⁹ Therefore, we are confident that the sample size will still be adequate to detect differences between treatment modalities among women on antiretroviral medications.

Data Management

Introduction

Data will be captured on paper forms or electronically using Open Data Kit (ODK) application which has a data collection component to be used on a mobile device, and a centrally managed data store component. ODK is supported by the University of Washington (UW). Data will be cleaned, and analyzed at UW project offices located at KNH in Nairobi, Kenya. This office, which supports several UW studies, will provide administrative and database management support to this research project. Study data will be collected separately from Hope clinic data and will not be linked to the Hope dataset. The forms have been used in a recent cervical screening study and improvements have been made as a result. Analysis of the data will be conducted using SPSS version 16.0 (SPSS Inc, Chicago, Illinois, USA) by UW statisticians and epidemiologists associated with the study. Data will be managed by the Data Manager and the data clerk.

See Appendix B for data collection forms

Facilities

Cervical Cancer Screening room at Coptic Hope Center for Infectious Diseases

The on-site facilities available for cervical cancer screening include two examination beds, one office desk, and one computer, and at least one tablet PC. Electronic and paper data collection forms and paper informed consent forms will be administered here. The address and intake form, which captures patient identifier data will be completed on paper forms only.

UW study clinic and laboratory at Coptic Hope Center for Infectious Diseases

At this site, there is one clinic room and laboratory dedicated to this study. The clinic will have one examination bed, one desk, and one computer. Data collection forms will be filled out at the study clinic and in the Cervical Cancer Screening room. Any paper data forms, including address and intake and lab request forms will be sent to the UW offices in KNH for scanning daily but returned for storage at this site, while the lab forms will be sent to the Lancet laboratory.

UW data and administrative offices at Kenyatta National Hospital

The study has an administrative and a data management office in the UNITID building at KNH. There are five computers, one laser printer, one scanner and a one copier. All computers except the data computer have internet access. All computers have antivirus software which is kept up to date by the data manager. Only non-identifiable data will be collected electronically and stored on a local ODK Aggregate (server) which is maintained by NASCOP. All data collected using the ODK will be downloaded from the nNASCOP server onto the data computer and converted in SPSS format.

Databases

The study data will be recorded in SPSS, MS Access, or written notebooks and include:

Data Collection Forms – this SPSS database includes all questionnaires that are electronically and manually filled at the study.clinic In total there are 16 questionnaires to be filled

- Enrollment
- Pap Smear
- Colposcopy
- Randomization

- Treatment
- Address and Intake
- Shedding
- Patient Contact
- Verbal Autopsy
- Exit
- Cytology Report
- Colposcopic Biopsy Histology Report
- Endocervical Curettage Histology Report
- LEEP Biospsy Histology Report
- CD4 Report
- Follow-up Form

Cervical Cancer Screening Appointment and Tracking – this MS Access database will track patients from the Hope Center who are referred to the study clinic by scheduling appointments and recording dates that screening tests are performed.

Laboratory – this database contains an inventory of all specimens collected, stored, and analyzed.

Specimen Collection – a written notebook provides a written record that tracks when the specimens have been collected and where they are going. Specimens include:

- Blood for CD4 count
- Blood for HIV viral level
- Cervical swab for HIV viral level
- Cervical swab for HPV

- Pap smear
- Colposcopy-directed biopsy
- Endocervical curettage specimen
- LEEP specimen

Storage – a written notebook which tracks specimens which are stored in the freezer and where they are kept. Stored specimens include:

- Cervical swab for HIV
- Cervical swab for HPV
- Blood plasma and cells for HIV

Laboratory Results – a written notebook which tracks the results of laboratory analyses that are performed in Kenya. Results tracked include:

- CD4 count
- Pap smear result
- Colposcopy-directed biopsy result
- LEEP biopsy result
- Endocervical curettage biopsy result

Management

Entry

The data collection forms will be completed either manually on paper forms or electronically on ODK. The following collection forms will be completed manually in paper forms: , address and intake form (which captures patient identifier data), Cytology Report, Colposcopic Biopsy Histology Report, Endocervical Curettage Histology Report, LEEP Biopsy Histology Report and CD4 report, all of which serves as lab request forms.

Each subject has a folder in which all paper forms and questionnaires are stored. These folders are stored in a locked cabinet at the UW study clinic at Coptic. Newly filled-out questionnaires are brought to the UW KNH data office from the study clinic daily. Questionnaires are scanned into the computer database the same afternoon, and folders are returned to the clinic the following morning. Scanned data is verified by Cardiff Teleform software and later cross-checked against the paper questionnaire on the same day by the data clerk after it is exported into SPSS.

For data captured electronically, only non-identifiable data will be collected electronically and stored on a local ODK Aggregate (server) maintained by NASCOP. The Address and Intake form which collects the patients identifiable information will be completed on paper, sent to the UW office in KNH for scanning daily, and returned for storage in the Coptic study clinic where they will be kept in a locked cabinet. Once the data is collected using the encrypted electronic forms, it will be transmitted to the Nascop server through an encrypted path.. As an added layer of security, all electronic data forms (even though they contain no identifiable data) are encrypted on the data collection devices prior to transmission to the local server. The Principal Investigator, the Data Manager and the data collectors will have access to the ODK server with different rights.*Quality*

The data clerk maintains a data logbook where all the data entry queries or errors encountered during data entry are recorded. Data entry queries or errors are then discussed with the study

staff at least once a week. Discussions are held with both study staff and data clerk together at the study clinic and corrections are made to the database by the data clerk upon returning to the data office. The data manager ensures that all errors are attended to on a regular basis.

The data manager ensures that all the data is clean at all times. Questionnaire (Address and Intake) data is checked for entry accuracy at the time of scanning. In addition, data checking is done on the databases every week and the data manager takes responsibility for organizing the data checking process. A summary report on the data quality and data entry accuracy is then produced by the data manager and distributed to the project investigators.

Data checking is done using the following methods:

Ranges and Validity rules

A range of acceptable values and skip patterns (checks) has been inbuilt for all the appropriate variables during the programming of the data collection form using the xml form. Any values that fall outside this range cannot be accepted by the database. Validity rules are also set where certain variables can only be entered if they comply with a particular rule. For example, the database does not allow outside limit or blank entries for the patient identification numbers and all unreal dates are rejected. The data manager works with the data clerk to identify any inconsistent data on a weekly basis. Inconsistency checks are done when the files are in SPSS. The manager then consults with study clinicians to resolve the inconsistencies.

Line listings for data captured non-electronically

The main objectives of line listings include:

- identifying any errors made during data entry

- estimating the accuracy rate of the data clerk, monitor and assess his/her data entry performance

The data manager produces 10% line listings of all the enrollment and follow-up files on a quarterly basis. The data team members then check the line listings against the hard copy questionnaires. Errors are highlighted on the line listings and the error rate approximated thereafter. If the error rate is higher than 0.3% then the whole database should be checked.

After the checking has been completed, the data manager lists all the corrections that need to be done in the patient charts on the data entry sheet; this helps the data entry clerk easily identify the corrections that need to be done.

Missing values

Restrictions have been put on all questions that are required to be answered. These checks prevent the interviewer from skipping any question that requires a response.

Safety, Security, & Storage

Data security in ODK

- All data collection devices(Tablets) will have a security code for unlocking to prevent unauthorized users
- All the electronic forms will be encrypted before loading to the ODK collect
- The data will be uploaded to Nascop server through a secured path Private key will be used to decrypt the data. ONLY the Data manager will have access to the private key
- Once the data is decrypted , it will be converted into SPSS format.
- All databases are backed up every Friday by saving the most current files on two CD disks and external harddrive and uploaded to the UW-server. All computers are password-protected, preventing access by any unauthorized persons. Data is backed

up on a weekly basis by the data clerk and the data manager. Data is saved on an external hard-drive that remains in the office as well as on a CD that is kept off-site. All data on CD disks will be password protected by the data clerk. All patient records are filed according to numerical order of the patients' identification. Folders are kept in a locked storage cabinet at the study clinic for reference by research personnel. Once the subject completes the study or is lost to follow-up, her records are brought to the KNH office for storage. Participant study ID will be generated by the data manager who will be responsible for linking and de-linking data.

Schedule

Daily

Electronic questionnaire entry – The nurses will complete the forms electronically and upload (send) the data to the ODK server. The data will then be downloaded from the server and saved onto the data computer in SPSS format

Specimen entry – Specimens that are collected in clinic are entered into the *Clinic Specimen CollectionBook*. Results are entered in the *CD4 Results Book*, *Pap Smear Results Book*, and *Biopsy Results Book*.

Laboratory Procedures

Sample collection

- Study nurse will collect Pap smear specimens and cervical swabs for HPV and HIV
- Study doctor will collect cervical biopsy specimen under colposcopy and perform endocervical curettage when necessary
- Study nurse or lab assistant will collect blood specimens
- Specimens will be gathered in the laboratory before transfer
- Lab assistant or technologist will record details in lab book. Information includes:
 - Specimen Study ID
 - Date
 - Visit type
 - Time sample was collected
- Pap smear, cervical biopsy, and endocervical curettage specimens will be placed in a cool box and transported to the designated laboratory before 4 pm by the lab assistant or technologist. Designated laboratory will record reception of samples in a book including: specimen study ID, date, time, and number of specimens. Results will be filled in the relevant “Histology Report” and “Cytology Report”.
- One set of blood samples will be sent to the Coptic Hospital for CD4 count. These results will be filled in the “CD4 Report”.
- One set of blood samples will be centrifuged at the study laboratory by the lab technologist located at the Hope Center and frozen in a -80oC freezer for storage. Shipment of samples for analysis of viral levels will be performed on these samples.
- Cervical swabs for HIV will be prepared in the study laboratory and frozen in a -80oC freezer for storage. Cervical swabs for HPV will be stored in storage medium

(preservCyt) at $<25^{\circ}$ C. Shipment of samples for viral analysis will be performed on these samples.

Pap smear and cervical tissue biopsy

- Papanicolaou test: after the application of a speculum, a cervex brush will be inserted to its full length into the endocervical canal so that the shorter outer brush hairs are in contact with the ectocervix. The cervex brush will then be gently pushed whilst being turned 5 times in an anticlockwise direction (2.5 complete turns), in order to collect cells from both the endo- and ecto-cervix. Care will be taken not to touch the tip of the brush in order to avoid cross-contamination. Using the brush, a Pap smear will then be prepared by smearing the brush on a clean glass slide and fixing immediately with 95% isopropyl alcohol for at least 30 minutes at the study clinic. The slide will then be transported to the laboratory to be stained, and the cytology will be reported by a study pathologist from Aga Khan University Hospital using the Bethesda classification.
- Samples from the colposcopy-directed biopsy of the cervix, endocervical curettage specimen and the LEEP specimen will be fixed in 10% buffered formaldehyde solution and transported for haematoxylin-eosin staining. The histopathology results will be read by a study pathologist from Aga Khan University Hospital, and the biopsy samples will subsequently be preserved and fixed in paraffin. Biopsy blocks will be collected and may be sent to the IARC for further analysis.

Cervical swabs for HIV and HPV

- Dacron swabs will be used to collect samples for HIV from the cervix. Endocervical secretions will be collected by rotating the swab 360 degrees in the outer part of the endocervix.
- For HIV-1 DNA testing, the dry swab will be stored as such in the -80°C freezer.

- For HIV RNA testing, the swabs will be collected in a cryovial containing 1 ml of freezing media and then stored at -80°C.
- Cervical samples for HPV will be taken using cervex brush. The brush will be inserted to its full length into the endocervical canal so that the shorter outer brush hairs are in contact with the ectocervix. The cervex brush will then be gently pushed whilst being turned 5 times (2.5 complete turns), in order to collect cells from both the endo- and ectocervix. Care will be taken not to touch the tip of the brush in order to avoid cross-contamination. The brush containing cervical cellular material will be placed in a vial containing PreservCyt media (Cytoc Corporation) and labeled with the subject identification number of the participant. The brush will be fully rinsed in the media by pressing 10 times against the bottom of the vial, forcing the brush hairs to separate. Finally the brush will be vigorously shaken in the media to remove any residual cells. The brush will NOT be left in the vial, but discarded. It is very important to close the vial very tightly to avoid possible leakage during transport. Cell samples will be stored at the Hope Center at 25°C, and then later transported to the US or to IARC in Europe.

Shipping

- The UW has extensive experience measuring HIV-1 viral RNA from cervical swabs and plasma. While measuring plasma HIV-1 RNA viral levels is available in Kenya, HIV-1 RNA analysis from cervical swabs has not been performed here. In order to maintain high quality standards, samples will be analyzed in Seattle. Plasma HIV-1 RNA will also be analyzed in the same laboratory so that the results, taken from two different body compartments on the same day, are comparable. The KNH IRB has approved laboratory analysis of cervical samples in Seattle for a current study on cervical cancer screening and is therefore expected to approve similar testing for this study. Analysis of samples locally

is dependent on significant funding to develop a laboratory locally that could perform PCR analysis.

- Samples for HPV testing will be shipped to the IARC in France and then to the Netherlands or to the US directly where the PCR testing will be conducted. The KNH IRB has also approved laboratory analysis of cervical samples in Europe for a current study (mentioned above) and is therefore expected to approve similar testing for this study.

Analysis for HIV-1 viral levels

- HIV swabs will be stored at -80°C until shipment to Seattle. The swabs will be tested for HIV RNA using a Gen-Probe HIV-1 viral load assay (San Diego, California, USA) which has been validated for use in Kenyan HIV subtypes.⁴⁰ The assay has a lower limit of 50 copies/swab in genital secretions. HIV-1 DNA will be examined in the cervical swabs by detecting proviral DNA using a nested PCR for the viral *gag* region which should detect as little as 1 copy of DNA. This technique has been validated in a number of other studies.^{39, 41}

Analysis for HPV subtypes

- Frozen pellets of exfoliated cervical cells collected in PreservCyt will be extracted and tested following standard operating procedures of the HPV Laboratory at CDC as implemented under the ongoing Quality Management System. The Linear Array HPV Genotyping Test (Roche Diagnostics), based on L1 consensus PCR with type-specific hybridization, will be used. The assay detects 37 HPV types: HPV6, 11, 16, 18, 26, 31, 33, 35, 39, 40, 42, 45, 51, 52, 53, 54, 55, 56, 58, 59, 61, 62, 64, 66, 67, 68, 69, 70, 71, 72, 73, 81, 82, 83, 84, 89, and IS39. The probe for HPV 52 (called XR) cross reacts with

HPV 33, 35, and 58. If the XR band and any cross-reacting types are identified, HPV 52 will be identified with a type-specific quantitative PCR assay. Samples negative for both, beta-globin and all HPV types will be reported as negative. Results will be recorded in database and securely transmitted to Dr. Chung for linking to epidemiologic and clinic data.

Quality Assurance and Quality Control (QA/QC) for Cytology and Histology Specimens

A Cytology and Histology QA/QC procedures document is attached to the protocol as an addendum.

- Loss of slides or blocks: If any cytology or histology slides or blocks are misplaced during the course of QA/QC procedures, the patient will be contacted to return for repeat Pap smear or biopsy, as necessary.

Quality Assurance/Quality Control of Study Procedures

- *LEEP*: The senior gynecologist on the study will work closely with the study doctor to ensure quality procedures with a low rate of complications through regular weekly meetings and contact through cell phone and e-mail. The complication rate and type will be reviewed on a bi-monthly basis. This information will be collected by the “Adverse Event” form which has been added. All LEEP specimens will also be evaluated by histopathology with confirmatory reads as detailed above.
- *Cryotherapy*: As with LEEP, the study doctor will be well trained and mentored by the study gynecologist to perform quality procedures with low rates of complications, and the complication rate and type will be reviewed on a bi-monthly basis. This information will be collected by the “Adverse Event” form which has been added.

- *Colposcopy and endocervical curettage:* The study gynecologist will supervise the study doctor to conduct colposcopy with biopsy and curettage. He will receive support and in-service trainings as deemed necessary.

Ethics

A. INTRODUCTION

Field and laboratory procedures will be performed in Nairobi, Kenya, while data analysis will be done in both Nairobi and Seattle, Washington. The study will be reviewed by the Institutional Review Board (IRB) at the University of Washington and the Kenyatta National Hospital (KNH) Ethical Review Committee (ERC). The study will not recruit subjects prior to approval from both the University of Washington IRB and the KNH ERC. In accordance with the International Conference on Harmonisation Good Clinical Practices (ICHGCP) section 4.5.4, the investigators may deviate from protocol prior to IRB approval in order to eliminate immediate hazard to trial subjects.

B. DECEPTION

If any deception or withholding of complete information is required for this activity, explain why this is necessary and attach a protocol explaining if, how, when, and by whom subjects will be debriefed.

No deception or withholding of complete information is required for this activity.

C. SUBJECTS

1. How many subjects will you need to **complete** this study?

Number: approximately 400

Age range: above 18

2. Explain how you will achieve equitable subject representation in the following categories.

If not applicable, justify exclusions.

a. **Age (minors, elderly):** N/A. Sexually active women are at greatest risk for cervical cancer. Adolescents will be excluded due to their high rate of regression of cervical lesions, with 90% clearing the HPV virus by 24 months. In terms of cervical dysplasia, it was been shown that in adolescents with high grade lesions, a majority will clear these lesions within 1 to 3 years. The current ASCCP guidelines state that wome with HSIL cytology should be managed with colposcopy.

b. **Gender:** N/A. Cervical cancer affects the female reproductive system only

c. **Ethnic and racial minority populations:** N/A. Study is performed in Kenya where the majority of the population is black

3. What characteristics (inclusion criteria) must subjects have to be in this study? (Answer for each subject group, if different.)

HIV-positive, female, over the age of 18, intact cervix, initiation of sexual intercourse

4. What characteristics (exclusion criteria) would exclude subjects who are otherwise eligible from this study? (Answer for each subject group, if different.)

HIV-negative, male, below 18 years of age, pregnant by clinical history or physical examination, post-hysterectomy,

post-cervical cancer treatment, history of a bleeding disorder, no prior initiation of sexual intercourse

5. Describe the subject recruitment strategies you will use for each group of subjects. (Attach advertisements, flyers, contact letters, telephone contact protocols, Health Sciences recruitment web site template, etc.)

Adult female subjects who attend the Hope Center for Infectious Diseases will be informed by a doctor at the clinic of the study and their potential eligibility using the attached form (Pre-Screening Talking Points). If the subject expresses interest, the doctor will contact the study nurse who will accompany the subject to the study clinic located less than 50 yards away. We will also use recruitment leaflets that will be distributed by the study staff at the Hope Center Reception. The leaflets will have the contact information of the study staff.

6. Explain who will approach subjects to take part in the study and how this will be done to protect subjects' privacy.

Doctors from the Hope Center who see the subjects at a medical follow-up or screening visit will approach subjects to take part in the study in a confidential, private room. We will also use recruitment leaflets that will be distributed by the study staff at the Hope Center Reception. The leaflets will have the contact information of the study staff.

7. Explain what steps you will take during the recruitment process to minimize potential coercion or the appearance of coercion.

Doctors or study staff who inform subjects of the study will follow the "Pre-Screening Talking Points" and let them know that their decision whether to participate in the study or not will not affect their clinical care at the Hope Center. In addition, women who wish to be screened but do not want to enter the study will receive free screening. The study nurse will reiterate these points during the informational interview using the "Screening Talking Points" to gauge the interest of the patient before signing an informed consent. Talking Points are enclosed in the Appendix.

8. Will you give subjects gifts, payments, services without charge, or extra course credit?

No Yes If yes, explain:

Subjects will receive free treatment for pre-cancerous lesions amenable to cryotherapy or LEEP. Medical personnel will provide free medical services, gynecological examination, treatment, and free condoms. Randomized subjects will also receive money for transportation to return to clinic.

9. Will any of the subjects or their third-party payers be charged for any study procedures?

No Yes If yes, explain:

10. Where will the study procedures be carried out?

The study will be carried out in Nairobi, Kenya at the Coptic Hospital. IRB approval for this PHE has been granted by Kenyatta National Hospital (KNH) in Kenya and the University of Washington (UW).

D. RISKS AND BENEFITS

1. Describe nature and degree of risk of possible injury, stress, discomfort, invasionofprivacy, and other sideeffects from all study procedures, drugs and devices (standard and experimental), interviews and questionnaires. Include psycho-social risks as well as physiological risks. Include risks of withholding standard care or procedures if this is the case. Do not reference the consent form.

The study may collect personal information that may be embarrassing for the subject to talk about. As part of the study, the subject may meet other patients from this clinic whom she may know from outside the clinic. We will be collecting blood samples using a needle and syringe. The puncture of the needle may be uncomfortable and leave a bruise and may cause infection or fainting. Collection of biopsy samples through the pelvic exam may cause: mild discomfort as the subject is examined, small amount of bleeding from the vagina for 1-2 days, and mild to moderate cramping for 5 minutes that is similar to mild menstrual pain. If the subject undergoes cryotherapy or LEEP, then she may experience mild abdominal cramps for less than 10 minutes, discharge from the vagina for about 2 weeks, and/or bleeding for several days. Possible serious complications

include excessive bleeding or infection. Additionally, some studies have shown that cervical treatment is associated with pregnancy complications including premature rupture of membranes, premature contractions, infections and cervical incompetence.

If the subject undergoes a biopsy, she is requested not have sex for 10 days. The reason is that if her partner is not infected with HIV, then he may be at greater risk of becoming infected with HIV after the biopsy procedure because of a possibility of increased HIV shedding from the cervix. The participant may also be at increased risk of infection at the site of the biopsy. If the subject has received cryotherapy or LEEP, then, for the same reasons as the biopsy, we suggest that she not have sex for 4 weeks after it is performed.

Subjects will be prescribed antibiotics to protect against infection after cervical treatment or as treatment for an sexually transmitted infection or other vaginal infection discovered on pelvic exam. Possible side-effects from antibiotic use include (but are not limited to) nausea or anorexia, vomiting, diarrhea, photosensitivity, rash, anaphylaxis possibly leading to death, dizziness, headache, confusion, tinnitus or hearing loss, seizures, arrhythmias, neutropenia, thrombocytopenia, hyper/hypoglycemia, tendon

rupture, liver disease, kidney disease and peripheral neuropathy.

2. Explain what steps you will take to minimize risks of harm and to protect subjects' rights and welfare. (If you will include protected groups of subjects (minors, fetuses in utero, prisoners, pregnant women, decisionally impaired or economically or educationally disadvantaged subjects) please identify the group(s) and answer this question for each group.)

The clinical procedures of collecting blood, fluid, and tissue samples will only be performed by certified medical staff trained in these tasks. Any complications arising from these procedures will be handled by a doctor and covered by the study.

Study investigators may decide to withdraw a study participant from the study if they find further enrollment may expose the participant to harm or the investigator determines the participant will not be able to abide by study safety requirements, e.g. in the case of mental health problems or drug/alcohol dependency problems.

3. Is it possible that you will discover a subject's previously unknown condition (disease, suicidal intentions, genetic predisposition, etc.) as a result of study procedures?

No Yes If yes, explain how you will handle this situation.

The intent of this cervical cancer screening study is to detect pre-cancerous and cancerous lesions of the cervix which may not be known to the subject and then treat those

lesions free of charge. If the lesions are too large or are cancerous then we will refer them to the neighboring government hospital for subsidized care.

4. Describe the anticipated benefits of this research for individual subjects in each subject group. If none, state “None.”

The benefit of this research for the individual is that we may be able to detect and treat pre-cancerous disease of the cervix before it becomes cancerous and deadly. By participating in this study, an individual may avoid the development of a life-threatening disease.

5. Describe the anticipated benefits of this research for society, and explain how the benefits outweigh the risks.

The benefit of this research for society is that it may identify the most effective method to treat cervical dysplasia in HIV-positive women who are at much higher risk for cervical cancer and who number in the millions in resource-constrained settings around the world. Results of this research may impact international guidelines and the way millions of dollars of donor funding is spent on the care of HIV-positive women.

As discussed above, the medical care offered through the research conforms to the standard of care established by the Kenyan MOH at tertiary care and provincial level health care facilities. Options are available to clients enrolling in the study to obtain Pap smears at these locations along with

cryotherapy and LEEP treatment. At the same time, the treatment interventions that are being offered in the study do not offer unreasonable risks and provide likely clinical benefit. Both procedures are widely recommended in both developing and developed settings and one method is not known to be better than the other among HIV-positive women. As previously discussed, given the paucity of evidence-based literature there is scientific and medical equipoise in addressing this question and offering these two treatment methods.

The benefits of the procedures being offered outweigh the risks. The risks of LEEP and cryotherapy are low. In the study previously mentioned from Zambia, Pfaendler, et al. found that the overall complication rate of LEEP to be 3.7%, all of which was managed on-site in the clinic.²⁰ Likewise, in a study of cryotherapy in India, the overall complication rate was found to be 3.0%.²¹ Both LEEP and cryotherapy may result in infection or bleeding, though the rates are low and the great majority can be managed in the clinic where the procedure was performed. Pfaendler et al. found bleeding as a complication in 14/697 (2.0%) and infection in 12/697 (2.0%) while performing LEEP. Nene et al. in the study from India mentioned above had 9 (1.9%) cases of mild bleeding and infection in 8 (1.4%) cases. The benefits of treatment through LEEP and cryotherapy, on the other hand,

are very high. It has long been known that HIV-positive women are at higher risk of cervical disease and faster progression of lesions.^{42, 43} From preliminary studies performed on our HIV-positive patient population, we have found that the prevalence of high-grade lesions is around 8% in women 30-39 years old and taking antiretroviral medications.⁴⁴ Additionally, studies have shown that either method of treatment is extremely effective in treating CIN.²⁷

E. CONFIDENTIALITY OF RESEARCH DATA

1. Will you record any direct subject identifiers (names, Social Security numbers, patient, hospital, laboratory or claim numbers, addresses, telephone numbers, locator information, etc.) No Yes If yes, explain why this is necessary and describe the coding system you will use to protect against disclosure.

We will be recording names and assigning a study number that will be used on all study visits. This is to ensure accurate follow-up of study participants. This will be handled by the senior data analyst.

2. Will you retain a link between study code numbers and direct identifiers after the data collection is complete? No Yes If yes, explain why this is necessary and for how long you will keep this link.

The link between the study participant's name and study number is necessary to facilitate follow-up during the 1 month study period. This link will no longer be needed after follow-up is completed and will be removed after 5

years. This will be handled by the senior data analyst. Data and specimens will be stored for 10 years after completion of study follow-up before being destroyed.

3. Describe how you will protect data against disclosure to the public or to other researchers or non-researchers. Explain who (other than members of the research team) will have access to data (e.g., sponsors, advisers, government agencies, etc.).

All names and numbers will remain in confidential files that are accessible only to the investigators and study staff. Computer databases containing information about study subjects will be protected by passwords which allow access to only the investigators.

Notebooks, folders and CDs will move between the study clinic site and the KNH data office in a direct manner to minimize handling of data. Information will be transported by private vehicle used by the study. There will be two CDs to back up study data, one to be kept with the administrator at an off-site location and one with the senior data analysis in a locked closet at KNH. Data files on the computer and CDs will be password protected, accessible only by data staff. Additionally, patient identifying data will be kept in a separate, locked folder from the clinical data to maximize confidentiality when reading clinical materials. All data, including physical medical files, will be physically locked when not in transport - both at the study site and at the office at KNH - to assure that only

designated staff will have access to the files. The study coordinator will be responsible for ensuring patient confidentiality for both electronic data and the medical files.

4. Will you place a copy of the consent form or other study information in the subject's medical or other personal record?

No Yes. If yes, explain why this is necessary.

5. Do you anticipate using any data (information, specimens, etc.) from this study for other studies in the future?

No Yes If "Yes," explain and include this information in the consent form.

Specimens may be tested for HPV at a later date. Data and specimens are requested to be stored from the participant for 10 years following enrollment for possible use in other HIV and cervical studies in the future. Approval from the UW and KNH IRBs will be obtained before any of these studies are performed.

F. ADDITIONAL INFORMATION

1. If the study will involve radiation exposure to subjects, e.g., X-rays, radioisotopes:

Pending Approved NA

2. Will you need access to subjects' medical, academic, or other personal records for screening purposes or during this study?

No Yes. If yes, specify types of records, what information you will take from the records and how you will use them.

Subjects will be enrolled at the Hope Center for their HIV care and their medical records contain information on what type of treatment they have received and the severity of their condition. We will access these records to determine what type of antiretroviral treatment they've been exposed to and their clinical status according to medical examination and laboratory values including CD4 count. This information is included in the consent form.

3. Will you make audio-visual or tape recordings or photographs of subjects? No Yes. If yes, explain what type of recordings you will make, how long you will keep them, and if anyone other than the members of the research team will be able to see them.
4. Will your study involve use of equipment involving energy input to the subjects (EMG, EKG, MRI, ultrasound, etc.)?
 No Yes. If yes, attach documentation that all equipment will be tested regularly or describe safety testing procedures you will use.

G. REPORTING OF ADVERSE EVENTS

1. Describe how unanticipated adverse events related to study participation will be reported to the local IRB/Ethics Committee.

The IRBs of host institutions and the sponsor, CDC, will be notified of adverse events by the PI within 72 hours of his becoming aware if the adverse events fall into one of two categories:

1. related to research procedures and unexpected and severe

2. related to research procedures and expected, but more severe or occurring at a greater frequency than expected

Severe adverse events related to the study include any death, any non-HIV-related hospitalization, severe infection including PID, severe bleeding or cramping and severe cervical stenosis. Any adverse events will be recorded by study staff on Adverse Events Reporting Forms at the time of the incident or at the next study visit. Documentation of severe adverse events which meet the criteria listed above will be brought to the attention of study PIs within 24 hours by the study doctor and/or study nurse via phone or email, and be assessed.

Other Adverse Events that do not meet the criteria above will be reported to the IRBs as per their specific guidelines.. All serious and non-serious adverse events will be reported to the DSMB during its regular meetings. All Adverse Event reports that are sent to the IRBs and DSMB will also be sent to the study sponsor.

Adverse Events

Specification of Safety Parameters

Safety parameters for this study will include signs and symptoms of local genital irritation and of systemic effects that could be related to either Pap smear screening or study treatment procedures (LEEP or cryotherapy), side effects from any medications prescribed as part of this study, and any other health complications that subjects may experience while enrolled in this study. Methods and Timing for Assessing, Recording, and Analyzing Safety Parameters.

Adverse Events (AEs)

An AE is any untoward medical occurrence or unintended clinical sign, including an abnormal laboratory finding, symptom or disease, in a clinical investigation subject that occurs during the course of the study. The occurrence is considered an AE whether it is associated with the use of a medical treatment or procedure, or considered unrelated to that medical treatment or procedure. The occurrence of an AE may come to the attention of study personnel in various ways--during study visits, during interviews of a study participant presenting for medical care, or during a review by a study monitor.

All AEs will be:

- recorded on the appropriate AE CRF by the study physician and nurses
- summarized by the data team
- followed through resolution by a study clinician
- reviewed and evaluated by a study clinician
- Study-related SAEs will be immediately reported to the host IRBs and ERCs, and to the study sponsor.

In this study, the following situations will be considered AEs:

1. Occurrences related to Pap smear, LEEP or cryotherapy procedures:

Symptoms such as abdominal pain that lasts longer than 2-3 days, vaginal bleeding and/or discharge, fever, or chills will be assessed at each study visit after obtaining either Pap smear or cervical biopsy.

2. Occurrences related to side effects from prescribed medications related to the study:

If a medication(s) are prescribed for treatment of an infection, subjects will be assessed for any potential side-effects at each visit. These symptoms may include: upset stomach, vomiting or diarrhea, sensitivity to light, rash, severe allergic reaction that could cause death, dizziness, headache, confusion, ringing in the ears or hearing loss, seizures, heart problems, blood disorders, problems with blood sugar, liver disease, kidney disease, and pain or numbness in the arms or legs.

3. New medical problem(s) or worsening of an existing medical problem(s)

Study staff will inquire about any new medical problem or worsening of an existing medical problem since the subject's last visit.

Any medical condition that is present at the time of the Enrollment Visit should be considered as the baseline for this pre-existing condition and not reported as an adverse event. However, if there is an increase in the frequency or severity of the condition, it will be recorded as an adverse event. Anticipated day-to-day fluctuations of pre-existing conditions, which do not represent clinically significant exacerbation, will not be considered adverse events.

All AEs will be graded for severity and relationship to study procedures or treatment.

Classification of Severity of Adverse Event:

For adverse events that do not fall under the three categories listed above (occurrences related to study procedures, medications, or new or worsening existing medical problems), the following guidelines will be used to quantify severity:

Mild: adverse events that require minimal or no treatment and do not interfere with the patient's daily activities.

Moderate: adverse events that result in a low level of inconvenience for the patient's daily activities or concern with the study treatment or procedures. Moderate events may cause some interference with functioning.

Severe: adverse events that interrupt a patient's usual daily activity and may require systemic drug therapy or other treatment. Severe events are usually incapacitating.

Life threatening: any adverse event that places the patient or subject, in the view of the investigator, at immediate risk of death from the reaction as it occurred, i.e., it does not include a reaction that, had it occurred in a more severe form, might have caused death.

Changes in the classification of severity of an AE should be documented to allow an assessment of the duration of the event at each level of intensity. Adverse events characterized as intermittent require documentation of onset and duration of each episode.

All AEs will be categorized by the study physician.

Relationship of Adverse Event to Study Procedures:

All AEs must have their relationship to study procedures assessed by the study clinician.

The terms used to assess the relationship of an AE to the study procedures are:

1. **Related-** There is a reasonable possibility that the AE may be related to the study agent.
2. **Not Related-** There is not a reasonable possibility that the AE is related to the study agent.

When an AE is assessed as “not related” to study agent(s), and alternative etiology, diagnosis, or explanation for the AE should be provided.

Outcome of Adverse Event:

All AEs must have their outcome assessed as either “resolved without sequelae”, “resolved with sequelae”, “ongoing”, “death”, “unknown.” The study clinician is responsible for assessing outcomes and recording them on the appropriate CRF.

Definition of Serious Adverse Event (SAE)

An SAE is defined as an AE that meets one of the following conditions:

- Death during the period of protocol defined surveillance
- Life-threatening event (defined as a subject at immediate risk of death at the time of the event)
- An event requiring inpatient hospitalization or prolongation of existing hospitalization during the period of protocol defined surveillance
- Results in congenital anomaly or birth defect
- Results in a persistent or significant disability/incapacity
- Any other important medical event that may not result in death, be life threatening, or require hospitalization, may be considered a serious adverse experience when, based

upon appropriate medical judgment, the event may jeopardize the subject and may require medical or surgical intervention to prevent one of the outcomes listed above.

Data Safety and Monitoring Plan

Introduction

The data safety and monitoring board (DSMB) will act in advisory capacity to the CDC, UW, KNH, UoN, Coptic Hospital, and IARC – WHO to monitor patient safety and evaluate treatment interventions for this study. Dr. Scott McClelland (UW) will be Chairman of the DSMB.

After its first meeting around study initiation, the DSMB will be responsible for reviewing interim safety and efficacy analyses at five time points during the 3 year study period. These meetings and reviews of interim analyses are expected to occur at approximately 4, 9, 15, 21, and 27 months after study initiation.

- 1) The first meeting will take place when 100 participants have received the study intervention, and this is expected to occur around 4 months after study initiation.
- 2) The second meeting will take place when 75 participants have received Month 6 Pap smear results, and this is expected to occur around 9 months after study initiation.
- 3) Thereafter, DSMB meetings and interim analyses will occur every 6 months after this second meeting or approximately at 15, 21, and 27 months after study initiation.

The DSMB will have the following responsibilities:

1. Review the research protocol and plans for data safety and monitoring;

2. Evaluate the progress of the trials, participant recruitment, accrual and retention, participant risk versus benefit, and other factors that can affect study outcome;
3. Protect the safety of the study participants and review interim or cumulative data for evidence of adverse events;
4. Review safety and progress report from an unblinded statistician who will use both blinded and unblinded data;
5. Make recommendations to the institutions involved and the PIs concerning continuation, termination or other modifications of the trials based on the observed beneficial or adverse effects of the treatment under study;
6. Review report on interim analysis of efficacy in accordance with stopping rules which are clearly defined in advance of data analysis and have the approval of the DSMB;
7. Ensure the confidentiality of the trial's data and the results of monitoring; and
8. Review issues that have been identified by the study team and upon request by the study team, review problems that are identified by the monitors in relation to patient safety.

DSMB Recommendations

The DSMB may conclude each review with recommendations to continue the trial without change, modification of the trial, or termination of the trial based on pre-defined criteria established at the beginning of the trial.

Recommendations for modification of the design and conduct of the trial may include:

1. Modifications of the study protocol based upon the review of the safety data;

2. Suspension or early termination of the study because of serious concerns about subjects' safety, inadequate performance or rate of enrollment;
3. Suspension or early termination of the study because study objectives have been obtained according to pre-established statistical guidelines;
4. Optional approaches for trial design when the DSMB determines that the incidence of primary study outcomes is substantially less than expected, such as recommendations to increase the number of trial centers or extend the recruitment period/follow-up period; and,

Confidentiality

Confidentiality will be maintained during all phases of DSMB review and deliberations. Only voting members of the DSMB will have access to interim analyses of outcome data by treatment group. Exceptions may be made when the DSMB deems it appropriate. DSMB members must maintain strict confidentiality concerning all privileged trial results ever provided to them. The DSMB will review data only by masked study group (such as 'Intervention 1' versus 'Intervention 2' rather than cryotherapy versus LEEP) unless the DSMB determines that the identities of the groups are necessary for their decision-making. Any request to unmask data must be made in writing.

Membership

The DSMB will be composed of four members chosen from both the U.S. and Kenya.

The members include:

1. R. Scott McClelland, MD, MPH, Associate Professor of Medicine, Epidemiology and Global Health, UW;
2. Elizabeth Brown, ScD, Associate Member, FHCRC, Research Associate Professor, Department of Biostatistics, UW;

3. James Kiarie, MBChB, MMed, MPH, Professor of Obstetrics/Gynecology, KNH; and
4. David Eschenbach, MD, Professor, Women's Health, Chairman, Department of Obstetrics and Gynecology, UW
5. *Ad hoc* specialists may be invited to participate as non-voting members at any time if additional expertise is desired.

Meetings

The first meeting will take place either prior to trial initiation or early after the trial has been initiated to discuss the protocol. A designated DSMB member and the PIs will prepare the agenda to review initiation of the trial and reporting of adverse events. The DSMB will also review monitoring guidelines and approve or give recommendations..

Once the trial has been initiated, the DSMB will meet, as outlined above, to examine the accumulated safety and enrollment data, review study progress and discuss other factors (internal or external to the study) that might impact continuation of the trials as designed. A DSMB meeting may be requested by DSMB members or the Principal Investigators at any time to discuss safety concerns, and includes the occurrence of any Significant Adverse Event (SAE) that is associated with the study. The study team will provide the logistical management and support of the DSMB meetings. The meetings will be convened by teleconference to decrease travel cost. An emergency meeting of the DSMB may be called at any time should questions of patient safety arise.

Meeting Format

The meetings will mainly be open sessions. These sessions will be attended by the Principal Investigators or designee. Other research staff may attend the open sessions but this is up to the discretion of the PI. Issues discussed at open sessions will include conduct and

progress of the study, including patient accrual, compliance with protocol, and problems encountered. Patient-specific data and treatment group data will not be presented in these sessions.

Closed sessions may be requested by the DSMB at any time and will be attended by voting DSMB members and the unblinded statistician. Any other trial staff may be requested to attend by the DSMB. All safety and efficacy data will be presented at this session. The discussion at the closed session is completely confidential.

Should the DSMB decide to issue a termination recommendation, the full vote of the DSMB will be required. In the event of a split vote, majority vote will rule and a minority report will be appended. An appeal may be filed by study PIs if a termination decision is made.

Study Stopping Criteria

The DSMB may recommend stopping the study for the following reasons:

- The data show a significantly increased risk of serious adverse effects in one of the treatment arms.
- Interim efficacy analyses show significant treatment benefits or futility in the one treatment group. The interim efficacy analyses are based on pre-specified stopping boundaries for the primary endpoint of the study which preserve the study wide Type I error rate.
- It becomes clear that successful completion of the study is not feasible (e.g. there is an excess of patient dropout, missing data, lack of recruitment etc).

Interim Efficacy Analyses

This study will employ interim analyses to assess accumulating study data for early evidence of treatment efficacy. The primary outcome, recurrence of cervical intraepithelial neoplasia grade 2 or 3, will be compared between groups at the following approximate times: months 9, 15, 21 and 27, and for DSMB safety, upon cumulative enrollment and intervention of 100 patients. The decision criteria for stopping the study at each interim analysis are based on O'Brien-Fleming superiority boundary with Type I error controlled at alpha-level $\alpha = 0.05$. In the event that interim analyses are conducted at times other than the preplanned times (e.g. unequal information accrual) the stopping criteria will be adjusted to maintain overall type I error rate of $\alpha = 0.05$. We will use Lan-DeMets alpha spending approach, to make the necessary adjustments. The decision criteria for stoppage are based on a power of 80% to detect a treatment group difference at the end of study.

Report

A formal report prepared by an assigned administrator will be reviewed and approved by the DSMB chair, who will prepare a summary. The minutes and summary will be sent to the full DSMB within three weeks of the meeting. Once approved by the DSMB, the chair of the DSMB will sign on behalf of the board and the report will be forwarded to the participating institutions within 4 weeks of each meeting. The PIs will submit the results of these meetings to the UW Institutional Review Board (IRB), the KNH Ethics Review Committee (ERC), and the Centers for Disease Control (CDC) which is the study sponsor. Each report will conclude with a recommendation to continue or to terminate the study. This recommendation will be made by formal majority vote. A termination recommendation may be made by the DSMB at any time by majority vote. The report will not include un-blinded data or any discussion of the un-blinded data.

Any new findings discovered by the DSMB during the course of the study that may affect the willingness of subjects to remain in the study will be shared by the study doctor or nurse through direct discussions and printed pamphlets.

Sponsor Monitoring

As the study sponsor, the Centers for Disease Control (CDC) may conduct monitoring or auditing of study activities to ensure the scientific integrity of the study and to ensure the rights and protection of study participants. Monitoring and auditing activities may be conducted by:

- CDC staff (“internal”)
- authorized representatives of CDC (e.g., a contracted party considered to be “external”)
- both internal and external parties

Monitoring or auditing may be performed by means of on-site visits to the Investigator’s facilities or through other communications such as telephone calls or written correspondence. The visits will be scheduled at mutually agreeable times, and the frequency of visits will be at the discretion of CDC. During the visit, any study-related materials may be reviewed and the Investigator along with study staff should be available for discussion of findings.

The study may also be subject to inspection by regulatory authorities (national or foreign) as well as the IECs/IRBs to review compliance and regulatory requirements.

Acronyms

AIDS – Acquired Immunodeficiency Syndrome

CCSP – Cervical Cancer Screening Program, Coptic Hope Center for Infectious Diseases

CDC – Centers for Disease Control

CIN – Cervical Intraepithelial Neoplasia

CIS – Carcinoma in situ

CVL – Cervicovaginal Lavage

HIV – Human Immunodeficiency Virus

HPV – Human Papillomavirus

IRB – Institutional Review Board

KNH – Kenyatta National Hospital

LEEP – Loop Electrosurgical Excision Procedure

OGAC – Office of the Global AIDS Coordinator

Pap – Papanicolaou

PEPFAR – President’s Emergency Plan for AIDS Relief

PHE – Public Health Evaluation

RNA – Ribonucleic Acid

UNITID – University of Nairobi Institute of Tropical and Infectious Diseases

UoN – University of Nairobi

USA – United States of America

USG – United States Government

UW – University of Washington

VIA – Visual Inspection with Acetic Acid

Pre-Screening Talking Points for Clinic Staff

Dr. Michael Chung is conducting a research study at the Hope Center. This study examines what methods may best treat disease in a woman's private parts called the cervix. This disease is like a wound on the skin and can go away by itself. But in some cases, especially in women who have HIV, these wounds might become cancer. The study wants to see how treatment can prevent this problem from becoming cancer and how treatment might affect HIV. Those patients who enroll in the study will be given free screening for cervical disease that may develop into cancer and will also provide free treatment. Dr. Chung and other doctors from the University of Washington in America, the University of Nairobi, and Coptic Hospital are leading this study. You appear to be eligible to be in this study and can possibly enroll in this trial if you like. The study offers free screening for cervical disease and if treatment is received, further follow-up free testing and treatment over 2 years.

Participation in the research study is voluntary and does not affect your medical care at the Hope Center in any way. Free screening for cervical disease is also available at the Hope Center Cervical Cancer Screening Program even if you do not participate in this study.

The study may help you detect cervical disease that may be treated and prevent cervical cancer in the future. Are you interested in learning more about the study from the study nurse or doctor? They can explain the study in more detail if you are interested. If you are, we will send you to the study clinic right now. If not, you may go home or continue receiving care here from the pharmacy and laboratory.

Screening Talking Points for Study Staff

You are being invited to participate in a research study at the Hope Center. The study is being conducted by Dr. Michael Chung and other doctors from the University of Washington in America, Kenyatta National Hospital, and the Coptic Hospital. This study examines what methods may best treat disease of the female private parts called the cervix, and how treatment affects HIV levels. This disease is like a wound on the skin and can go away by itself. But in some cases, especially in women who have HIV, these wounds might become cancer.

You do not have to join the study. Whether or not you join the study will not impact your care at the Hope Center in any way. If you are eligible and participate in the study, you will receive free screening and treatment that will help detect and remove disease from your cervix that may lead to cancer. You will be asked questions and undergo a pelvic examination where our doctor and nurse can examine you to detect any areas that look like disease. After undergoing screening, we will ask you to return in 2 weeks to receive the results from the test. Most likely you will not need further treatment or tests after this. If, however, we find a result that might be disease, we will conduct another test to confirm the disease and the need for follow-up treatment.

If you are positive for cervical disease, you may enroll in a study to receive treatment. If you do, we will ask you to return every week for 3 weeks to examine you and test for any HIV that may be shed from your cervix after treatment. We will follow you for the next 2 years at 6 months intervals to test if there is any further disease. If it does, we will treat you. We will also draw blood from you in order to

see how much HIV is in your blood and what is your CD4 count. All of this is free of cost.

Free screening for cervical disease is also available at the Hope Center Cervical Cancer Screening Program regardless of whether you participate in this study or not. You are referred to us because the clinical officer has determined that you are a woman who may be eligible for this study.

- Are you over 18 years of age?
- Are you pregnant?
- Have you had a hysterectomy?
- Have you ever received treatment for cervical disease?
- Have you ever had a problem with bleeding?
- Have you initiated sexual intercourse?

If you are interested in the study, I will explain more about it from the informed consent form which I will give or read to you. If you still want to be in the study after reading or being read the informed consent, you can sign the form and we can enroll you in the study

Post-Medical Care Information

Cervical Cancer Screening

Pap smear or biopsy

After receiving a Pap smear or biopsy, you may have any of the following symptoms which are normal:

- Slight belly discomfort (like menstrual cramps, should not last more than 1-2 days)
- Slight bleeding from the vagina

We recommend you take paracetamol or ibuprofen for pain or cramps.

It is very uncommon to have severe problems but if you experience any of the following, **please notify the clinic or screening staff** (see contacts below) right away:

- Pain in the belly that lasts longer than 2-3 days
- Much bleeding from the vagina
- Bleeding from the vagina that is increasing in amount, or comes with belly pain
- Fever, chills

After a biopsy, it is important to wait 10 days before having sex. This will protect you and your partner from infection. (Please discuss with the screening nurse or doctor if you cannot wait and don't forget to take condoms!)

Treatment (cryotherapy or LEEP)

After receiving treatment, you may have any of the following symptoms which are normal:

- Belly pain like during your period (should not last longer than 1-2 days)
- Slight bleeding from the vagina for up to 1 week
- Clear fluid from the vagina (as long as 2 weeks)

The following symptoms are not normal and you should **contact screening staff** and **have medical attention as soon as possible** if you have:

- Severe belly pain
- Bleeding from the vagina that continues or is a large amount
- Fever, chills
- Cloudy (white) fluid from the vagina

After treatment, it is important to wait 4 weeks before having sex. This will protect you and your partner from infection. (Please discuss with the screening nurse or doctor if you cannot wait and don't forget to take condoms!)

Medications

You may receive medications (antibiotics) after treatment or to treat an infection found during your exams. Please follow the directions about how to take the medications carefully. If you have a severe problem after taking the medication or have any of the follow symptoms, please **stop the medication and contact screening staff**:

- upset stomach, vomiting
- severe diarrhea or diarrhea with blood
- sensitivity to light, rash
- dizziness, severe headache, confusion, ringing in the ears or hearing loss
- seizures (uncontrolled jerking of the body)
- heart problems

Screening Staff Contacts:

****EMERGENCY NUMBERS (24 HOURS): 020-272-2710 or 0733-771-288**

- Office 1: 0728-456-540
- Office 2: 020-271-2947
- Dr. Michael Chung: 020-271-2947
- Dr. Nelly Mugo: 020-273-6744
- Dr. Samir Sakr: 020-272-4737

Information for Spouses and Partners **Cervical Cancer Screening and Treatment**

Treatment (cryotherapy or LEEP)

Your wife or partner has undergone the following treatment as part of a cervical cancer screening and treatment program:

Cryotherapy

LEEP

After treatment, it is important to wait 4 weeks before having sex. This is because HIV shedding may increase substantially (but temporarily) at the site of cryotherapy or LEEP. This shedding may increase the risk of HIV transmission to an uninfected partner or lead to HIV re-infection. If abstinence is impossible during the healing period, it is important to use a condom every time you have sex for at least 4 weeks after treatment.

Screening Staff Contacts:

If you need more information, you may call the following numbers Mon-Fri between 8am and 5pm: Office 1: 0728-456-540

- Office 2: 020-271-2947
- Cell: 0721-289-733
- Dr. Samir Sakr: 020-272-4737

Maelezo kwa mabwana na wapenzi.

Ukaguzi na matibabu ya saratani ya mlango wa kizazi.

Matibabu ya Cryotherapy au LEEP

Mke au mpenzi wako amefanyiwa matibabu yafuatayo kama mojawapo wa mpangilio wa ukaguzi na matibabu ya saratani ya mlango wa kizazi:

Cryotherapy

LEEP

Baada ya matibabu ni muhimu usionane kimwili na mke au mpenzi wako kwa mda wa wiki nne. Hi ni kwa sababu idadi ya virusi vinanyosababisha ukimwi inaweza kuongezeka (kwa muda tu) katika sehemu iliyofanyiwa Cryotherapy au LEEP. Ongezeko la idadi ya virusi kwa mlango wa kizazi inaweza kuongezea hadhari ya kuambukiza mpenzi ambaye hajaambukizwa, au ongezeko la uambukizi mpya wa virusi vinavyosababisha ukimwi. Iwapo haiwezekani kujinyima kufanya mapenzi wakati huu ambapo mke au mpenzi wako anaendelea kupata nafuu, ni muhimu kutumia mpira wa kondomu kila wakati mnapoonana kimwili kwa mda wa wiki nne, baada ya kupokea matibabu.

Numbari za mawasiliano :

Iwapo ungependa kupewa maelezo zaidi, unaweza kupiga simu kwa numbari zifuatazo Jumatatu hadi ijumaa kuanzia saa mbili asubuhi mpaka saa kumi na moja jioni.

Afisi 1: 0728-456-540

Afisi 2: 020-271-2947

- Rununu: 0721-289-733

STUDY WRITTEN CONSENT FORM

Cervical Treatment Study: Screening *INITIAL CONSENT*

Full Title:

Impact of Cryotherapy versus Loop Electrosurgical Excision Procedure (LEEP) on Recurrence of Cervical Intraepithelial Neoplasia and HIV-1 Cervical Shedding among HIV-positive Women

Study Investigators:

Michael H. Chung, MD, MPH, Assistant Professor, Department of Global Health, University of Washington, 020 271-2947

Nelly Mugo, MBChB, MMed, MPH, Gynecologist, Department of Obstetrics and Gynecology, Kenyatta National Hospital, 020 273-6744

Samah Sakr, MBChB, Medical Director, Coptic Hospital of Kenya, 020 272-4737

Hugo De Vuyst, MD, PhD, Epidemiologist, International Agency for Research on Cancer, World Health Organization, +33-472738521

Silvia Franceschi, MD, Epidemiologist, International Agency for Research on Cancer, World Health Organization, +33-4728404

Barbra Richardson, PhD, Research Professor, Department of Biostatistics, University of Washington, +1-206-731-2425

Grace John-Stewart, MD, PhD, Professor, Department of Medicine, University of Washington, +1-206-543-4278

Emergency telephone number staffed 24 hours a day: 020 271 2947, 0723 914 057 or 0721 289 733

Ethical Review Committee Chairperson: Professor A. N. Guantai, 020 2726300, Ext. 44102, 4435544355, can be contacted for questions about research subject rights

Researchers' Statement

We are asking you to be in a research study. The purpose of this consent form is to give you information so you can decide if you want to be in the study. Please read the form carefully. You may ask questions about the purpose of the research and what we would ask you to do for the study. You may ask about possible risks and benefits, your rights as a volunteer, and anything else about the research or this form. You may ask questions at any time (before, during, and after the study) about anything. When we have answered all your questions, you can decide if you want to be in the study or not. This process is called 'informed consent.' If you wish, we will give you a copy of this form. Please let us know if you would rather use the Kiswahili consent form.

Purpose of the Study

The reason we are doing this research project is to screen for disease which may lead to cervical cancer. Participation in this project lasts for up to 1 month. You were asked to participate in this project because you are:

- HIV-positive,
- currently receiving care at the Hope clinic,
- not pregnant,
- do not have a history of problems with bleeding,
- have not had a hysterectomy (an operation to remove the uterus),
- and have initiated sexual intercourse.

Cervical cancer is the most common cancer among young and middle-aged women in Kenya. More women who have HIV get cervical cancer than those who do not have HIV. Even in HIV-infected women, cervical cancer is not common. However, in HIV-infected women, cervical disease that is not cancer is common. This disease is caused by a virus called the human papillomavirus (HPV). It is very important to find this disease and treat it before it becomes cancer. This study will screen you for cervical disease. If you screen positive for cervical disease, you may choose to enroll in a treatment study, or you will be referred elsewhere for treatment. The treatment study will be explained to you in detail before you decide. This screening study will have over 2,400 participants. By doing this study, we hope to provide free and comprehensive screening for HIV-positive women to prevent cervical cancer.

Procedures (see Appendix)

All participants will be asked to come for 1 to 3 visits over 1 month. All study visits will take from 20 to 40 minutes (Visit 1: consent process – 10 minutes, Pap smear – 20 minutes; Visit 2: review of Pap smear results – 10 minutes, colposcopy and biopsy – 30 minutes; Visit 3: review of biopsy results and discussion of treatment options – 30 minutes)

Visit 1:

If you agree, you will first undergo a pelvic exam with a Pap smear to see if you have a something like a wound (called a lesion). This lesion may possibly develop into cancer. A pelvic exam means that a doctor or nurse will examine your female parts. The Pap smear involves brushing the cervix with a small brush to collect material that can be looked at under a microscope (like a large magnifying glass).

Visit 2:

You will then return to this clinic 2 weeks later to find out the results of this test. Most of the time the results will show no disease or that we just have to repeat the Pap smear or visual inspection (VIA) every 6 months. If this is the case, then you will exit the study at this point and receive further follow-up at the Coptic Hope Cervical Cancer Screening Program (CCSP) or be asked to come back to the study clinic for a repeat Pap smear after 6 months. If we find lesions that need to be treated at these visits, we will discuss your treatment options with you in detail.

If you are screened and we find a lesion that needs to be looked at carefully, you will first have a pregnancy test. If you are pregnant, nothing more will be done until after you deliver. We will send you for special care at the Hope center to make sure you do not pass HIV to your baby (this care is called PMTCT), and you will exit the study and receive further follow-up at the CCSP. If you are not pregnant, we will look at your cervix with a type of magnifying lens called a colposcope. We will look closely at the cervix and take a piece out, about the size of a grain of rice. This is called a biopsy. We will look at this tissue biopsy under a microscope and share the results with you 4-6weeks later. This test is very good and will help us decide whether you need to have treatment.

There are some cells found in the cervix and the lining of the uterus. These are called ‘glandular cells.’ When Pap smear results show that your glandular cells are abnormal, the doctor will perform a procedure referred to as ‘Endocervical Curettage (ECC).’ This is a procedure where a spoon-shaped instrument called a ‘curette’ is used to scrape abnormal material from the passageway between the cervix and the uterus. This procedure obtains a small sample, which is then sent to the lab to be examined for abnormal cells. ECC is performed during colposcopy and takes just a few minutes to perform. You can expect to feel mild cramping, much like menstrual cramps following the procedure. If you are found to have abnormal cells but the doctor cannot see them by means of a colposcope, you will undergo ECC as described above.

Visit 3: You will return to this clinic in 4-6weeks for the biopsy results. If we find cervical disease that needs treatment, we will discuss your treatment options with you. You may choose to enroll in a further treatment study, which will be explained in detail at this time. If you prefer not to enroll in another study, you will be referred for treatment at Kenyatta National Hospital (KNH) and standard care at Coptic Hope Center. Whether you choose to accept study treatment or referral, you will be exited from this screening study at your third visit.

Contacting Participants

We will ask you to give us contact information like a phone number so we can call you if you do not come to a scheduled visit. We may ask about your health or where you are during these calls. It is important for you to come to all of your scheduled visits. If we cannot contact you by phone, we may try to visit you at your house. If this happens, we will not wear clothes that show we are health workers. You should tell us if you do not want to be contacted in this way. If you do not return for your Pap smear or biopsy results, we will also “flag” your Coptic Hope file so that you can be notified that you have available study results at your next clinic visit.

If study quality control procedures indicate that your Pap smear result is discordant from the original result you were given, we will ask you to return to the clinic to receive your new results and to discuss potential treatment options. At this point, you may be asked if you would like to re-enroll in the study.

Risks and discomforts of being in the study

This is an explanation of problems you may have through your participation in this study. Other problems not listed may happen as well.

Screening

Collection of samples through the pelvic exam may cause:

- mild discomfort as you are examined,
- a small amount of bleeding from the vagina for 1-2 days afterwards and
- mild to moderate cramping for around 5 minutes that is similar to mild period pain.

If you receive a biopsy, we ask that you do not have sex for 10 days. The reason is that if you are HIV-infected and your partner is not, then he may be at greater risk of becoming infected with HIV. If your partner is HIV positive, he may be at greater risk of getting re-infected with resistant HIV. Also, you need time to heal and will put yourself at higher risk of infection unless you give yourself this time. If you need help talking about these issues with your partner, a nurse, doctor or counselor can help you. We will give you free condoms to use if it is impossible for you to not have sex.

Another possible discomfort you may face is the worry or anxiety that you may have disease on your cervix. You may talk about this with a study nurse or doctor or if you would like talk to a counselor, we can help to arrange this.

We may find on screening that you have an infection. In this case we will give you a prescription for antibiotics. Side effects of these antibiotics may include: upset stomach, vomiting or diarrhea, sensitivity to light, rash, severe allergic reaction that could cause death, dizziness, headache, confusion, ringing in the ears or hearing loss, seizures (jerking of the body), heart problems, blood disorders, problems with blood sugar, liver disease, kidney disease and pain or numbness in your legs or arms. If you experience any of these problems, please report to study staff right away.

Confidentiality

The study staff will ask you for personal information that may be embarrassing to talk about like how many sexual partners you have had. You may choose not to answer any question. As part of the study, you may meet other patients from this clinic whom you know from outside the clinic. We have no plans to release your information to anyone other than the study researchers or appointed monitors. Sometimes committees that oversee research will examine study information to make sure nothing illegal or unethical is being done. Your personal information will be protected if this happens and will not be shown to anyone outside of this review. As the study sponsor, CDC may monitor or audit study activities. The reason for this would be to make sure that the study is being done the way it is supposed to be done. It would also make sure that your rights and health are protected. Your personal medical information will be kept confidential.

Alternative to taking part in this study

If you choose not to take part in this study, you will continue to receive medical care at the Coptic Hope Center and free antiretroviral medications. You may also receive free cervical cancer screening at this clinic without having to enroll in this study.

Benefits of the study

Your participation will help us understand more about cervical disease. By participating in this study, you receive free screening for cervical disease. If you would like to know the results of the study you can contact the study office 6 months or 1 year after you leave the study.

Compensation for injury

There is no cost to you for participating in this study other than your time. The study will pay for all screening costs for tests provided at the study clinic. If any physical injuries happen to you as a result of study participation, the study will cover the costs of care. If you think you have an injury or illness related to this study, contact the study staff (Dr. Michael Chung (020-272-2710) or Dr. Nelly Mugo (020-273-6744) or Dr. Evans Malava (0721 289 733) right away. They will treat you or refer you for treatment.

Specimen and Data Storage and Use of your Samples for Future Studies

We would like to save your medical information and Pap smear, colposcopy and endocervical curettage samples in Kenya at the Coptic Hospital and Kenyatta National Hospital for future research. This research may be done by the University of Washington or by other researchers who are working with us on this study for ten years after the end of follow-up in the study. We will use these data and samples only for research related to cervical cancer and HIV. Before your samples leave the clinic, they will be assigned a code. Your name will not be on them. Your name will be linked to the code only for five years after the study is completed. After that time, the link between your name and the code on your samples and data will be destroyed. The Institutional Review Boards are committees that watch over the safety and rights of research participants at Kenyatta National Hospital and the University of Washington. They must approve any future research studies using your samples. If you do not want to have your samples saved for future research, you can still be in this study and your samples will be destroyed once testing for the study is completed. If you agree to store your samples now, but change your mind before the end of the study, let the study staff know and we will make sure that your samples do not get stored for future research. We will not sell your samples. Tests done on your samples may lead to a new invention or discovery. We have no plans to share any money or other benefits resulting from this invention or discovery with you.

Other information

Your medical information is confidential (or kept secret) and we will keep your records in a locked office. Your medical records and information about your participation in the research will be available to you and to the study team but not to anyone outside of the study without your agreement. If you agree, we will share the information from this study with your doctors at the Hope Center. This information may help them treat your HIV and give you better care. All of your records will be kept in locked areas and all computer information will be password protected.

University of Washington staff sometimes review studies such as this one to make sure they are being done safely and legally. If a review of this study takes place, your records may be examined. You may refuse to participate or may leave from the study at any time without penalty or loss of benefit or help to which you have a right to. We will tell you if there is any new information about the treatments we are studying so you can decide if you want to leave the study. Your relationship with staff and services at Coptic Hope Center for Infectious Diseases will not be affected in any way if you do or do not participate or if you enter the program and leave later. Please inform study staff if you decide you would like to leave the study. You may be asked to give some final samples but you may refuse.

Study staff may decide to take you out the study if they find you may be harmed if you continue to participate. You may also be taken out of the study if the staff think you will not be able to follow study safety requirements.

Please contact Dr. Michael Chung (020 271-2947), Dr. Samah Sakr (020 272-4737), or Dr. Nelly Mugo (020 273-6744) for questions about the study or to report any problems.

Signature of study staff _____ Date _____

Printed name of study staff _____

Subject's statement

This study has been explained to me. I volunteer to take part in this research. I give permission to the researchers to use my medical records as described in this consent form. I have had a chance to ask questions. If I have questions later about the research, I can ask one of the researchers listed above. If I have questions about my rights as a research subject, I can call the Ethical Review Committee at Kenyatta National Hospital 726-300. I will receive a copy of this consent form if I would like one.

Please mark, initial and date one option:

_____ I DO agree to store my samples and data for future research

_____ I DO NOT agree to store my samples and data for future research

Signature or thumbprint of participant _____ Date _____

Printed name of participant _____

Copies to: Investigator and Subject

STUDY WRITTEN CONSENT FORM

Cervical Treatment Study: Screening *RE-ENROLLMENT CONSENT*

Full Title:

Impact of Cryotherapy versus Loop Electrosurgical Excision Procedure (LEEP) on Recurrence of Cervical Intraepithelial Neoplasia and HIV-1 Cervical Shedding among HIV-positive Women

Study Investigators:

Michael H. Chung, MD, MPH, Assistant Professor, Department of Global Health, University of Washington, 020 271-2947

Nelly Mugo, MBChB, MMed, MPH, Gynecologist, Department of Obstetrics and Gynecology, Kenyatta National Hospital, 020 273-6744

Samah Sakr, MBChB, Medical Director, Coptic Hospital of Kenya, 020 272-4737

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Barbra Richardson, PhD, Research Professor, Department of Biostatistics, University of Washington, +1-206-731-2425

Grace John-Stewart, MD, PhD, Professor, Department of Medicine, University of Washington, +1-206-543-4278

Emergency telephone number staffed 24 hours a day: 020 271 2947, 0723 914 057 or 0721 289 733

Ethical Review Committee Chairperson: Professor A. N. Guantai, 020 2726300, Ext. 44102, 44355, can be contacted for questions about research subject rights

Researchers' Statement

We are asking you to be in a research study. The purpose of this consent form is to give you information so you can decide if you want to be in the study. Please read the form carefully. You may ask questions about the purpose of the research and what we would ask you to do for the study. You may ask about possible risks and benefits, your rights as a volunteer, and anything else about the research or this form. You may ask questions at any time (before, during, and after the study) about anything. When we have answered all your questions, you can decide if you want to be in the study or not. This process is called 'informed consent.' If you wish, we will give you a copy of this form. Please let us know if you would rather use the Kiswahili consent form.

Purpose of the Study

The reason we are doing this research project is to screen for disease which may lead to cervical cancer. Participation in this project lasts for up to 1 month. You were asked to participate in this project because you are:

- HIV-positive,
- currently receiving care at the Hope clinic,
- not pregnant,
- do not have a history of problems with bleeding,
- have not had a hysterectomy (an operation to remove the uterus),
- and have initiated sexual intercourse.

Cervical cancer is the most common cancer among young and middle-aged women in Kenya. More women who have HIV get cervical cancer than those who do not have HIV. Even in HIV-infected women, cervical cancer is not common. However, in HIV-infected women, cervical disease that is not cancer is common. This disease is caused by a virus called the human papillomavirus (HPV). It is very important to find this disease and treat it before it becomes cancer. This study will screen you for cervical disease. If you screen positive for cervical disease, you may choose to enroll in a treatment study, or you will be referred elsewhere for treatment. The treatment study will be explained to you in detail before you decide. This screening study will have over 2,400 participants. By doing this study, we hope to provide free and comprehensive screening for HIV-positive women to prevent cervical cancer.

Procedures (see Appendix)

All participants will be asked to come for 1 to 3 visits over 1 month. All study visits will take from 20 to 40 minutes (Visit 1: consent process – 10 minutes, Pap smear – 20 minutes; Visit 2: review of Pap smear results – 10 minutes, colposcopy and biopsy – 30 minutes; Visit 3: review of biopsy results and discussion of treatment options – 30 minutes)

Visit 1:

If you agree, you will first undergo a pelvic exam with a Pap smear to see if you have a something like a wound (called a lesion). This lesion may possibly develop into cancer. A pelvic exam means that a doctor or nurse will examine your female parts. The Pap smear involves brushing the cervix with a small brush to collect material that can be looked at under a microscope (like a large magnifying glass).

Visit 2:

You will then return to this clinic 2 weeks later to find out the results of this test. Most of the time the results will show no disease or that we just have to repeat the Pap smear or visual inspection (VIA) every 6 months. If this is the case, then you will exit the study at this point and receive further follow-up at the Coptic Hope Cervical Cancer Screening Program (CCSP) or be asked to come back to the study clinic for a repeat Pap smear after 6 months. If we find lesions that need to be treated at these visits, we will discuss your treatment options with you in detail.

If you are screened and we find a lesion that needs to be looked at carefully, you will first have a pregnancy test. If you are pregnant, nothing more will be done until after you deliver. We will send you for special care at the Hope center to make sure you do not pass HIV to your baby (this care is called PMTCT), and you will exit the study and receive further follow-up at the CCSP. If you are not pregnant, we will look at your cervix with a type of magnifying lens called a colposcope. We will look closely at the cervix and take a piece out, about the size of a grain of rice. This is called a biopsy. We will look at this tissue biopsy under a microscope and share the results with you 4-6 weeks later. This test is very good and will help us decide whether you need to have treatment.

There are some cells found in the cervix and the lining of the uterus. These are called ‘glandular cells.’ When Pap smear results show that your glandular cells are abnormal, the doctor will perform a procedure referred to as ‘Endocervical Curettage (ECC).’ This is a procedure where a spoon-shaped instrument called a ‘curette’ is used to scrape abnormal material from the passageway between the cervix and the uterus. This procedure obtains a small sample, which is then sent to the lab to be examined for abnormal cells. ECC is performed during colposcopy and takes just a few minutes to perform. You can expect to feel mild cramping, much like menstrual cramps following the procedure. If you are found to have abnormal cells but the doctor cannot see them by means of a colposcope, you will undergo ECC as described above.

Visit 3: You will return to this clinic in 4-6 weeks for the biopsy results. If we find cervical disease that needs treatment, we will discuss your treatment options with you. You may choose to enroll in a further treatment study, which will be explained in detail at this time. If you prefer not to enroll in another study, you will be referred for treatment at Kenyatta National Hospital (KNH) and standard care at Coptic Hope Center. Whether you choose to accept study treatment or referral, you will be exited from this screening study at your third visit.

Contacting Participants

We will ask you to give us contact information like a phone number so we can call you if you do not come to a scheduled visit. We may ask about your health or where you are during these calls. It is important for you to come to all of your scheduled visits. If we cannot contact you by phone, we may try to visit you at your house. If this happens, we will not wear clothes that show we are health workers. You should tell us if you do not want to be contacted in this way. If you do not return for your Pap smear or biopsy results, we will also “flag” your Coptic Hope file so that you can be notified that you have available study results at your next clinic visit.

If study quality control procedures indicate that your Pap smear result is discordant from the original result you were given, we will ask you to return to the clinic to receive your new results and to discuss potential treatment options. At this point, you may be asked if you would like to re-enroll in the study.

Risks and discomforts of being in the study

This is an explanation of problems you may have through your participation in this study. Other problems not listed may happen as well.

Screening

Collection of samples through the pelvic exam may cause:

- mild discomfort as you are examined,
- a small amount of bleeding from the vagina for 1-2 days afterwards and
- mild to moderate cramping for around 5 minutes that is similar to mild period pain.

If you receive a biopsy, we ask that you do not have sex for 10 days. The reason is that if you are HIV-infected and your partner is not, then he may be at greater risk of becoming infected with HIV. If your partner is HIV positive, he may be at greater risk of getting re-infected with resistant HIV. Also, you need time to heal and will put yourself at higher risk of infection unless you give yourself this time. If you need help talking about these issues with your partner, a nurse, doctor or counselor can help you. We will give you free condoms to use if it is impossible for you to not have sex.

Another possible discomfort you may face is the worry or anxiety that you may have disease on your cervix. You may talk about this with a study nurse or doctor or if you would like talk to a counselor, we can help to arrange this.

We may find on screening that you have an infection. In this case we will give you a prescription for antibiotics. Side effects of these antibiotics may include: upset stomach, vomiting or diarrhea, sensitivity to light, rash, severe allergic reaction that could cause death, dizziness, headache, confusion, ringing in the ears or hearing loss, seizures (jerking of the body), heart problems, blood disorders, problems with blood sugar, liver disease, kidney disease and pain or numbness in your legs or arms. If you experience any of these problems, please report to study staff right away.

Confidentiality

The study staff will ask you for personal information that may be embarrassing to talk about like how many sexual partners you have had. You may choose not to answer any question. As part of the study, you may meet other patients from this clinic whom you know from outside the clinic. We have no plans to release your information to anyone other than the study researchers or appointed monitors. Sometimes committees that oversee research will examine study information to make sure nothing illegal or unethical is being done. Your personal information will be protected if this happens and will not be shown to anyone outside of this review. As the study sponsor, CDC may monitor or audit study activities. The reason for this would be to make sure that the study is being done the way it is supposed to be done. It would also make sure that your rights and health are protected. Your personal medical information will be kept confidential.

Alternative to taking part in this study

If you choose not to take part in this study, you will continue to receive medical care at the Coptic Hope Center and free antiretroviral medications. You may also receive free cervical cancer screening at this clinic without having to enroll in this study.

Benefits of the study

Your participation will help us understand more about cervical disease. By participating in this study, you receive free screening for cervical disease. If you would like to know the results of the study you can contact the study office 6 months or 1 year after you leave the study.

Compensation for injury

There is no cost to you for participating in this study other than your time. The study will pay for all screening costs for tests provided at the study clinic. If any physical injuries happen to you as a result of study participation, the study will cover the costs of care. If you think you have an injury or illness related to this study, contact the study staff (Dr. Michael Chung (020-272-2710) or Dr. Nelly Mugo (020-273-6744) or Dr. Evans Malava (0721 289 733) right away. They will treat you or refer you for treatment.

Specimen and Data Storage and Use of your Samples for Future Studies

We would like to save your medical information and Pap smear, colposcopy and endocervical curettage samples in Kenya at the Coptic Hospital and Kenyatta National Hospital for future research. This research may be done by the University of Washington or by other researchers who are working with us on this study for ten years after the end of follow-up in the study. We will use these data and samples only for research related to cervical cancer and HIV. Before your samples leave the clinic, they will be assigned a code. Your name will not be on them. Your name will be linked to the code only for five years after the study is completed. After that time, the link between your name and the code on your samples and data will be destroyed. The Institutional Review Boards are committees that watch over the safety and rights of research participants at Kenyatta National Hospital and the University of Washington. They must approve any future research studies using your samples. If you do not want to have your samples saved for future research, you can still be in this study and your samples will be destroyed once testing for the study is completed. If you agree to store your samples now, but change your mind before the end of the study, let the study staff know and we will make sure that your samples do not get stored for future research. We will not sell your samples. Tests done on your samples may lead to a new invention or discovery. We have no plans to share any money or other benefits resulting from this invention or discovery with you.

Other information

Your medical information is confidential (or kept secret) and we will keep your records in a locked office. Your medical records and information about your participation in the research will be available to you and to the study team but not to anyone outside of the study without your agreement. If you agree, we will share the information from this study with your doctors at the Hope Center. This information may help them treat your HIV and give you better care. All of your records will be kept in locked areas and all computer information will be password protected.

University of Washington staff sometimes review studies such as this one to make sure they are being done safely and legally. If a review of this study takes place, your records may be examined. You may refuse to participate or may leave from the study at any time without penalty or loss of benefit or help to which you have a right to. We will tell you if there is any new information about the treatments we are studying so you can decide if you want to leave the study. Your relationship with staff and services at Coptic Hope Center for Infectious Diseases will not be affected in any way if you do or do not participate or if you enter the program and leave later. Please inform study staff if you decide you would like to leave the study. You may be asked to give some final samples but you may refuse.

Study staff may decide to take you out the study if they find you may be harmed if you continue to participate. You may also be taken out of the study if the staff think you will not be able to follow study safety requirements.

Please contact Dr. Michael Chung (020 271-2947), Dr. Samah Sakr (020 272-4737), or Dr. Nelly Mugo (020 273-6744) for questions about the study or to report any problems.

Signature of study staff _____ Date _____

Printed name of study staff _____

Subject's statement

This study has been explained to me. I volunteer to take part in this research. I give permission to the researchers to use my medical records as described in this consent form. I have had a chance to ask questions. If I have questions later about the research, I can ask one of the researchers listed above. If I have questions about my rights as a research subject, I can call the Ethical Review Committee at Kenyatta National Hospital 726-300. I will receive a copy of this consent form if I would like one.

Please mark, initial and date one option:

_____ I DO agree to store my samples and data for future research

_____ I DO NOT agree to store my samples and data for future research

Signature or thumbprint of participant _____ Date _____

Printed name of participant _____

Copies to: Investigator and Subject

STUDY WRITTEN CONSENT FORM

Cervical Treatment Study: Cryotherapy vs. LEEP

Full Title:

Impact of Cryotherapy versus Loop Electrosurgical Excision Procedure (LEEP) on Recurrence of Cervical Intraepithelial Neoplasia and HIV-1 Cervical Shedding among HIV-positive Women

Study Investigators:

Michael H. Chung, MD, MPH, Assistant Professor, Department of Global Health, University of Washington, 020 271-2947

Nelly Mugo, MBChB, MMed, MPH, Gynecologist, Department of Obstetrics and Gynecology, Kenyatta National Hospital, 020 273-6744

Samah Sakr, MBChB, Medical Director, Coptic Hospital of Kenya, 020 272-4737

Hugo De Vuyst, MD, PhD, Epidemiologist, International Agency for Research on Cancer, World Health Organization, +33-472738521

Silvia Franceschi, MD, Epidemiologist, International Agency for Research on Cancer, World Health Organization, +33-4728404

Barbra Richardson, PhD, Research Professor, Department of Biostatistics, University of Washington, +1-206-731-2425

Grace John-Stewart, MD, PhD, Professor, Department of Medicine, University of Washington, +1-206-543-4278

Emergency telephone number staffed 24 hours a day: 020 271 2947, 0723 914 057 or 0721 289 733

Ethical Review Committee Chairperson: Professor A. N. Guantai, 020 2726300, Ext. 44102, 4435544355, can be contacted for questions about research subject rights

Researchers' Statement

We are asking you to be in a research study. The purpose of this consent form is to give you information so you can decide if you want to be in the study. Please read the form carefully. You may ask questions about the purpose of the research and what we would ask you to do for the study. You may ask about possible risks and benefits, your rights as a volunteer, and anything else about the research or this form. You may ask questions at any time (before, during, and after the study) about anything. When we have answered all your questions, you can decide if you want to be in the study or not. This process is called 'informed consent.' If you wish, we will give you a copy of this form. Please let us know if you would rather use the Kiswahili consent form.

Purpose of the Study

The reason we are doing this research project is to find the best treatment of disease which may lead to cervical cancer. Participation in this project lasts for 2 years. You were asked to participate in this project because you are:

- HIV-positive,
- currently receiving care at the Hope clinic,
- not pregnant,
- do not have a history of problems with bleeding,
- have not had a hysterectomy (an operation to remove the uterus),
- have initiated sexual intercourse
- have not received treatment for cervical disease in the past
- and have received a positive result for cervical disease from Pap smear and biopsy screening.

Cervical cancer is the most common cancer among young and middle-aged women in Kenya. More women who have HIV get cervical cancer than those who do not have HIV. Even in HIV-infected women, cervical cancer is not common. However, in HIV-infected women, cervical disease that is not cancer is common. This disease is caused by a virus called the human papillomavirus (HPV). It is very important to find this disease and treat it before it becomes cancer. This study will compare two ways of treating cervical disease: cryotherapy and loop electrosurgical excision procedure (LEEP). Both treatments are commonly done for women around the world and are not new. We will explain these treatments for you. This study will have 400 participants. By doing this study, we hope to find the best way to treat HIV-positive women to prevent cervical cancer.

Procedures (see Appendix)

All participants will be asked to come for 5 to 8 visits over 2 years. All study visits will take from 15 to 40 minutes.

Randomization (Visit 1: review of biopsy results – 10 minutes, randomization and treatment– 30 minutes)

Based on your Pap smear and biopsy results, we have found cervical disease that needs treatment. If you decide to participate in this study, we will offer you one of two common and effective treatment methods. One method is called cryotherapy. It is a procedure which will freeze and remove the diseased part of your cervix. We freeze by touching your cervix with a small stick that is very cold. For cryotherapy, you will be offered an oral painkiller. The other method is called LEEP and uses a heated wire to do the same thing after the cervix is numbed by medication. The heated wire will scoop out the disease from the cervix.

You will be randomly assigned to one of these methods. Random assignment is like “flipping a coin.” You have an equal chance of receiving either of these methods. We won’t know which treatment you will receive until we open an envelope that has a sheet of paper telling us which treatment you will get. Neither of us will choose your treatment. We are randomly assigning one of these methods because we do not know if one treatment is better than another for women who have HIV.

If we see that the diseased part of your cervix or ‘lesion’ is too large and cannot be treated well by cryotherapy, then we will not randomize you and will choose to treat you with LEEP. You will receive LEEP treatment free of charge and we will follow you every 6 months for 2 years. If the lesion cannot be treated well by either cryotherapy or LEEP, then we will refer you to Kenyatta National Hospital (KNH). At KNH you can receive different types of treatments at the lower cost of a government hospital. We will send copies of forms with you that will be important for your care. These forms will not show that you are part of a study. We will provide follow-up for you for 2 years after your care at KNH. If you agree to be treated by either cryotherapy or LEEP at the study clinic, 2 teaspoons (10 mls) of your blood will be taken with a needle from your arm. We will check a CD4 count and the levels of HIV in your blood. We will also brush your cervix and later test the sample for levels of HIV and for HPV. We would like to look at your medical records at the Hope Center. We want to gather a full picture of your prescribed medications and medical condition, the results of your laboratory tests, and your attendance in clinic.

Follow-up after Randomization (Visits 2-8: review results and cervical swab – 30 minutes)

If you are randomized to cryotherapy or LEEP, we will ask you to return again at the 1, 2, and 3-week visits after treatment. At these visits, we will draw 2 teaspoons (10mls) of blood and again brush your cervix to see how much HIV is there. We will want to see whether the level of virus is increased in your cervix after treatment and when it returns to usual levels.

If you are randomized or offered LEEP but not randomized, we will also ask you to return at 6, 12, 18, and 24 months after treatment for repeat Pap smears. This is to make sure that the diseased part of your cervix was completely removed and/or no new abnormal tissue has formed. We will also brush your cervix to test for HPV at this time and take 2 teaspoons (10 mls) of blood to measure your CD4 count and HIV viral levels. If we find more abnormal lesions at this time, we will treat you with LEEP or refer you for further care at KNH. The study will not pay for care you receive at KNH.

Contacting Participants

We will ask you to give us contact information like a phone number so we can call you if you do not come to a scheduled visit. We may ask about your health or where you are during these calls. It is important for you to come to all of your scheduled visits. We want to follow you carefully to find and treat any cervical disease you might have or develop later on. If we cannot contact you by phone, we may try to visit you at your house. If this happens, we will not wear clothes that show we are health workers. You should tell us if you do not want to be contacted in this way.

Risks and discomforts of being in the study

This is an explanation of problems you may have through your participation in this study. Other problems not listed may happen as well.

Treatment

There may be risks or discomforts from receiving treatment. If you have cryotherapy, then you may have:

- mild abdominal (or belly) cramps (usually last less than 10 minutes),
- fluid from the vagina for about 2 weeks (may last longer),

- bleeding,
- infection that we will need to treat here at the clinic or in rare cases, at the hospital. Infection may cause fevers, chills, night sweats, or white fluid from the vagina.

If you have LEEP, you may develop:

- bleeding during or after the procedure (you may have to return to the clinic if the bleeding starts and continues after the procedure is done)
- infection.

All of these complications can be treated with medications or treatments that will be provided free of charge by the clinic. Possible treatments may include:

- antibiotic medications to treat infection,
- packing the vagina with bandages to stop bleeding,
- putting stitches in the cervix to stop bleeding or
- hospitalization for severe infection or bleeding.

Please tell us if you have any of these problems after treatment. If you have received cryotherapy or LEEP, then we ask that you do not have sex for 4 weeks after it is performed. The reason is the same that your partner may be at greater risk of becoming infected with HIV. Also, you may be at greater risk of getting an infection. We can help you to discuss this with your partner.

All participants who have treatment (cryotherapy or LEEP) will get a prescription for antibiotics. These antibiotics include doxycycline, metronidazole or norfloxacin.

Effects of these antibiotics may include upset stomach (must avoid alcohol for 2 days), vomiting or diarrhea, sensitivity to light, rash, dizziness, headache, confusion, seizures (jerking of the body), heart problems, blood disorders, problems with blood sugar, problems, liver disease and pain or numbness in your legs or arms. Any reaction to medications should be reported right away to study staff.

After having the treatment, there is a small chance that you may have problems later with pregnancy. After having LEEP, some women have problems when they are pregnant including infections, early contractions or problems with the cervix. Some women develop a tightened opening of their cervix that must be stretched. Other women have had their water break early or had babies born early possibly because of the operation on their cervix.

Blood draw

We will be collecting blood samples, 2 teaspoons (10 mls), from you using a new needle and syringe. The puncture of the needle may be uncomfortable and leave a bruise. It may also cause infection or fainting.

Confidentiality

The study staff will ask you for personal information that may be embarrassing to talk about like how many sexual partners you have had. You may choose not to answer any question. As part of the study, you

may meet other patients from this clinic whom you know from outside the clinic. We have no plans to release your information to anyone other than the study researchers or appointed monitors. Sometimes committees that oversee research will examine study information to make sure nothing illegal or unethical is being done. Your personal information will be protected if this happens and will not be shown to anyone outside of this review. As the study sponsor, CDC may monitor or audit study activities. The reason for this would be to make sure that the study is being done the way it is supposed to be done. It would also make sure that your rights and health are protected. Your personal medical information will be kept confidential.

Alternative to taking part in this study

If you choose not to take part in this study, you will continue to receive medical care at the Coptic Hope Center and free antiretroviral medications. You may also receive free cervical cancer screening at this clinic without having to enroll in this study. The Coptic Hope Clinic can provide you with Cryotherapy treatment even if you do not enroll in this study. We can also refer you to Kenyatta National Hospital for different types of treatments at the lower cost of a government hospital.

Benefits of the study

Your participation will help us understand more about cervical disease. This may change the way cervical disease is found and treated in developing countries like Kenya. If you would like to know the results of the study you can contact the study office 6 months or 1 year after you leave the study.

Compensation for injury

There is no cost to you for participating in this study other than your time. The study will pay for all screening and treatment costs for tests and therapy provided at the study clinic. If any physical injuries happen to you as a result of study participation, the study will cover the costs of care. Treatment includes antibiotics, pain relief, and methods to stop bleeding. If you think you have an injury or illness related to this study, contact the study staff (Dr. Michael Chung (020-272-2710) or Dr. Nelly Mugo (020-273-6744) or Dr. Evans Malava (0721 289 733) right away. They will treat you or refer you for treatment.

Specimen and Data Storage and Use of your Samples for Future Studies

We would like to save your medical information and samples of your blood and cervix in Kenya at the Coptic Hospital and Kenyatta National Hospital for future research. This research may be done by the University of Washington or by other researchers who are working with us on this study for ten years after the end of follow-up in the study. We will use these data and samples only for research related to cervical cancer and HIV. Before your samples leave the clinic, they will be assigned a code. Your name will not be on them. Your name will be linked to the code only for five years after the study is completed. After that time, the link between your name and the code on your samples and data will be destroyed. The Institutional Review Boards are committees that watch over the safety and rights of research participants at Kenyatta National Hospital and the University of Washington. They must approve any future research studies using your samples. If you do not want to have your samples saved for future research, you can still be in this study and

your samples will be destroyed once testing for the study is completed. If you agree to store your samples now, but change your mind before the end of the study, let the study staff know and we will make sure that your samples do not get stored for future research. We will not sell your samples. Tests done on your samples may lead to a new invention or discovery. We have no plans to share any money or other benefits resulting from this invention or discovery with you.

Other information

Your medical information is confidential (or kept secret) and we will keep your records in a locked office. Your medical records and information about your participation in the research will be available to you and to the study team but not to anyone outside of the study without your agreement. If you agree, we will share the information from this study with your doctors at the Hope Center. This information may help them treat your HIV and give you better care. Some of your samples including blood and samples from the cervix may be sent to the USA or Europe for testing. Your information and samples will still be protected if this happens. All of your records will be kept in locked areas and all computer information will be password protected.

University of Washington staff sometimes reviews studies such as this one to make sure they are being done safely and legally. If a review of this study takes place, your records may be examined.

You may refuse to participate or may leave from the study at any time without penalty or loss of benefit or help to which you have a right to. We will tell you if there is any new information about the treatments we are studying so you can decide if you want to leave the study. Your relationship with staff and services at Coptic Hope Center for Infectious Diseases will not be affected in any way if you do or do not participate or if you enter the program and leave later. Please inform study staff if you decide you would like to leave the study. You may be asked to give some final samples but you may refuse.

Study staff may decide to take you out the study if they find you may be harmed if you continue to participate. You may also be taken out of the study if the staff think you will not be able to follow study safety requirements.

Transportation costs of Ksh 300 will be given to you when you return to the clinic for a study-related visit. You will receive transport money from the study receptionist.

Please contact Dr. Michael Chung (020 271-2947), Dr. Samah Sakr (020 272-4737), or Dr. Nelly Mugo (020 273-6744) for questions about the study or to report any problems.

Signature of study staff _____ Date _____

Printed name of study staff _____

Subject's statement

This study has been explained to me. I volunteer to take part in this research. I give permission to the researchers to use my medical records as described in this consent form. I have had a chance to ask questions. If I have questions later about the research, I can ask one of the researchers listed above. If I have

questions about my rights as a research subject, I can call the Ethical Review Committee at Kenyatta National Hospital 726-300. I will receive a copy of this consent form if I would like one.

Please mark, initial and date one option:

_____ I DO agree to store my samples and data for future research

_____ I DO NOT agree to store my samples and data for future research

Signature or thumbprint of participant _____ Date _____

Printed name of participant _____

Copies to: Investigator and Subject

MAKUBALIANO YA PAMOJA**KISWAHILI CONSENT FORM****MATIBABU YA SARATANI YA MLANGO WA KIZAZI: UCHUNGUZI****IDHINI YA AWALI****WATAFITI**

Michael H Chung, MD. MPH, Mkufunzi, Idara ya utabibu, Chuo Kikuu cha Washington 272-2710

Nelly Mugo, MBChB, MMed, MPH, Gainakolojia, Mhadhiri, Idara ya Ukunga na Gainakologia, Hospitali kuu ya Kenyatta, 020-273-6744

Sarah Sakr, MBChB, Msimamizi wa afya, Hospitali ya Coptic, Kenya 020-272-4737

Hugo De Vuyst, MD, PhD, Epidemiologisti, Idara ya Kimataifa ya Utafiti wa Saratani, Shirika la Afya la Dunia (WHO), +33-472738521

Silvia Franceschi, MD, Epidemiologisti, Idara ya Kimataifa ya Utafiti wa Saratani, Shirika la Afya la Dunia (WHO), +33-4728404

Barbara Richardson, PhD, Profesa Msaidizi katika Utafiti, Idara ya Biostatistiki, Chuo Kikuu cha Washington, +1-206-731-2425

Grace John Stewart, MD, PhD, Profesa, Kitivo Cha Utabibu, Chuo Kikuu Cha Washington, +1-206-543-4278

Nambari za simu ya dharura; 072-2710 au 0733-711-288

Mwenyekiti wa kamati ya uchunguzi wa maadili; Proffesa A. N. Guantai, 020 2726300, Ext. 44102, 44355 unaweza kuwasiliana naye kwa maswala ya utafiti na maadili ya washiriki wa utafiti huu.

Ujumbe wa Watafiti

Tunakuuliza kushiriki katika utafiti huu wa kitaalamu. Lengo la fomu hii ya idhini ni kukupa habari itakayokufahamisha na kukusaidia kuamua kama ungelipenda kushiriki katika utafiti huu au la. Tafathali soma maelezo haya kwa makini. Unaweza kuuliza maswali kuhusu nia/kusudi ya utafiti huu; unavyohitajika kufanya katika utafiti, uwezekano wa kuwepo na madhara au manufaa yeyote, haki yako kama aliyejitolea, na mambo mengine kuhusu utafiti huu au chochote usicho elewa kwenye fomu hii. Tukishajibu maswali yako yote, unapaswa kuamua kushiriki kwenye utafiti huu au la. Kukubali kushiriki kwenye utafiti huu kwa hiari yako kunamaanisha kuwa umeelezwa na umeelewa yote yanayohusika na kukubaliana nayo. Ukipenda, tutakupua nakala ya fomu hii kujihifadhia. Tafadhali tujulishe iwapo ungependa kutumia lugha ya Kingereza.

Lengo la Utafiti huu.

Mathumuni ya mradi huu wa utafiti ni kukupima njia yako ya uzazi kuthibitisha ikiwa una dalili inayoashiria kuwa unaweza kupata saratani ya mlango wa uzazi (cervical cancer). Umeulizwa kushiriki katika utafiti huu kwa sababu;

- Umeambukizwa maradhi ya UKIMWI
- Unapata matibabu katika kituo cha matibabu cha Hope
- Hauna Mimba
- Hauna historia ya shida ya kuvuja damu
- Haujapata kutolewa sehemu yako ya uzazi (uterasi)
- Umewahi fanya ngono

Saratani ya aina hii huambukiza wanawake wengi wenye umri mdogo na hata wa makamu nchini Kenya. Wanawake walioambukizwa na virusi vinavyosababisha ukimwi (HIV) wana hatari kubwa ya kuambukizwa saratani ya mlango wa kizazi (cervical cancer), kuliko wanawake wengine ambao bado hawajaambukizwa na HIV. Hata hivyo, kwa wanawake walioambukizwa na virusi vinavyosababisha ukimwi, saratani ya mlango wa kizazi si ya kawaida. Lakini katika wanawake walioambukizwa na virusi vinavyosababisha ukimwi, maambukizo ya mlango wa kizazi usio saratani ya mlango wa kizazi ni wa kawaida. Maambukizo haya husababishwa na virusi vya “Papiloma” (HPV). Ni muhimu kutambua na kutibu maambukizo haya kabla hayajageuka kuwa saratani ya mlango wa kizazi.

Utafiti huu utakupima mlango wako wa uzazi. Iwapo wachunguzi wakipata chembe chembe zisizo za kawaida ambazo zinahitaji kutibiwa katika kiwango hiki, unaweza kujiunga na utafiti wa matibabu ya ugonjwa unaosababisha saratani ya mlango wa uzazi, ama unaweza kutumwa upate matibabu kwingine. Tutakuelezea njia za matibabu kabla hujaamua kujiunga na utafiti huu. Katika utafiti huu, tutakuwa na washiriki takriban 2,400. Utafiti huu utatuwezesha kupima wanawake bila malipo ili kuwawezesha wale wanoishi na virusi vinavyosababisha ukimwi (HIV) kuishi maisha bila saratani ya mlango wa kizazi.

Hatua ya kushiriki katika utafiti

Washirika wote watahitajika kutembelea kiliniki mara 1 au 2 au 3. Kila mara, mshirika atatumia dakika 20 hadi 40 katika kiliniki.

Uteuzi; (Kutembelea kiliniki Mara Ya 1; Makubaliano - dakika 10, Pap Smear-dakika 20; Kutembelea kiliniki Mara Ya 2; Marejeleo ya Matokeo ya Pap Smear- dakika 10, Colposcopy na Biopsy (ikihitajika)- dakika 30; Kutembelea kiliniki mara ya 3; Marejeleo ya matokeo ya biopsy na kujadili juu ya njia za matibabu- dakika 30)

Kutembelea kiliniki mara ya 1:

Ukikubali kushiriki katika utafiti huu, kwanza utahitajika kuchunguzwa fupanyonga (Pelvic exam) kwa kupanguzwa sehemu yako ya siri kutumia burashi ili kuthibitisha ikiwa una dalili

inayoashiria kuwa na virusi vinavyoweza kusababisha saratani ya mlango wa uzazi (Pap Smear). Kuchunguzwa fupanyonga, kunamaanisha kuwa muuguzi atakagua sehemu zako za siri (uke wako). Pap smear itajumuisha utumiaji wa burashi ndogo iingizwayo kwenye kizazi ili kuchukua chembe chembe zitakazochunguzwa kwa kutumia “mikroskope”. Kisha utahitajika kurudi katika kiliniki baada ya majuma mawili ili kuchukua matokeo ya uchunguzi huu.

Kutembelea kiliniki mara ya 2:

Utarejea kwenye kiliniki baada ya wiki mbili ili kupata matokeo ya pap smear. Mara nyingi, matokeo huwa ni sawa (hakuna kasoro) na mtu hahitaji matibabu ya ziada, ila kurudia “Pap smear” tena baada ya kila miezi sita au kufanyaukaguzi na asidi asetiki (VIA) baada ya miezi sita. Ikiwa hujambo utaondoka utafitini katika kiwango hiki na kupokea matibabu mengine katika Kiliniki ya ukaguzi wa Ugonjwa wa saratani ya mlango wa kizazi ya Coptic Hope Centre, [*Coptic Hope Cervical Cancer Screening Program (CCSP)*] ampapo utapokea VIA, au utaulizwa kurudi katika kiliniki ya utafiti baada ya miezi sita ili kurudia “Pap smear”.

Wachunguzi wakipata chembe chembe zisizo za kawaida ambazo zinahitaji kutibiwa katika kiwango hiki, watakuelezea juu ya matibabu.

Lakini watafiti wakigundua chembe chembe zisizo za kawaida, watachunguza sehemu yako ya siri kwa makini. Kwanza utapimwa kama wewe ni mjamzito. Kama wewe ni mjamzito, Hakuna utafiti mwingine utaendelea hadi utakapojifungua mtoto. Watafiti watakutuma kupata matibabu ya dharura katika kituo cha Hope ili kuhakikisha hauambukizi mtoto wako virusi vinavyosababisha ukimwi (PMTCT). Kisha utaondoka utafitini katika kiwango hiki na kupokea matibabu mengine katika Kiliniki ya ukaguzi wa Ugonjwa wa saratani ya mlango wa kizazi ya Coptic Hope Centre, [*Coptic Hope Cervical Cancer Screening Program (CCSP)*].

Ikiwa wewe si mjamzito, watafiti wakigundua chembe chembe zisizo za kawaida, watachunguza sehemu yako ya siri kwa makini wakitumia chombo kiitwacho “colposcope”, ili kuchukua tishu, kiwango kama kipande kimoja cha mchele, utarabu huu unajulikana kama “Biopsy”. Tishu iliyochukuliwa, itachunguzwa kwa mikroskope na utajulishwa matokeo baada ya wiki nne hadi sita. Mwenendo huu wa kupima ambao majibu yake huwa ni sahihi utatuwezesha kujua kana kwamba unahitaji matibabu.

Kuna chembe chembe za ndani ya njia ya kizazi katikati ya mlango wa kizazi na kizazi na pia zinaendelea mpaka ndani ya kizazi. Hizi chembe chembe zinauitwa ‘glandular cells’. Ikiwa majibu ya pap smear itaonyesha ‘glandular’ cells si kawaida, daktari ata gwaruza kwa njia ya kizazi kutumia chombo chenye umbo cha kijiko ili sampuli ndogo ipatikane ya kukaguliwa. Utaratibu huu unaitwa ‘Endocervical curettage (ECC)’. Kukwaruza inafanywa wakati ‘colposcopy’ na inachukuwa dakika chache. Unaweza pata maumivu ya tumbo kidogo kama wakati unapo pata damu ya mwezi. Pia, kama chembe chembe za mlango wa kizazi kwa pap smear inaonyesha si kawaida na mabadiliko ya ugonjwa kwa mlango wa kizazi wakati wa colposcopy haionekani, hii kukwaruza itafanywa kama vile imeelezwa hapo mbeleni.

Kutembelea kiliniki mara ya 3:

Utarejea kwenye kiliniki kati ya wiki nne na wiki sitaili kupata matokeo ya biopsy. Iwapo tutagundua maambukizi yanayohitaji matibabu, tutakujulisha matibabu ambayo unaweza pata. Unaweza kujiunga na utafiti wa matibabu ya ugonjwa unaosababisha saratani ya mlango wa uzazi, ama unaweza chagua matibabu kwingine, au tunaweza kukutuma Hospitali Kuu ya Kenyatta (KNH) na uendelee kupokea matibabu yako ya kawaida katika Coptic Hope Center. Ukichagua kuendelea na utafiti wa matibabu au la, katika kiwango hii, tutakuondoa katika utafiti wa kupimwa mlango wa kizazi.

Tutatumia rekodi zako za afya ziliko HOPE Center ili kukusanya habari zote kuhusu afya yako ili kuelewa vyema dawa ambazo umekuwa ukitumia, hali ya afya yako kwa sasa na majibu ya vipimo kutoka maabara na pia jinsi umekuwa ukihudhuria kliniki.

Kuwasiliana na Washiriki

Tutakuuliza utupatie jinsi ya kuwasiliana nawe kama vile nambari yako ya simu ili tuweze kukupigia usipotembelea kiliniki kama ilivyopangwa. Tunaweza kukuuliza kuhusu afya yako au ulipokuwa katika mawasiliano haya. Ni muhimu kwako kutembelea kiliniki kama ilivyopangwa. Iwapo hatuwezi kuwasiliana nawe kwa njia ya simu, tunaweza kukutembelea kwako nyumbani. Iwapo tutakutembelea kwako nyumbani, hatutavalia mavazi yatakayoashiria kuwa sisi ni wahudumu katika hospitali. Unaweza kutuarifu iwapo hupendi njia hii ya mawasiliano. Usiporejea kupokea matokeo yako ya pap smear au biospy, tutawasiliana na wafanyi kazi wa mapokezi ya Coptic waweke mawaidha kwa faili yako ili unaporejea kwa matibabu yako ya kawaida, utakumbushwa kupitia kwenye kiliniki ya utafiti kupokea matokeo yako.

Kamataratibu zakudhibiti ubora wautafitizinaonyesha kuwa Papsmearyakomatokeonitofautinamotokeoya awali uliyo pewa, tutakuuliza urudiklinikikupokeamatokeo yakompya nakujadilimatibabu inayoweze kama. Katika hatua hii, unaweza kuulizwakamaungependakujandikisha tena katika utafiti.

Athari na usumbufu wa kushiriki kwenye utafiti huu.

Haya ni maelezo ya athari na usumbufu unayoweza kukumbana nayo kwa kushiriki katika utafiti huu. Athari na usumbufu mwingine ambao haujatajwa unaweza kutokea pia.

Kiingilio (Screening)

Kuchukua sampuli kutoka kwenye fupanyonga (pelvic) kwaweza sababisha;

- kukerwa kidogo wakati ukaguzi ukiendelea,
- kutokwa na damu kidogo kwenye uke wako baadaye kwa siku moja au mbili, na
- kupata uchungu mdogo wa tumbo kama ule wa damu ya mwezi kwa dakika tano hivi.

Ukifanyiwa uchunguzi wa “Biopsy” kwenye fupanyonga (Pelvic), tunakusihia usishiriki ngono kwa muda wa siku kumi. Hii ni kwa sababu ikiwa una virusi vinavyosababisha ukimwi na (HIV) ili hali mpenzi hana, atakuwa katika hatari kubwa zaidi ya kuambukizwa na virusi hivi vya ukimwi. Ikiwa mpenzi wako ana virusi vinavyosababisha ukimwi, atakuwa katika hatari kubwa ya kuambukizwa tena na virusi stahimilivu vinavyosababisha ukimwi. Pia, wewe unahitaji muda wa kupona na utajiweka katika hatari ya maambukizi usipojipatia muda huu. Iwapo unahitaji msaada wowote wa kuongelea mambo haya na mpenzi wako, muuguzi, daktari au mshaurikatika hospital atakusaidia. Tutakupatia mipira ya kondomu bila malipo iwapo hutaweza kutoshiriki katika ngono.

Athari nyengine unayoweza kukumbana nayo ni wasiwasiya kuwa una ugonjwa katika mlango wako wa uzazi. Unaweza kuongea kuhusu jambo hili na muuguzi wa kitafiti, daktari, au ukitaka kuongea na mshaurikatika hospitali, tutakusaidia kupanga haya.

Tunaweza kugundua katika kiingilio kuwa una maambukizo. Iwapo utapatikana na maambukizo, tutakupa maagizo ya kupata dawa. Madhara ya dawa hizi yanaweza kuwa kuumwa na tumbo, kutapika na kuharisha, usikivu wa mwangaza, mwasho, mzio mkali mmenyuko unaoweza kusababisha kifo, kizunguzungu, kuumwa na kichwa, kuchanganyikiwa, kupigapiga kwenyemasikio au kupoteza usikivu, mshtuko wa mwili (jerking of the body), maumivu ya moyo, machafuko ya damu, shida ya kisukari, ugonjwa wa maini, ugonjwa wa figo na uchungu au kuganda kwenye mikono na miguu. Ikiwa utahisi mojawepo ya athari hizi, mfahamishe mkaguzi katika utafiti huu mara moja.

Uwekaji wa siri

Wahudumu katika utafiti huu, watakusanya habari ya binafsi na waweza kuona aibu kuongea kuhusu habari zako za ndani kama vile; idadi ya wapenzi ambao ushakuwa nao. Unaweza kuchagua kutojibu maswali yoyote. Pia, ukija kiliniki, waweza kukutana na wagonjwa wengine ambao umewajua mbeleni. Hatuna mpango wa kupeana habari yako ya kibinafsi kwamtu yeyote nje ya utafiti huusipokuwa watafiti au wachunguzi walioteuliwa. Wakati mwingine kamati inayosimamia utafiti huu watachunguza habari ya utafiti huu ili kuhakikisha hakuna jambo lolote ambalo ni kinyume na sheria au kinyume na maadili litakalotendwa. Habari yako ya kibinafsi italindwa ipasavyo ikiwa hayo yatatendeka na hakuna mtu yeyote atakayeona habari hiyo nje ya uchunguzi huu. Kamamdamini wautafiti, CDC wanaweza kufuatilia aukukaguashughuli zautafiti. Sababu hii itakuwahakikishakuwa utafiti unafanyika kwanji inayotakiwa, napia kuhakikishakwamba hakiyakona afya yako inalindwa. Matibabuyakotaarifa binafsi itakuwasiri.

Mbadala wa kujiunga na utafiti huu

Ukichagua kutoshiriki kwenye utafiti huu, utaendelea kuhudumiwa kikamilifu na kupata matibabu na dawa za ART kutoka “Coptic Hope Centre” bila malipo. Unaweza pia kupata uchunguzi wa saratani ya mlango wa nyumba ya uzazi bila malipo yeyote katika kiliniki hii bila kushiriki katika utafiti huu.

Manufaa kutokana na utafiti huu

Kushiriki kwako katika utafiti huu, kutatuwezesha kuelewa zaidi kuhusu ugonjwa wa mlango wa nyumba ya uzazi. Kwa kujiunga na utafiti huu, utapata kupimwa mlango wako wa uzazi bila malipo. Kama ungependa kujua matokeo ya utafiti huu, unaweza kuwasiliana na ofisi ya utafiti huu miezi sita au mwaka mmoja baada ya kumaliza utafiti huu.

Gharama na Fidia ya Majeraha

Hakuna gharama yoyote ya kushiriki katika utafiti huu, ila muda wako pekee. Utafiti huu utalipia gharama ya uchanguzi wowote utakayopokea kwenye kiliniki ya utafiti. Ikiwa Utapata majeraha kutokana na kushiriki katika utafiti huu, utafiti huu utafidia gharama yote ya malipo ya utunzaji. Ikiwa unafikiri kuwa umepata jeraha ama maumivu kutokana na utafiti huu, wasiliana na wahudumu wa utafiti huu, (Daktari Michael Chung (020-272-2710) au Daktari Nelly Mugo (020-273-6744) au Daktari Evans Malava (0721 289 733)) mara moja. Watakuhudumia kwa matibabu au kukupendekeza kwa matibabu.

Kuhifadhi kwa Sampuli na Data na Matumizi ya sampuli hizi kwa utafiti wa baadaye

Tungependa kuhifadhi habari yako ya matibabu na sampuli za pap smear na biopsy katika Hospitali ya Coptic na katika Hospitali kuu ya Kenyatta kwa madhumuni ya utafiti wa baadaye. Utafiti huu waweza kufanywa na Chuo kikuu cha Washington au na watafiti wengine wanaofanya kazi nasi katika utafiti huu kwa muda wa miaka kumi baada ya mwisho wa kufuatiliwa katika utafiti. Tutatumia data na sampuli hizi kwa minajili ya utafiti unaoegemea na saratani ya mlango wa kizazi na UKIMWI. Kabla sampuli yako kutoka kwenye kiliniki, itapewa kifichomaalumu Jina lako halitakuwemo. Jina lako litahusishwa na kificho hiki kwa miaka mitano pekee yake baada ya kukamilisha utafiti huu. Baada ya muda huu, kihusishi baina ya jina lako na kificho kilichoko katika sampuli na data kitaangamizwa. Bodi za taasisi ya mapitio ni kamati zinazoagalia usalama na haki za washiriki wa utafiti katika Hospitali kuu ya Kenyatta na Chuo kikuu cha Washington. Lazima ziidhinishe utafiti wowote wa baadaye utakaotumia sampuli na data yako. Ikiwa hupendi sampuli zako zihifadhiwe kwa minajili ya utafiti wa baadaye, unaweza kushiriki katika utafiti huu na sampuli zako zitaharibiwa mara tu uchunguzi utakapokamilika. Ukikubali uhufadhi wa sampuli zako leo, kisha ubadilishe nia yako baadaye kabla ya utafiti kukamilika, wafahamishe watafiti ambao watahakikisha kuwa sampuli zako hazitahifadhiwa kwa minajili ya utafiti wa baadaye. Hatutauza sampuli zako. Uchunguzi utakao fanya kwenye sampuli zako waweza kutuelekeza katika uvumbuzi mpya. Hatuna mpango wowote wa kugawana nawe pesa zozote au faida nyinginezo zitakazotokana na uvumbuzi huu -

Maelezo ya ziada

Habari kuhusu utambuzi wa ugonjwa wako ni siri na tutaweka rekodi ya afya yako kwa afisi inayofungwa. Habari kuhusu uchunguzi na ushiriki wako kwenye utafiti huu utatolewa kwako na kikundi cha watafiti pekee wala si mtu mwengine yeyote nje ya utafiti huu bila idhini yako. Ikiwa utapeana idhini, tutawafahamisha habari yako daktari wa Coptic Hope Center. Habari hii inaweza kuwasaidia kukupa matibabu na huduma bora ya UKIMWI. Rekodi zako zote zitahifadhiwa katika sehemu zilizofungwa na kifuli na tarakilishi zote zitakuwa zina neno la siri la kuhakikisha ulinzi.

Wakati mwengine, wafanyikazi wa chuo kikuu cha Washington wanakagua utafiti kama huu ili kuhakikisha unafanywa kwa njia inayofaa na iliyo halali. Ikiwa ukaguzi wa utafiti huu utafanyika, rekodi zako zaweza kuchunguzwa. Waweza kukataa kushiriki au waweza kuondoka kwenye utafiti huu wakati wowote bila kuadhibiwa au kupoteza faida iliyokuwa haki yako. Tutakueleza ikiwa kuna habari geni zinazohusu matibabu ambayo tunatafiti ili uweze kuamua iwapo utajiondoa kwenye utafiti huu. Uhusiano wako na wafanyikazi wa huduma za Coptic Hope Centre for infectious diseases, hautaadhiriwa kwa vyovyote ikiwa utakubali kushiriki kwenye utafiti huu ama ukiingia kwenye mpangilio halafu baadaye uondoke kabla ya kumaliza uchunguzi. Tafadhali wafahamishe wahudumu

wa utafiti huu ikiwa utaamua kuondoka katika utafiti huu. Unaweza kuulizwa kupeana sampuli za mwisho lakini waweza kukataa.

Wahudumu katika utafiti huu wanaweza kuamua kukuondoa kwenye utafiti huu wakigundua kuwa unaweza kupata madhara ukiendelea kushiriki. Unaweza kuondolewa kwenye utafiti huu ikiwa wahudumu watagundua kuwa huwezi kufuata masharti ya usalama.

Maswali yeyote kuhusu utafiti huu, ama athari mbaya kutokana na uchunguzi huu yapaswa kuelekezwa kwa mtafiti anayekuhudumia au Daktari Michael Chung (272-2710), Daktari Samah Sakr (272-4737), au Daktari Nelly Mugo (273-6744).

Je, una maswali yeyote?

Sahihi ya mtafiti..... Tarehe.....

Jina la Mtafiti.....

Muhusika

Nimeelezwa juu ya utafiti huu. Najitolea kwa hiari kushiriki kwenye utafiti huu. Nimewapa watafiti ruhusa ya kutumia rekodi zangu za utabibu kama ilivyopendekezwa kwenye fomu hii ya makubaliano. Nimekuwa na fursa ya kuuliza maswali. Nikiwa na maswali yeyote kuhusu utafiti huu baadaye, naweza kumuuliza mojawapo wa watafiti waliotajwa kwenye fomu hii. Nikiwa na maswali kuhusu haki yangu kama mhusika kwenye utafiti, naweza piga simu kwa kamati ya uchunguzi wa maadili walioko Hospitali kuu ya Kenyatta kwa kutumia nambari 020-726-300. Nikitaka, nitapewa nakala ya fomu hii yangu binafsi.

Tafadhali tia alama, ufupi wa jina na anwani katika chaguo moja:

____ Nimekubali kuweka sampuli na data kwa minajili ya utafiti wa baadaye

____ Sijakubali kuweka sampuli na data kwa minajili ya utafiti wa baadaye

Sahihi au alama ya kidole cha gumba cha mshiriki _____

Tarehe _____

Jina la Mshiriki lililochapishwa _____

Nakala kwa Mtafiti na Mshiriki

MAKUBALIANO YA PAMOJA**KISWAHILI CONSENT FORM****MATIBABU YA SARATANI YA MLANGO WA KIZAZI: UCHUNGUZI****UANDIKISHAJI UPYA****WATAFITI** _____

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Nambari za simu ya dharura; 072-2710 au 0733-711-288

Mwenyekiti wa kamati ya uchunguzi wa maadili; Proffesa A. N. Guantai, 020 2726300, Ext. 44102, 44355 unaweza kuwasiliana naye kwa maswala ya utafiti na maadili ya washiriki wa utafiti huu.

Ujumbe wa Watafiti

Tunakuuliza kushiriki katika utafiti huu wa kitaalamu. Lengo la fomu hii ya idhini ni kukupa habari itakayokufahamisha na kukusaidia kuamua kama ungelipenda kushiriki katika utafiti huu au la. Tafathali soma maelezo haya kwa makini. Unaweza kuuliza maswali kuhusu nia/kusudi ya utafiti huu; unavyohitajika kufanya katika utafiti, uwezekano wa kuwepo na madhara au manufaa yeyote, haki yako kama aliyejitolea, na mambo mengine kuhusu utafiti huu au chochote usicho elewa kwenye fomu hii. Tukishajibu maswali yako yote, unapaswa kuamua kushiriki kwenye utafiti huu au la. Kukubali kushiriki kwenye utafiti huu kwa hiari yako kunamaanisha kuwa umeelezwa na umeelewa yote yanayohusika na kukubaliana nayo. Ukipenda, tutakupua nakala ya fomu hii kujihifadhi. Tafadhali tujulishe iwapo ungependa kutumia lugha ya Kingereza.

Lengo la Utafiti huu.

Mathumuni ya mradi huu wa utafiti ni kukupima njia yako ya uzazi kuthibitisha ikiwa una dalili inayoashiria kuwa unaweza kupata saratani ya mlango wa uzazi (cervical cancer). Umeulizwa kushiriki katika utafiti huu kwa sababu;

- Umeambukizwa maradhi ya UKIMWI
- Unapata matibabu katika kituo cha matibabu cha Hope
- Hauna Mimba
- Hauna historia ya shida ya kuvuja damu
- Haujapata kutolewa sehemu yako ya uzazi (uterasi)
- Umewahi fanya ngono

Saratani ya aina hii huambukiza wanawake wengi wenye umri mdogo na hata wa makamu nchini Kenya. Wanawake walioambukizwa na virusi vinavyosababisha ukimwi (HIV) wana hatari kubwa ya kuambukizwa saratani ya mlango wa kizazi (cervical cancer), kuliko wanawake wengine ambao bado hawajaambukizwa na HIV. Hata hivyo, kwa wanawake walioambukizwa na virusi vinavyosababisha ukimwi, saratani ya mlango wa kizazi si ya kawaida. Lakini katika wanawake walioambukizwa na virusi vinavyosababisha ukimwi, maambukizo ya mlango wa kizazi usio saratani ya mlango wa kizazi ni wa kawaida. Maambukizo haya husababishwa na virusi vya “Papiloma” (HPV). Ni muhimu kutambua na kutibu maambukizo haya kabla hayajageuka kuwa saratani ya mlango wa kizazi.

Utafiti huu utakupima mlango wako wa uzazi. Iwapo wachunguzi wakipata chembe chembe zisizo za kawaida ambazo zinahitaji kutibiwa katika kiwango hiki, unaweza kujiunga na utafiti wa matibabu ya ugonjwa unaosababisha saratani ya mlango wa uzazi, ama unaweza kutumwa upate matibabu kwingine. Tutakuelezea njia za matibabu kabla hujaamua kujiunga na utafiti huu. Katika utafiti huu, tutakuwa na washiriki takriban 2,400. Utafiti huu utatuwezesha kupima wanawake bila malipo ili kuwawezesha wale wanoishi na virusi vinavyosababisha ukimwi (HIV) kuishi maisha bila saratani ya mlango wa kizazi.

Hatua ya kushiriki katika utafiti

Washirika wote watahitajika kutembelea kiliniki mara 1 au 2 au 3. Kila mara, mshirika atatumia dakika 20 hadi 40 katika kiliniki.

Uteuzi; (Kutembelea kiliniki Mara Ya 1; Makubaliano - dakika 10, Pap Smear-dakika 20; Kutembelea kiliniki Mara Ya 2; Marejeleo ya Matokeo ya Pap Smear- dakika 10, Colposcopy na Biopsy (ikihitajika)- dakika 30; Kutembelea kiliniki mara ya 3; Marejeleo ya matokeo ya biopsy na kujadili juu ya njia za matibabu- dakika 30)

Kutembelea kiliniki mara ya 1:

Ukikubali kushiriki katika utafiti huu, kwanza utahitajika kuchunguzwa fupanyonga (Pelvic exam) kwa kupanguzwa sehemu yako ya siri kutumia burashi ili kuthibitisha ikiwa una dalili

inayoashiria kuwa na virusi vinavyoweza kusababisha saratani ya mlango wa uzazi (Pap Smear). Kuchunguzwa fupanyonga, kunamaanisha kuwa muuguzi atakagua sehemu zako za siri (uke wako). Pap smear itajumuisha utumiaji wa burashi ndogo iingizwayo kwenye kizazi ili kuchukua chembe chembe zitakazochunguzwa kwa kutumia “mikroskope”. Kisha utahitajika kurudi katika kiliniki baada ya majuma mawili ili kuchukua matokeo ya uchunguzi huu.

Kutembelea kiliniki mara ya 2:

Utarejea kwenye kiliniki baada ya wiki mbili ili kupata matokeo ya pap smear. Mara nyingi, matokeo huwa ni sawa (hakuna kasoro) na mtu hahitaji matibabu ya ziada, ila kurudia “Pap smear” tena baada ya kila miezi sita au kufanyaukaguzi na asidi asetiki (VIA) baada ya miezi sita. Ikiwa hujambo utaondoka utafitini katika kiwango hiki na kupokea matibabu mengine katika Kiliniki ya ukaguzi wa Ugonjwa wa saratani ya mlango wa kizazi ya Coptic Hope Centre, [*Coptic Hope Cervical Cancer Screening Program (CCSP)*] ampapo utapokea VIA, au utaulizwa kurudi katika kiliniki ya utafiti baada ya miezi sita ili kurudia “Pap smear”.

Wachunguzi wakipata chembe chembe zisizo za kawaida ambazo zinahitaji kutibiwa katika kiwango hiki, watakuelezea juu ya matibabu.

Lakini watafiti wakigundua chembe chembe zisizo za kawaida, watachunguza sehemu yako ya siri kwa makini. Kwanza utapimwa kama wewe ni mjamzito. Kama wewe ni mjamzito, Hakuna utafiti mwingine utaendelea hadi utakapojifungua mtoto. Watafiti watakutuma kupata matibabu ya dharura katika kituo cha Hope ili kuhakikisha hauambukizi mtoto wako virusi vinavyosababisha ukimwi (PMTCT). Kisha utaondoka utafitini katika kiwango hiki na kupokea matibabu mengine katika Kiliniki ya ukaguzi wa Ugonjwa wa saratani ya mlango wa kizazi ya Coptic Hope Centre, [*Coptic Hope Cervical Cancer Screening Program (CCSP)*].

Ikiwa wewe si mjamzito, watafiti wakigundua chembe chembe zisizo za kawaida, watachunguza sehemu yako ya siri kwa makini wakitumia chombo kiitwacho “colposcope”, ili kuchukua tishu, kiwango kama kipande kimoja cha mchele, utarabu huu unajulikana kama “Biopsy”. Tishu iliyochukuliwa, itachunguzwa kwa mikroskope na utajulishwa matokeo baada ya wiki nne hadi sita. Mwenendo huu wa kupima ambao majibu yake huwa ni sahihi utatuwezesha kujua kana kwamba unahitaji matibabu.

Kuna chembe chembe za ndani ya njia ya kizazi katikati ya mlango wa kizazi na kizazi na pia zinaendelea mpaka ndani ya kizazi. Hizi chembe chembe zinaitwa ‘glandular cells’. Ikiwa majibu ya pap smear itaonyesha ‘glandular’ cells si kawaida, daktari ata gwaruza kwa njia ya kizazi kutumia chombo chenye umbo cha kijiko ili sampuli ndogo ipatikane ya kukaguliwa. Utaratibu huu unaitwa ‘Endocervical curettage (ECC)’. Kukwaruza inafanywa wakati ‘colposcopy’ na inachukuwa dakika chache. Unaweza pata maumivu ya tumbo kidogo kama wakati unapo pata damu ya mwezi. Pia, kama chembe chembe za mlango wa kizazi kwa pap smear inaonyesha si kawaida na mabadiliko ya ugonjwa kwa mlango wa kizazi wakati wa colposcopy haionekani, hii kukwaruza itafanywa kama vile imeelezwa hapo mbeleni.

Kutembelea kiliniki mara ya 3:

Utarejea kwenye kiliniki kati ya wiki nne na wiki sitaili kupata matokeo ya biopsy. Iwapo tutagundua maambukizi yanayohitaji matibabu, tutakujulisha matibabu ambayo unaweza pata. Unaweza kujiunga na utafiti wa matibabu ya ugonjwa unaosababisha saratani ya mlango wa uzazi, ama unaweza chagua matibabu kwingine, au tunaweza kukutuma Hospitali Kuu ya Kenyatta (KNH) na uendelee kupokea matibabu yako ya kawaida katika Coptic Hope Center. Ukichagua kuendelea na utafiti wa matibabu au la, katika kiwango hii, tutakuondoa katika utafiti wa kupimwa mlango wa kizazi.

Tutatumia rekodi zako za afya ziliko HOPE Center ili kukusanya habari zote kuhusu afya yako ili kuelewa vyema dawa ambazo umekuwa ukitumia, hali ya afya yako kwa sasa na majibu ya vipimo kutoka maabara na pia jinsi umekuwa ukihudhuria kliniki.

Kuwasiliana na Washiriki

Tutakuuliza utupatie jinsi ya kuwasiliana nawe kama vile nambari yako ya simu ili tuweze kukupigia usipotembelea kiliniki kama ilivyopangwa. Tunaweza kukuuliza kuhusu afya yako au ulipokuwa katika mawasiliano haya. Ni muhimu kwako kutembelea kiliniki kama ilivyopangwa. Iwapo hatuwezi kuwasiliana nawe kwa njia ya simu, tunaweza kukutembelea kwako nyumbani. Iwapo tutakutembelea kwako nyumbani, hatutavalia mavazi yatakayoashiria kuwa sisi ni wahudumu katika hospitali. Unaweza kutuarifu iwapo hupendi njia hii ya mawasiliano. Usiporejea kupokea matokeo yako ya pap smear au biospy, tutawasiliana na wafanyi kazi wa mapokezi ya Coptic waweke mawaidha kwa faili yako ili unaporejea kwa matibabu yako ya kawaida, utakumbushwa kupitia kwenye kiliniki ya utafiti kupokea matokeo yako.

Kamataratibu zakudhibiti ubora wautafitizinaonyesha kuwa Papsmearyakomatokeonitofautinamatokeoya awali uliyo pewa, tutakuuliza urudiklinikikupokeamatokeo yakompya nakujadilimatibabu inayoweze kana. Katika hatua hii, unaweza kuulizwakamaungependakujandikisha tena katika utafiti.

Athari na usumbufu wa kushiriki kwenye utafiti huu.

Haya ni maelezo ya athari na usumbufu unayoweza kukumbana nayo kwa kushiriki katika utafiti huu. Athari na usumbufu mwingine ambao haujatajwa unaweza kutokea pia.

Kiingilio (Screening)

Kuchukua sampuli kutoka kwenye fupanyonga (pelvic) kwaweza sababisha;

- kukerwa kidogo wakati ukaguzi ukiendelea,
- kutokwa na damu kidogo kwenye uke wako baadaye kwa siku moja au mbili, na
- kupata uchungu mdogo wa tumbo kama ule wa damu ya mwezi kwa dakika tano hivi.

Ukifanyiwa uchunguzi wa “Biopsy” kwenye fupanyonga (Pelvic), tunakusihia usishiriki ngono kwa muda wa siku kumi. Hii ni kwa sababu ikiwa una virusi vinavyosababisha ukimwi na (HIV) ili hali mpenzi hana, atakuwa katika hatari kubwa zaidi ya kuambukizwa na virusi hivi vya ukimwi. Ikiwa mpenzi wako ana virusi vinavyosababisha ukimwi, atakuwa katika hatari kubwa ya kuambukizwa tena na virusi stahimilivu vinavyosababisha ukimwi. Pia, wewe unahitaji muda wa kupona na utajiweka katika hatari ya maambukizi usipojipatia muda huu. Iwapo unahitaji msaada wowote wa kuongelea mambo haya na mpenzi wako, muuguzi, daktari au mshaurikatika hospital atakusaidia. Tutakupatia mipira ya kondomu bila malipo iwapo hutaweza kutoshiriki katika ngono.

Athari nyengine unayoweza kukumbana nayo ni wasiwasiya kuwa una ugonjwa katika mlango wako wa uzazi. Unaweza kuongea kuhusu jambo hili na muuguzi wa kitafiti, daktari, au ukitaka kuongea na mshaurikatika hospitali, tutakusaidia kupanga haya.

Tunaweza kugundua katika kiingilio kuwa una maambukizo. Iwapo utapatikana na maambukizo, tutakupaa maagizo ya kupata dawa. Madhara ya dawa hizi yanaweza kuwa kuumwa na tumbo, kutapika na kuharisha, usikivu wa mwangaza, mwasho, mzio mkali mmenyuko unaoweza kusababisha kifo, kizunguzungu, kuumwa na kichwa, kuchanganyikiwa, kupigapiga kwenyemasikio au kupoteza usikivu, mshtuko wa mwili (jerking of the body), maumivu ya moyo, machafuko ya damu, shida ya kisukari, ugonjwa wa maini, ugonjwa wa figo na uchungu au kuganda kwenye mikono na miguu. Ikiwa utahisi mojawepo ya athari hizi, mfahamishe mkaguzi katika utafiti huu mara moja.

Uwekaji wa siri

Wahudumu katika utafiti huu, watakusanya habari ya binafsi na waweza kuona aibu kuongea kuhusu habari zako za ndani kama vile; idadi ya wapenzi ambao ushakuwa nao. Unaweza kuchagua kutojibu maswali yoyote. Pia, ukija kiliniki, waweza kukutana na wagonjwa wengine ambao umewajua mbeleni. Hatuna mpango wa kupeana habari yako ya kibinafsi kwamtu yeyote nje ya utafiti huusipokuwa watafiti au wachunguzi walioteuliwa. Wakati mwingine kamati inayosimamia utafiti huu watachunguza habari ya utafiti huu ili kuhakikisha hakuna jambo lolote ambalo ni kinyume na sheria au kinyume na maadili litakalotendwa. Habari yako ya kibinafsi italindwa ipasavyo ikiwa hayo yatatendeka na hakuna mtu yeyote atakayeona habari hiyo nje ya uchunguzi huu. Kamamdamini watafiti, CDC wanaweza kufuatilia au kukaguashughuli zautafiti. Sababu hii itakuwahakikishakuwa utafiti unafanyika kwanji inayotakiwa, napia kuhakikishakwamba hakiyakona afya yako inalindwa. Matibabuyakotaarifa binafsi itakuwasiri.

Mbadala wa kujiunga na utafiti huu

Ukichagua kutoshiriki kwenye utafiti huu, utaendelea kuhudumiwa kikamilifu na kupata matibabu na dawa za ART kutoka “Coptic Hope Centre” bila malipo. Unaweza pia kupata uchunguzi wa saratani ya mlango wa nyumba ya uzazi bila malipo yeyote katika kiliniki hii bila kushiriki katika utafiti huu.

Manufaa kutokana na utafiti huu

Kushiriki kwako katika utafiti huu, kutatuwezesha kuelewa zaidi kuhusu ugonjwa wa mlango wa nyumba ya uzazi. Kwa kujiunga na utafiti huu, utapata kupimwa mlango wako wa uzazi bila malipo. Kama ungependa kujua matokeo ya utafiti huu, unaweza kuwasiliana na ofisi ya utafiti huu miezi sita au mwaka mmoja baada ya kumaliza utafiti huu.

Gharama na Fidia ya Majeraha

Hakuna gharama yoyote ya kushiriki katika utafiti huu, ila muda wako pekee. Utafiti huu utalipia gharama ya uchanguzi wowote utakayopokea kwenye kiliniki ya utafiti. Ikiwa Utapata majeraha kutokana na kushiriki katika utafiti huu, utafiti huu utafidia gharama yote ya malipo ya utunzaji. Ikiwa unafikiri kuwa umepata jeraha ama maumivu kutokana na utafiti huu, wasiliana na wahudumu wa utafiti huu, (Daktari Michael Chung (020-272-2710) au Daktari Nelly Mugo (020-273-6744) au Daktari Evans Malava (0721 289 733)) mara moja. Watakuhudumia kwa matibabu au kukupendekeza kwa matibabu.

Kuhifadhi kwa Sampuli na Data na Matumizi ya sampuli hizi kwa utafiti wa baadaye

Tungependa kuhifadhi habari yako ya matibabu na sampuli za pap smear na biopsy katika Hospitali ya Coptic na katika Hospitali kuu ya Kenyatta kwa madhumuni ya utafiti wa baadaye. Utafiti huu waweza kufanywa na Chuo kikuu cha Washington au na watafiti wengine wanaofanya kazi nasi katika utafiti huu kwa muda wa miaka kumi baada ya mwisho wa kufuatiliwa katika utafiti. Tutatumia data na sampuli hizi kwa minajili ya utafiti unaoegemea na saratani ya mlango wa kizazi na UKIMWI. Kabla sampuli yako kutoka kwenye kiliniki, itapewa kifichomaalumu Jina lako halitakuwemo. Jina lako litahusishwa na kificho hiki kwa miaka mitano pekee yake baada ya kukamilisha utafiti huu. Baada ya muda huu, kihusishi baina ya jina lako na kificho kilichoko katika sampuli na data kitaangamizwa. Bodi za taasisi ya mapitio ni kamati zinazoagalia usalama na haki za washiriki wa utafiti katika Hospitali kuu ya Kenyatta na Chuo kikuu cha Washington. Lazima ziidhinishe utafiti wowote wa baadaye utakaotumia sampuli na data yako. Ikiwa hupendi sampuli zako zihifadhiwe kwa minajili ya utafiti wa baadaye, unaweza kushiriki katika utafiti huu na sampuli zako zitaharibiwa mara tu uchunguzi utakapokamilika. Ukikubali uhufadhi wa sampuli zako leo, kisha ubadilishe nia yako baadaye kabla ya utafiti kukamilika, wafahamishe watafiti ambao watahakikisha kuwa sampuli zako hazitahifadhiwa kwa minajili ya utafiti wa baadaye. Hatutauza sampuli zako. Uchunguzi utakao fanya kwenye sampuli zako waweza kutuelekeza katika uvumbuzi mpya. Hatuna mpango wowote wa kugawana nawe pesa zozote au faida nyinginezo zitakazotokana na uvumbuzi huu ..

Maelezo ya ziada

Habari kuhusu utambuzi wa ugonjwa wako ni siri na tutaweka rekodi ya afya yako kwa afisi inayofungwa. Habari kuhusu uchunguzi na ushiriki wako kwenye utafiti huu utatolewa kwako na kikundi cha watafiti pekee wala si mtu mwengine yeyote nje ya utafiti huu bila idhini yako. Ikiwa utapeana idhini, tutawafahamisha habari yako daktari wa Coptic Hope Center. Habari hii inaweza kuwasaidia kukupa matibabu na huduma bora ya UKIMWI. Rekodi zako zote zitahifadhiwa katika sehemu zilizofungwa na kifuli na tarakilishi zote zitakuwa zina neno la siri la kuhakikisha ulinzi.

Wakati mwengine, wafanyikazi wa chuo kikuu cha Washington wanakagua utafiti kama huu ili kuhakikisha unafanywa kwa njia inayofaa na iliyo halali. Ikiwa ukaguzi wa utafiti huu utafanyika, rekodi zako zaweza kuchunguzwa. Waweza kukataa kushiriki au waweza kuondoka kwenye utafiti huu wakati wowote bila kuadhibiwa au kupoteza faida iliyokuwa haki yako. Tutakueleza ikiwa kuna habari geni zinazohusu matibabu ambayo tunatafiti ili uweze kuamua iwapo utajiondoa kwenye utafiti huu. Uhusiano wako na wafanyikazi wa huduma za Coptic Hope Centre for infectious diseases, hautaadhiriwa kwa vyovyote ikiwa utakubali kushiriki kwenye utafiti huu ama ukiingia kwenye mpangilio halafu baadaye uondoke kabla ya kumaliza uchunguzi. Tafadhali wafahamishe

wahudumu wa utafiti huu ikiwa utaamua kuondoka katika utafiti huu. Unaweza kuulizwa kupeana sampuli za mwisho lakini waweza kukataa.

Wahudumu katika utafiti huu wanaweza kuamua kukuondoa kwenye utafiti huu wakigundua kuwa unaweza kupata madhara ukiendelea kushiriki. Unaweza kuondolewa kwenye utafiti huu ikiwa wahudumu watagundua kuwa huwezi kufuata masharti ya usalama.

Maswali yeyote kuhusu utafiti huu, ama athari mbaya kutokana na uchunguzi huu yapaswa kuelekezwa kwa mtafiti anayekuhudumia au Daktari Michael Chung (272-2710), Daktari Samah Sakr (272-4737), au Daktari Nelly Mugo (273-6744).

Je, una maswali yeyote?

Sahihi ya mtafiti..... Tarehe.....

Jina la Mtafiti.....

Muhusika

Nimeelezwa juu ya utafiti huu. Najitolea kwa hiari kushiriki kwenye utafiti huu. Nimewapa watafiti ruhusa ya kutumia rekodi zangu za utabibu kama ilivyopendekezwa kwenye fomu hii ya makubaliano. Nimekuwa na fursa ya kuuliza maswali. Nikiwa na maswali yeyote kuhusu utafiti huu baadaye, naweza kumuuliza mojawapo wa watafiti waliotajwa kwenye fomu hii. Nikiwa na maswali kuhusu haki yangu kama mhusika kwenye utafiti, naweza piga simu kwa kamati ya uchunguzi wa maadili walioko Hospitali kuu ya Kenyatta kwa kutumia nambari 020-726-300. Nikitaka, nitapewa nakala ya fomu hii yangu binafsi.

Tafadhali tia alama, ufupi wa jina na anwani katika chaguo moja:

_____ Nimekubali kuweka sampuli na data kwa minajili ya utafiti wa baadaye

_____ Sijakubali kuweka sampuli na data kwa minajili ya utafiti wa baadaye

Sahihi au alama ya kidole cha gumba cha mshiriki _____

Tarehe _____

Jina la Mshiriki lililochapishwa _____

Nakala kwa Mtafiti na Mshiriki

MAKUBALIANO YA PAMOJA

KISWAHILI CONSENT FORM

**MATIBABU YA SARATANI YA MLANGO WA KIZAZI; CRYOTHERAPY AU
LEEP**

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Mwenyekiti wa kamati ya uchunguzi wa maadili; Proffesa A. N. Guantai, 020 2726300, Ext. 44102, 44355 unaweza kuwasiliana naye kwa maswala ya utafiti na maadili ya washiriki wa utafiti huu.

Ujumbe wa Watafiti

Tunakuuliza kushiriki katika utafiti huu wa kitaalam. Lengo la fomu hii ya idhini ni kukupa habari itakayokufahamisha na kukusaidia kuamua kama ungelipenda kushiriki katika utafiti huu au la. Tafathali soma maelekezo haya kwa makini. Unaweza kuuliza maswali kuhusu nia/kusudi ya utafiti huu; unavyohitajika kufanya katika utafiti, uwezekano wa kuwepo na madhara au manufaa yeyote, haki yako kama aliyejitolea, na mambo mengine kuhusu utafiti huu au chochote usichoelewa kwenye fomu hii. Tukishajibu maswali yako yote, unapaswakuamua kushiriki kwenye utafiti huu au la. Kukubali kushiriki kwenye utafiti huu kwa hiari yako kunamaanisha kuwa umeelezwa na umelewa yote yanayohusika na kukubaliana nayo. Ukipenda, tutakupa nakala ya fomu hii kujihifadhia. Tafadhali tujulishe iwapo ungependa kutumia lugha ya Kingereza.

Lengo la Utafiti huu.

Mathumuni ya mradi huu wa utafiti, ni kutafuta njia iliyo bora ya kutibu ugonjwa unaosababisha saratani ya mlango wa kizazi (cervical cancer). Umeulizwa kushiriki katika utafiti huu kwa sababu;

- Umeambukizwa maradhi ya UKIMWI
- Unapata matibabu katika kituo cha matibabu cha Hope
- Hauna Mimba
- Hauna historia ya shida ya kuvuja damu
- Haujapata kutolewa sehemu yako ya uzazi (uterasi)
- Umewahi fanya ngono
- Haujawahi kupata matibabu yoyote ya mlango wa kizazi hapo mbeleni
- Umepatikana kuwa na chembe chembe zisizo za kawaida kwa mlango wa uzazi kwenye pap smear ua biopsy.

Saratani ya aina hii huambukiza wanawake wengi wenye umri mdogo na hata wa makamo nchini Kenya. Wanawake walioambukizwa na virusi vinavyosababisha ukimwi (HIV) wana hatari kubwa ya kuambukizwa saratani ya mlango wa kizazi (cervical cancer), kuliko wanawake wengine ambao bado hawajaambukizwa na HIV. Hata hivyo, kwa wanawake walioambukizwa na virusi vinavyosababisha ukimwi, saratani ya mlango wa kizazi si ya kawaida. Lakini katika wanawake walioambukizwa na virusi vinavyosababisha ukimwi, maambukizo ya mlango wa kizazi usio saratani ya mlango wa kizazi ni wa kawaida. Maambukizo haya husababishwa na virusi vya “Papiloma” (HPV). Ni muhimu kutambua na kutibu maambukizo haya kabla hayajageuka kuwa saratani ya mlango wa kizazi.

Utafiti huu utalinganisha namna mbili za ukaguzi wa mlango wa kizazi; Cryotherapy na LEEP (loop electrosurgical excision procedure). Njia hizi mbili hutumika sana kwa wanawake wengi duniani na sio geni. Tutakuelezea njia hizi mbili za utabibu. Katika utafiti huu, tutakuwa na washiriki 400. Utafiti huu utatuwezesha kutambua matibabu bora itakayowawezesha wanawake wanaoishi na virusi vinavyosababisha ukimwi (HIV) kuishi maisha bila saratani ya mlango wa kizazi.

Hatua ya kushiriki katika utafiti

Washirika wote watahitajika kutembelea kliniki mara 5 au 8 kwa muda usiozidi miaka 2. Kila mara, mushirika atatumia dakika 15 hadi 40 katika kiliniki.

Utaratibu wa kupeana matibabu bila mapendeleo (Randomization)

(Marejeleo ya Matokeo ya Biopsy – dakika 10, Utaratibu wa kupeana matibabu bila mapendeleo (ikihitajika) – dakika 30)

Matokeo ya pap smear na biopsy yameonyesha ya kuwa uko na maambukizo yanayohitaji matibabu. ukiamua kujiunga na utafiti huu, tutatumia mojawapo ya njia mbili ambazo ni za kutumainika. Njia moja inaitwa “Cryotherapy”. Njia hii inatia barafu sana (freeze) kwenye sehemu yako ya uzazi iliyoambukizwa na ugonjwa. Unaweza pewa dawa ya kumeza kupunguza maumivu baada ya Cryotherapy. Njia ya pili inajulikana kama, “LEEP”. Njia hii inatumia chuma kidogo kilichopashwajoto ili kutoa sehemu iliyoambukizwa baada ya kugandishwa kwa dawa. Utahudumiwa kwa njia mojawapo ya hizi bila mapendeleo ili kudhibitisha ni njia ipi iliyo bora kuliko nyengine.

Njia ya kuchagua matibabu bila mapendeleo ni kama “kuzungusha peni”, kila upande wa peni una uwezo uliyo sawa, kama ilivyo njia hizi mbili za matibabu. Hatujui kwa hakika ni njia gani ya matibabu utakayopata hadi tutakapofungua bahasha iliyo na kijikaratasi kitakachotujulisha ni njia gani ya matibabu utakayopata. Sio sisi wala wewe utakayechagua njia ya matibabu, bali ni bahasha iliyo na vijikaratasi vitakavyo tuelekeza ni njia ipi itakayotumika baada ya kufungua ile bahasha. Tutakuashiria kutumia mojawapo ya njia hizi mbili bila mapendeleo (kuegemea njia moja) kwa sababu hatujui ni njia ipi iliyo bora kutumiwa na wanawake walioambukizwa na virusi vinavyosababisha ukimwi (HIV).

Tukigundua kuwa sehemu ya kizazi iliyoambukizwa ni kubwa mno, na haiwezi kutibika sawa sawa kwa “Cryotherapy”, tutakutibu tukitumia njia ya “LEEP”. Utapata matibabu ya LEEP bila malipo yoyote.

Ikiwa sehemu iliyoambukizwa haiwezi kutibika vizuri kwa “Cryotherapy” au “LEEP”, utaondoka kwenye utafiti huu na kuelekezwa kwenye hospitali kuu ya Kenyatta (KNH) utakapopata matibabu ya aina nyengine ya matibabu kwa bei nafuu katika hospitali ya serikali. Watafiti watakupatia fomu zako za matibabu ambazo ni muhimu kwa matibabu yako. Hizi fomu hazitaonyesha kuwa wewe ni mshirika wa utafiti huu. Baada ya matibabu utakayopokea kutoka KNH, tutakufuatilia kwa miaka 2 na kukupa matibabu.

Tukikupa njia mojawapo ya matibabu yaliyotajwa (Cryotherapy au LEEP), vijiko viwili vya chai (tea spoons [10mls]) vya damu vitatolewa ili kuhesabu chembe chembe za CD4 na kuthibitisha kiwango cha virusi vinavyosababisha ukimwi (HIV) katika damu yako kwa wakati huu. Kisha tutapanguza kwa burashi sehemu yako ya kizazi na kukagua kiwango cha virusi vinavyosababisha ukimwi. Mwisho, tutapangusa sehemu yako ya kizazi ili kukagua kiwango cha virusi vya “Papiloma” (HPV) ambavyo ni viini vinavyoambukiza sehemu ya kizazi na kusababisha saratani ya uzazi (Cervical cancer).

Tutatumia rekodi zako za afya ziliko HOPE Center ili kukusanya habari zote kuhusu afya yako ili kuelewa vyema dawa ambazo umekuwa ukitumia, hali ya afya yako kwa sasa na majibu ya vipimo kutoka maabara na pia jinsi umekuwa ukihudhuria kiliniki.

Mpangilio baada ya kupata matibabu bila mapendeleo

(Kutembelea kiliniki Mara Ya 2-8; Marejeleo ya Matokeo na usufi wa mlango wa kizazi-dakika30)

Ukipata mojawapo ya matibabu (Cryotherapy au LEEP) bila mapendeleo, watafiti watakuuliza urudi kwenye kiliniki tena wiki ya kwanza (1), pili (2) na tatu (3) baada ya matibabu.

Kila mara utakaporudi kiliniki, vijiko viwili vidogo vya damu vitatolewa na kisha kupanguza sehemu yako ya kizazi ili kuthibitisha kiwango cha virusi vinavyosababisha ukimwi (HIV). Tungependa kujua kama kiwango hiki kimeongezeka baada ya matibabu na kitarudi katika hali yake ya kawaida lini?

Ukipata mojawapo ya matibabu (Cryotherapy au LEEP), tutakuuliza kurudi kiliniki ya utafiti mwezi wa sita (6), kumi na mbili (12), kumi na nane (18) na pia mwezi wa ishirini na nne (24) baada ya matibabu yako ya kwanza ili kurejelea “Pap smear” kudhibitisha Ikiwa sehemu yako ya kizazi iliyoambukizwa imeondolewa kabisa na hakuna tishu iliyojitenga au isiyo ya kawaida iliyojitengeza. Tutapanguza sehemu yako ya kizazi (cervix), ili kukagua virusi vya “Papiloma” (HPV) kwa wakati huu na kutoa vijiko viwilividogo vya damu ili kuhesabu chembe chembe za CD4 na kudhibitisha viwango vya virusi vinavyosababisha ukimwi (HIV). Tukigundua hali isiyo ya kawaida, tutakutibu kwa kutumia LEEP, kisha tutakuelekeza katika hospitali kuu ya Kenyatta (KNH) kwa matibabu ya ziada yaliyo nafuu.

Kuwasiliana na Washiriki

Tutakuuliza utupatie jinsi ya kuwasiliana nawe kama vile nambari yako ya simu ili tuweze kukupigia usipotembelea kiliniki kama ilivyopangwa. Tunaweza kukuuliza kuhusu afya yako au ulipokuwa katika mawasiliano haya. Ni muhimu kwako kutembelea kiliniki kama ilivyopangwa. Tunataka kukufuatilia vyema na kwa makini ili kutibu ugonjwa wowote katika mlango wa kizazi ulio nao sasa au utakaoupata baadaye. Iwapo hatuwezi kuwasiliana nawe kwa njia ya simu, tunaweza kukutembelea kwako nyumbani. Iwapo tutakutembelea kwako nyumbani, hatutavalia mavazi yatakayoashiria kuwa sisi ni wahudumu katika hospitali. Unaweza kutuarifu iwapo hupendi njia hii ya mawasiliano.

Athari na usumbufu wa kushiriki kwenye utafiti huu.

Haya ni maelezo ya athari na usumbufu unayoweza kukumbana nayo kwa kushiriki katika utafiti huu. Athari na usumbufu mwingine ambao haujatajwa unaweza kutokea pia.

Matibabu

Kuna uwezekano wa madhara au maumivu ukipata matibabu haya. Ukishiriki katikautibabu wa “Cryotherapy”, waweza kuhisi;

- maumivu madogo ya tumbo kwa muda unaopungua dakika kumi,
- kutokwa na majimaji kwenye uke wako kwa muda wa wiki mbili (au zaidi).
- Kuvuja damu
- Maambukizo ambayo tunaweza kutibu hapa katika kiliniki au katika hospitali kwa nadra sana. Maambukizo haya yanaweza kusababisha homa, baridi, kutokwa na jasho usiku, au majimaji meupe kutoka kwenye uke wako.

Ukitibiwa kwa LEEP, unaweza;

- kuvuja damu baada au wakati matibabu haya yanapoendelea. (Unashauriwa kurudi kiliniki Ikiwa uvujaji huu wa damu utaanza, kisha kuendelea baada ya matibabu kwa ukaguzi wa ziada.)
- Kuna uwezekano wa kuambukizwa kwenye eneo la matibabu.

Madhara haya yote yanaweza kutibiwa bila malipo yeyote katika kiliniki hii. Matibabu haya yanaweza kuwa;

- kupewa dawa ya kuuguza maambukizi.
- Kuingiza pamba kwenye uke wako kuzuia kuvuja damu,
- Kushona kwenye mlango wako wa uzazi ili kuzuia kuvuja kwa damu au
- Kulazwa hospitalini kwa makali ya maambukizo au uvujaji wa damu.

Tafadhali tufahamisha ikiwa una madhara haya baada ya matibabu.

Baada ya kupokea matibabu ya “Cryotherapy”, au LEEP tunakusihia usishiriki katika kitendo cha gono kwa muda wa juma nne, sababu ni kuwa waweza kumwambukiza mwenzi kwa urahisi. Pia kuna uwezekano wa wewe kupata maambukizo. Tunaweza kukusaidia kuongelea jambo hili na mpenzi wako.

Washiriki wote watakaopata matibabu ya “Cryotherapy” au “LEEP” watapata maagizo ya dawa ya “antibiotiki”. Dawa hii ya antibiotiki itashirikisha doxycycline, metronidazole or norfloxacin.

Madhara ya dawa hizi yanaweza kuwa usumbufu wa tumbo (Lazima usikunywe tembo/pombe kwa siku 2) kutapika na kuharisha usikivu wa mwangaza, mwasho, mzio mkali mmenyuko unaoweza kusababisha kifo, kizunguzungu, kuumwa na kichwa, kuchanganyikiwa, kupigapiga kwenyemasikio au kupoteza usikivu, mshutuko wa mwili (jerk of the body), maumivu ya moyo, machafuko ya damu, shida ya kisukari, ugonjwa wa maini, ugonjwa wa figo na uchungu au kuganda kwenye mikono na miguu. Ikiwa utahisi mojawepo ya athari hizi, mfahamisha mkaguzi katika utafiti huu mara moja.

Baada ya kupata matibabu haya, kuna uwezekano wa kupata matatizo utakapobeba mimba baadaye. Baada ya kupata matibabu ya “LEEP”, wanawake wengine wamepata matatizo wanapobeba mimba kama vile; maumivu ya mapema wakati wa kujifungua au shida katika mlango wa mfuko wa uzazi. Katika Wanawake wachache, mlango wa mfuko wa uzazi huziba na inabidi ulegezwe. Wanawake wengine huvuja majimaji ya uzazi kabla ya muda ufaao wa kuzaa mtoto pengine kutokana na operesheni ya mlango wa mfuko wa uzazi (cervix).

Utoaji wa damu

Tutakusanya sampuli za damu, vijiko 2 (10 mls), kutoka kwako kwa kutumia shindano na “sirinji”. Kudungwa shindano kwaweza kuwa chungu na yaweza kuacha alama kidogo kwa muda mfupi. Yaweza kusababisha maambukizo au kuzirai.

Uwekaji wa siri

Wahudumu katika utafiti huu, watakusanya habari ya binafsi na waweza kuona aibu kuongea kuhusu habari zako za ndani kama vile; idadi ya wapenzi ambao ushakuwa nao. Unaweza kuchagua kutojibu maswali yoyote. Pia, ukija kiliniki, waweza kukutana na wagonjwa wengine ambao umewajua mbeleni. Hatuna mpango wa kupeana habari yako ya kibinafsi kwamtu yeyote nje ya utafiti huu isipokuwa watafiti au wachunguzi walioteuliwa. Wakati mwingine kamati inayosimamia utafiti huu watachunguza habari ya utafiti huu ili kuhakikisha hakuna jambo lolote ambalo ni kinyume na sheria au kinyume na maadili litakalotendwa. Habari yako ya kibinafsi italindwa ipasavyo ikiwa hayo yatatendeka na hakuna mtu yeyote atakayeona habari hiyo nje ya uchunguzi huu. Kamamdamini watafiti, CDC wanaweza kufuatilia aukukaguashughuli zautafiti. Sababu hii itakuwahakikishakuwa utafiti unafanyika kwanja inayotakiwa, napia kuhakikishakwamba hakiyakonaafya yako inalindwa. Matibabu yako tarifa binafsi itakuwasiri.

Mbadala wa kujiunga na utafiti huu

Ukichagua kutoshiriki kwenye utafiti huu, utaendelea kuhudumiwa kikamilifu na kupata matibabu na dawa za ART kutoka “Coptic Hope Centre” bila malipo. Unaweza pia kupata uchunguzi wa saratani ya mlango wa nyumba ya uzazi bila malipo yeyote katika kiliniki hii bila kushiriki katika utafiti huu. Unaweza kutibiwa na Cryotherapy katika Hospitali ya kiliniki ya Hope hata ukichagua kutojiunga na utafiti huu. Pia, tunaweza kukutuma kupewa huduma katika hospitali kuu ya Kenyatta, ambapo malipo ni kama ya hospitali za serikali.

Manufaa kutokana na utafiti huu

Kushiriki kwako katika utafiti huu, kutatuwezesha kuelewa zaidi kuhusu ugonjwa wa mlango wa nyumba ya uzazi. Pia, kushiriki kwako katika utafiti huu kutachangia katika elimu ya nyanja hii ambayo yaweza kutoa mabadiliko katika jinsi ya ugonjwa wa saratani ya mlango wa uzazi unavyogunduliwa na kutibiwa. Kwenye nchi inayoendelea kama Kenya. Kama ungependa kujua matokeo ya utafiti huu, unaweza kuwasiliana na ofisi ya utafiti huu miezi sita au mwaka mmoja baada ya kumalizika utafiti huu.

Gharama na Fidia ya Majeraha

Hakuna gharama yeyote ya kushiriki katika utafiti huu, ila muda wako pekee. Utafiti huu utalipia gharama ya uchunguzi na matibabu yoyote utakayopokea kwenye kiliniki ya utafiti. Ikiwa utapata majeraha kutokana na kushiriki katika utafiti huu, utafiti huu utafidia gharama yote ya malipo ya utunzaji. Matibabu yatahusisha dawa zinazopambana na vidudu vilivyo hai (antibiotics), dawa zinazopunguza maumivu na njia zinazoachisha kuvuja damu. Ikiwa unafikiri kuwa umepata jeraha ama maumivu kutokana na utafiti huu, wasiliana na wahudumu wa utafiti huu, (Daktari Michael Chung (020-272-2710) au Daktari Nelly Mugo (020-273-6744) au Daktari Evans Malava (0721 289 733)) mara moja. Watakuhudumia kwa matibabu au kukupendekeza kwa matibabu.

Kuhifadhi kwa Sampuli na Data na Matumizi ya sampuli hizi kwa utafiti wa baadaye

Tungependa kuhifadhi habari yako ya matibabu na sampuli za damu yako na mlango mfuko wa uzazi, Kenya katika Hospitali ya Coptic na katika Hospitali kuu ya Kenyatta kwa madhumuni ya utafiti wa baadaye. Utafiti huu waweza kufanywa na Chuo kikuu cha Washington au na watafiti wengine wanaofanya kazi nasi katika utafiti huu kwa muda wa miaka kumi baada ya mwisho wa kufuatiliwa katika utafiti. Tutatumia data na sampuli hizi kwa minajili ya utafiti unaoegemea na

saratani ya mlango wa kizazi na UKIMWI. Kabla sampuli yako kutoka kwenye kiliniki, itapewa kifichomaalumu Jina lako halitakuwemo. Jina lako litahusishwa na kificho hiki kwa miaka mitano pekee yake baada ya kukamilisha utafiti huu. Baada ya muda huu, kihusishi baina ya jina lako na kificho kilichoko katika sampuli na data kitaangamizawa. Bodi za taasisi ya mapitio ni kamati zinazoangalia usalama na haki za washiriki wa utafiti katika Hospitali kuu ya Kenyatta na Chuo kikuu cha Washington. Lazima ziidhinishe utafiti wowote wa baadaye utakaotumia sampuli na data yako. Ikiwa hupendi sampuli zako zihifadhiwe kwa minajili ya utafiti wa baadaye, unaweza kushiriki katika utafiti huu na sampuli zako zitaharibiwa mara tu uchunguzi utakapokamilika. Ukikubali uhifadhi wa sampuli zako leo, kisha ubadilishe nia yako baadaye kabla ya utafiti kukamilika, wafahamisha watafiti ambao watahakikisha kuwa sampuli zako hazitahifadhiwa kwa minajili ya utafiti wa baadaye. Hatutauza sampuli zako. Uchunguzi utakao fanya kwenye sampuli zako waweza kutuelekeza katika uvumbuzi mpya. Hatuna mpango wowote wa kugawana nawe pesa zozote au faida nyinginezo zitakazotokana na uvumbuzi huu ..

Maelezo ya ziada

Habari kuhusu utambuzi wa ugonjwa wako ni siri na tutaweka rekodi ya afya yako kwa afisi inayofungwa. Habari kuhusu uchunguzi na ushiriki wako kwenye utafiti huu utatolewa kwako na kikundi cha watafiti pekee wala si mtu mwingine yeyote nje ya utafiti huu bila idhini yako. Ikiwa utapeana idhini, tutawafahamisha habari yako daktari wa Coptic Hope Center. Habari hii inaweza kuwasaidia kukupa matibabu na huduma bora ya UKIMWI. Baadhi ya sampuli zako ikiwa ni pamoja na sampuli za damu na sampuli kutoka kwenye mlango wa mfuko wa uzazi zaweza kutumwa USA au Uropa kwa uchunguzi. Habari yako na sampuli zako zitalindwa ikiwa haya yatatendeka. Rekodi zako zote zitahifadhiwa katika sehemu zilizofungwa na kifuli na tarakilishi zote zitakuwa zina neno la siri la kuhakikisha ulinzi.

Wakati mwingine, wafanyikazi wa chuo kikuu cha Washington wanakagua utafiti kama huu ili kuhakikisha unafanywa kwa njia inayofaa na iliyo halali. Ikiwa ukaguzi wa utafiti huu utafanyika, rekodi zako zaweza kuchunguzwa. Waweza kukataa kushiriki au waweza kuondoka kwenye utafiti huu wakati wowote bila kuadhibiwa au kupoteza faida iliyokuwa haki yako. Tutakueleza ikiwa kuna habari geni zinazohusu matibabu ambayo tunatafiti ili uweze kuamua iwapo utajiondoa kwenye utafiti huu. Uhusiano wako na wafanyikazi wa huduma za Coptic Hope Centre for infectious diseases, hautaadhiriwa kwa vyovyote ikiwa utakubali kushiriki kwenye utafiti huu ama ukiingia kwenye programu halafu baadaye uondoke kabla ya kumaliza uchunguzi. Tafadhali wafahamisha wahudumu wa utafiti huu ikiwa utaamua kuondoka katika utafiti huu. Unaweza kuulizwa kupeana sampuli za mwisho lakini waweza kukataa.

Wahudumu katika utafiti huu wanaweza kuamua kukuondoa kwenye utafiti huu wakigundua kuwa unaweza kupata madhara ukiendelea kushiriki. Unaweza kuondolewa kwenye utafiti huu ikiwa wahudumu watagundua kuwa huwezi kufuata masharti ya usalama.

Utarejeshewa gharama ya usafiri ya Ksh 300 mara utakaporudi katika kiliniki kwa sababu zinazohusiana na utafiti huu. Utapata pesa za usafiri kutoka kwa mhudumu wa mapokezi.

Maswali yeyote kuhusu utafiti huu, ama athari mbaya kutokana na uchunguzi huu yapaswa kuelekezwa kwa mtafiti anayekuhudumia au Daktari Michael Chung (272-2710), Daktari Samah Sakr (272-4737), au Daktari Nelly Mugo (273-6744).

Je, una maswali yeyote?

Sahihi ya mtafiti.....

Tarehe.....

Jina la Mtafiti.....

Mhusika

Nimeelezwa juu ya utafiti huu. Najitolea kwa hiari kushiriki kwenye utafiti huu. Nimewapa watafiti ruhusa ya kutumia rekodi zangu za utabibu kama ilivyopendekezwa kwenye fomu hii ya makubaliano. Nimekuwa na fursa ya kuuliza maswali. Nikiwa na maswali yeyote kuhusu utafiti huu baadaye, naweza kumuuliza mojawapo wa watafiti waliotajwa kwenye fomu hii. Nikiwa na maswali kuhusu haki yangu kama mhusika kwenye utafiti , naweza piga simu kwa kamati ya uchunguzi wa maadili walioko Hospitali kuu ya Kenyatta kwa kutumia nambari 020-726-300. Nikitaka, nitapewa nakala ya fomu hii yangu binafsi. Tafadhali tia alama , ufupi wa jina na anwani katika chaguo moja:

_____Nimekubali kuweka sampuli na data kwa minajili ya utafiti wa baadaye

_____Sijakubali kuweka sampuli na data kwa minajili ya utafiti wa baadaye

Sahihi au alama ya kidole cha gumba cha mshiriki _____

Tarehe _____

Jina la Mshiriki lililochapishwa _____

Nakala kwa Mtafiti na Mshiriki

ENROLLMENT

Study ID Number ___ ___ **Hope ID Number** ___ ___ **Interviewer Number** ___

Date of interview (day/month/year) ___ ___ / ___ ___ / ___ ___

Agreed to store samples: Yes No Don't know Other, specify

A: SOCIODEMOGRAPHIC

1. Date of birth (day/month/year) ___ ___ / ___ ___ / ___ ___

2. Age _____ years

3. How many years of education did you complete? _____ years

4. What is highest education level you have completed? None

Primary

Secondary

Higher education/ University

Don't know

Refused

Other, specify

5. Marital status (tick one): Married (monogamous)

Married (polygamous)

Single

Divorced/Separated

Widowed

Refused

Cohabiting

Other, specify

6. Employment (tick one): Salaried job

Self-employed

- Housewife
- Unemployed
- Casual laborer
- Refused
- Other, specify

7. Household income per month (tick one): None
- < 5000 Ksh
- 5001 – 10000 Ksh
- 10001 – 15,000 Ksh
- >15,000 Ksh
- Don't know
- Refused

B: SEXUAL HISTORY

8. How old were you when you first had vaginal intercourse? ___ ___ Don't know
 refused never Other, specify
9. How many sex partners have you ever had? ___ ___ Don't know refused
 Other, specify _____
10. How many different sex partners did you have in the last year? ___ ___
 Don't know refused Other, specify
11. Have you had sex in the last month? (If no, don't know, refused, skip to 14)
 Yes No Don't know Refused Other, specify
12. How often have you used condoms during sex in the last month?
 Always (100%) Most of the time (75-99%)
 Half of the time (50%-74%) Sometimes (25-49%)

 Rarely (1-25%) Never (0%)
 Refused
 Don't know Other, specify
13. Do you think you may be currently pregnant? Yes No Don't know
 Refused Other, specify

C: CERVICAL CANCER SCREENING HISTORY

14. Have you been previously screened for cervical cancer? Yes No

Don't know Refused Other, specify

If no, don't know or refused skip to 20.

15. What cervical cancer screening test did you undergo most recently (tick one)?

Pap smear

Visual inspection with Acetic acid

HPV

Other, specify: _____

Don't know

Refused

16. Where was the most recent test performed?

Coptic

KNH

Refused

Don't know

Other, specify

17. When was the test performed?

Don't know

18. What were the results?

Normal

Abnormal, specify _____

Don't know

Refused

19. Did you receive any cervical treatment or surgery as a result of this test?

Yes No Don't know Refused Other, specify _____

If yes, specify _____

20. Was Pap Smear performed today? Yes No
 If no, explain Refused Don't know
21. Comments.

PAP SMEAR

Study ID Number _____

Interviewer Number _____

Date of interview (day/month/year) ____/____/____

Visit (tick one) Initial Visit Repeat Month 6 FU Month 12 FU Month 18 FU Month 24 Other, specify _____

A: PHYSICAL EXAMINATION

1. Temperature _____°C
2. Is there inguinal node enlargement Yes No
3. Is there abdominal tenderness Yes No
4. Are there any abdominal masses Yes No
5. Were there any other abnormalities found on general physical exam Yes No
If yes, specify _____

B: PELVIC EXAMINATION

6. What was found on the external genital exam (tick all that apply)?

<input type="checkbox"/> Abnormal visible discharge at introitus	<input type="checkbox"/> Ulcers
<input type="checkbox"/> Excoriations	<input type="checkbox"/> Vesicles
<input type="checkbox"/> Oedema	<input type="checkbox"/> Papules
<input type="checkbox"/> Sores	<input type="checkbox"/> Normal
<input type="checkbox"/> Other (Specify) _____	
7. Were there any perineal warts on external genital exam? Yes No
If yes, specify
 - a. Size _____ mm
 - b. Number of warts _____
 - c. Location of warts _____
8. Did the cervix appear abnormal on gross pelvic exam? Yes No
If yes, tick all that apply

<input type="checkbox"/> Abnormal discharge	<input type="checkbox"/> Warts
<input type="checkbox"/> Bleeds easily on touch	<input type="checkbox"/> Cervicitis
<input type="checkbox"/> Visible lesion	<input type="checkbox"/> Condylomata

- Bloody discharge Ulcers
- Fungating mass
- Leukoplakia Cervical polyp
- Bartholian cysts Blisters
- Overt cervical cancer Other abnormality, specify

9. Did you palpate the uterus? Yes No
If yes, then specify

- a. Estimated uterine size ___ . ___ cm
- b. Was the uterus tender? Yes No
- c. Were there possible fibroids? Yes No

10. Was there any adnexal tenderness? Yes No
If yes, specify location Right Left Both

C: Pap

11. Were you able to take an adequate Pap smear? Yes No

- a. If no, specify why
- Patient discomfort
- Excessive bleeding
- Excessive discharge or inflammation
- Other, specify

D: DIAGNOSIS

12. Normal exam Yes No
13. Candidiasis Yes No
14. Cervicitis Yes No
15. Pelvic inflammatory disease Yes No
16. Vulval warts Yes No
17. Vaginal warts Yes No
18. Genital ulcerations Yes No
19. Lower genital tract infection Yes No
20. Other Yes No
If others, specify

E: OTHER

21. Did you give any treatment to the participant? Yes No
If yes, specify (treatment)

22. Did you give referral to the participant? Yes No
If yes, specify (diagnosis and referral institution)

23. Comment.

COLPOSCOPY

Study ID Number ___ ___ ___

Interviewer Number ___ ___

Date of interview (day/month/year) ___ ___ / ___ ___ / ___ ___ ___ ___

Visit (tick one) Initial Visit Month 6 FU Month 12 FU Month 18 FU Month 24 FU LSIL FU CIN 1 FU Other, specify _____

A: PAP SMEAR DIAGNOSIS

1. What date was the Pap smear performed?
2. What was the Pap smear diagnosis (tick all that apply)?
 - No dysplasia (NIL)
 - ASCUS
 - LSIL (CIN 1)
 - HSIL (CIN 2 & 3)
 - ASC-H
 - Invasive carcinoma
 - ACG (Atypical Glandular Cells)
 - Cervicitis
 - Yeast infection
 - Indeterminate/insufficient sample
 - Other, specify _____
 - Unknown, specify reason _____
3. Is cervical biopsy with colposcopy indicated based on Pap smear cytology?

Yes No

If no, [skip to Q24 and](#) fill exit form where necessary

If yes, and colposcopy had been done previously, skip to Q24.

If yes and colposcopy had not been done previously, do a pregnancy test

4. Result of pregnancy test: Pregnant Not Pregnant

(If pregnant fill exit form, If not pregnant refer for colposcopy)

Not pregnant

B: PHYSICAL EXAMINATION

5. Temperature ___ °C

6. Is there inguinal node enlargement Yes No

7. Is there abdominal tenderness Yes No

8. Are there any abdominal masses Yes No

9. Were there any other abnormalities found on general physical exam Yes No
If yes, specify

C: PELVIC EXAMINATION

10. What was found on the external genital exam (tick all that apply)?

Abnormal visible discharge at introitus Ulcers

Excoriations Vesicles

Oedema Papules

Sores Normal

11. Were there any perennial warts on external genital exam? Yes No
If yes, specify

d. Size ___ mm

e. Number of warts ___

f. Location of warts _____

12. Did the cervix appear abnormal on gross pelvic exam? Yes No
If yes, tick all that apply

Abnormal discharge Warts

Bleeds easily on touch Cervicitis

Visible lesion Condylomata

Bloody discharge Ulcers

Fungating mass

Leukoplakia Cervical polyp

Blisters

Overt cervical cancer

13. Did you palpate the uterus? Yes No
If yes, the specify

a. Estimated uterine size ___ . ___ cm

b. Was the uterus tender? Yes No

c. Were there possible fibroids? Yes No

d. Was there any adnexal tenderness? Yes No

d(i) If yes, specify location Right Left Both

D: COLPOSCOPIC BIOPSY

15. Did you see the entire squamocolumnar junction (SCJ)? Yes No

16. Was it a satisfactory colposcopy? Yes No

If no, specify

17. Were there colposcopic findings within the transformation zone? Yes No
If yes, specify (tick all that apply)

Flat acetowhite epithelium

Micropapillary or microconvoluted acetowhite epithelium

Leukoplakia

Punctation

Mosaic

Atypical vessels

Iodine-negative epithelium

Lesion extended into endocervix

18. Draw SCJ (acetowhite, punctation, mosaics, atypical vessels, and other lesions):

19. Were there any other colposcopic findings? Yes No

If yes, specify (tick all that apply)

Mucosal bleeding easily induced

Purulent cervicitis

Opaque discharge

Yellow discharge

Other, specify:

20. Were there colposcopic findings consistent with invasive carcinoma?
Yes No

E: COLPOSCOPY DIAGNOSIS

21. Is patient eligible for cryotherapy if necessary? Yes No
If no, indicate reason (tick all that apply)

- Lesion >75% of cervix
- Lesion is larger than cryoprobe tip
- Lesion suspicious for cancer
- Polyp or anatomic defect preventing access to cervix
- Previous treatment with cryotherapy in this study
- Other, specify

22. What was your diagnosis based on colposcopy examination (tick all that apply)?

- Normal colposcopic findings
- Unsatisfactory, specify:
- Inflammation/infection, specify
- Leukoplakia
- Condyloma
- LSIL (CIN 1)
- HSIL (CIN 2 & 3)
- Invasive cancer
- Other, specify:

D: SPECIMEN COLLECTION

23. Was a biopsy taken? Yes No
- a. If yes, how many biopsies were taken ___
 - b. Draw: (mark site(s) with an 'X' on colposcopy drawing)
 - c. If no, specify why biopsy was not taken

E: Treatment

24. Was any treatment given to the patient? Yes No
If yes, specify treatment
25. Was a referral given to the patient? Yes No
If yes, specify diagnosis, and the referral institution

26. Comments:

RANDOMIZATION

Study ID Number _____ Hope ID Number _____

Randomization Number __ Interviewer Number _

Date of interview (day/month/year) ____/____/____

A: CURRENT MEDICAL HISTORY

1. Do you have pain when passing urine? Yes No Other, specify Refused Don't know
2. Do you have any lower abdominal pain? Yes No Other, specify Refused Don't know
3. Do you have any abnormal vaginal discharge? Yes No Other, specify Refused Don't know
4. Have you noticed any growths around your vagina? Yes No Don't know Other, specify Refused Don't know

B: REPRODUCTIVE HEALTH

5. How old were you when you had your first menstrual period? ____ Don't know refused Other, specify
6. Date of last menstrual period (day/month/year) ____/____/____ Don't know refused Other, specify
7. Do you have history of abnormal vaginal bleeding Yes No Don't know Refused Other, specify

If yes specify the type of bleeding

- Irregular
- Heavy
- Menorrhagia Other, specify
- Don't know
- Refused

8. Have you ever used any form of family planning method? Yes No
Don't know

Refused

Other, specify _____

If yes, specify (tick all that apply)

Injectable IUCD Natural

Condoms OCP Norplant/ Implant

BTL others, specify _____

9. Are you using any form of family planning method now?

Yes No Don't know Refused

Other, specify _____

If yes, specify (tick all that apply)

Injectable IUCD Natural

Condoms OCP Norplant/ Implant

BTL others, specify _____

10. How many times have you been pregnant? ___ ___ Refused Don't know
Other

11. How many times have you had live births? ___ ___ Refused Don't know
Other

12. How many abortions, miscarriages, and/or stillbirths have you had? ___ ___
Refused Don't know Other

13. Have you ever been admitted to the hospital with a gynecological problem?
Yes No Don't know Refused Other, specify

14. Have you ever had abdominal surgery? Yes No Refused
Don't know Other, specify

15. Have you ever had vaginal surgery? Yes No
Refused Don't know Other, specify

16. Do you currently smoke cigarettes Yes No
Refused Don't know Other, specify

C: HIV HISTORY

17. When were you diagnosed as having HIV? (day/month/year) ___ ___ / ___
___ / _____

18.1. How was HIV detected? (tick only one)

- At the occasion of a VCT: Yes
- During pre-natal check-up: Yes
- Because of a sickness, specify: _____ Yes
- Other, specify: _____
- Don't know:
- Refused:

19. Are you currently on antiretroviral medications? Yes No
 Don't know
 Refused
 Other, specify _____

If yes,

a) specify current medications: _____

b) original start date: ____ / ____ / ____ Don't know

c) Do you know why you were started on antiretroviral medication?

- because of sickness, specify: _____ Yes
- because of low CD4 count: Yes
- because of high viral load: Yes
- Other, specify: _____
- Don't know:
- Refused:

20. Comment

4. Is patient indicated for LEEP or cryotherapy treatment based on histopathology?

Yes No

If no, continue to 14

5. Is patient's lesion amenable to cryotherapy? Yes No (If yes, go to question 6)

5(i) If no, indicate reason (tick all that apply)

- Lesion >75% of cervix
- Lesion is larger than cryoprobe tip
- Invasive cervical cancer on histology
- Lesion suspicious for cancer
- Polyp or anatomic defect preventing access to cervix
- Patient declines procedure
- Other, specify

5(ii) If no, is patient's lesions amenable to LEEP? Yes No

If yes, skip to 7

If no, skip to 14

6. Was patient randomized to LEEP or cryotherapy today? Yes No
If yes, to which treatment? LEEP Cryotherapy

If no, why?

- Patient refuses
- Patient not eligible for treatment, specify reason
- Other, specify

C: TREATMENT

7. Did you perform LEEP or cryotherapy today? (tick one)
LEEP Cryotherapy Neither, explain

(If neither, explain, skip to Q14)

- 8. Did you visualize the full extent of lesion? Yes No
- 9. Was the squamocolumnar junction fully visualized? Yes No
- 10. Draw position of lesion and treatment performed:
- 11. Did the patient experience any pain during the procedure? Yes No
- 12. Was analgesia provided? Yes No
- 13. Were there any complications? Yes No

If yes, specify

14. Was any antibiotics given? Yes No

If yes, specify

D: OTHER

15. Was the patient referred for further treatment at another institution? Yes

No

If yes, specify institution and reason

16. Was treatment or a referral given for something other than cervical disease? Yes No

If yes, specify (give diagnosis and treatment)

17. Comments

ADDRESS AND INTAKE

Study ID number _ _ _ _

Interviewer number _ _ _ _

Date of interview (day/month/year) _ _ _ / _ _ _ / _ _ _ _ _ _

Format(tick one)

 New Update (fill only updated info)

A. PERSONAL INFORMATION

1. What is your name?
 - a. First name
 - b. Middle name
 - c. Last name
2. How are you called in your home area?
3. What is the current physical location where you live?
 - a. District
 - b. City
 - c. Village
 - d. Estate
 - e. Plot number
 - f. Door number
 - g. Road name
4. Public Transportation to the house:
 - 4a. Type: F Bus F Boda boda F Matatu F Taxi F Other (specify)
 - 4b. Route number:
 - 4c. Stage Name
 - 4d. General Name of the Area
5. Walking directions to house from the stage?
6. Landmarks that aide in locating the household: (Names of schools, churches, businesses etc.)
7. Can you be reached by phone? Yes No
If yes,
 - 7 (i)What is the phone number 1?
 - 7 (ii)What is the phone number 2?
 - 7iii. Who carries the phone (tick one) self other, specify

B. SIGNIFICANT CONTACT

8. Is there another person who is ware of your HIV status that we can contact through phone if we are unable to reach you directly?
If no, skip to 9.
9. What is the name of this contact person?
a. First name
b. Middle name
c. Last name
10. What is the relationship of this person to you?
11. (a)What is the phone number 1?
(b) What is the phone number 2?

C. UPCOUNTRY INFORMATION

12. Do you have an upcountry home? Yes No
If no, skip to 19
13. What is the physical location of your upcountry home?
a. District
b. City
c. Village
d. Estate
e. Plot number
f. Door number
14. Specific directions to residence
15. Is there person located in your upcountry home whom we can contact if we are unable to reach you directly? Yes No
If no, skip to 19
16. What is the name of this contact person?
a. First name
b. Middle name
c. Last name
17. What is the relationship of this person to you?
18. Does this contact have a phone number? Yes No
If yes, specify: phone number 1, phone number 2

D. OTHER

Comment:

SHEDDING

Study ID Number ___ ___ ___

Interviewer Number ___ ___

Date of interview (day/month/year) ___ ___ / ___ ___ / ___ ___ ___ ___

Visit (tick one) Week 1 FU Week 2 FU Week 3 FU Other, specify _____

A: MEDICAL HISTORY

1. Has the patient experienced any lower abdominal pain since the last visit?

Yes No

If yes, indicate:

a. Duration ___ ___ days

b. Severity (scale from 1 to 5; 5 being most severe) ___ ___

2. Has the patient experienced any vaginal bleeding since the last visit?

Yes No

If yes, indicate:

a. Duration ___ ___ days

b. Volume

 Stain pants Requires sanitary pad Other, specify

3. Has the patient experienced any vaginal discharge since the last visit?

Yes No

If yes, indicate:

c. Duration ___ ___ days

d. Color

 Yellow Brown White Clear

Other, specify

e. Smell

Malodorous

No odor

Other, specify

f. Volume

Stain pants

Requires sanitary pad

Other, specify

4. Has the patient experienced any fever after the last visit?

Yes No

If yes, specify duration ___ days

5. Did the patient seek medical care for these or other complaints?

Yes No

If yes, specify what complaint prompted the participant to seek care:

Abdominal pain

Fever

Vaginal Bleeding

Vaginal discharge

a. Other, specify

If yes, specify where the participant sought care:

Study clinic

Hope Center

Coptic Hospital

KNH

b. Other, specify

6. Was the participant's condition possibly due to a study procedure? (Ask if any of questions 1-4 is yes)

Yes No

If any question 1-4 is Yes and the condition is not expected as per study protocol, or is more severe than expected, fill an AE form.

7. Date of last menstrual cycle (dd/mm/yyyy)

8. Have you ever had vaginal sex since treatment? Yes No Refused

Other (specify) _____

8 (a). If yes, how many times? ___

8(b) If you have had vaginal sex since treatment, how often did you use condoms during sex?

Always (100%) Most of the time (75-99%)
 Half of the time (50%-74%) Sometimes (25-49%)
 Rarely (1-25%) Never (0%)
 Refused Don't know Other, specify _____

9. Do you think you may be currently pregnant? Yes No

10. Are you currently on antiretroviral medications?

Yes Refused Other, specify

No Don't know

If No, Refused, or Don't Know, skip to Q14

If yes, specify current medications and original start date.

- d4t, 3tc, nvp
- d4t, 3tc, efv
- azt, 3tc, nvp
- azt, 3tc, efv
- Other specify

Original ARV start date (dd/mm/yyyy)

11. During the last 7 days, how many antiretroviral pills did the patient MISS taking?

12. During the last 30 days, how many antiretroviral pills did the patient MISS taking?

(If Q11 and Q12 patient did not MISS taking any pills skip to Q14)

13. If the patient missed any doses, please specify reasons (check all that apply)

- Toxicity/ side effect
- Share with others
- Forgot
 - c. Felt better
- Too ill
- Stigma, disclosure or privacy issues
- Drug out of stock
- Patient lost or ran out of pills
- Delivery /travel problem
- Inability to pay
- Alcohol
- Depression
 - d. Other specify

B: DIAGNOSIS

14. Is there any new infection that was related to the procedure since the last visit?

Yes No

If yes, specify and fill an Adverse Event questionnaire:

15. Were there any new complications diagnosed today related to the treatment?

Yes No

If yes, specify and fill an Adverse Event questionnaire:

C: TREATMENT

16. Was any treatment provided today? Yes No

If yes, specify

17. Was the patient referred for further cervical treatment at another institution?

Yes No

17 (i) If yes, specify institution

17(ii) If yes, specify reason for referral

D: SPECIMEN COLLECTION

18. Did you collect cervical HIV swab? Yes No
 If no, specify reason:

19. Did you collect blood? Yes No
 If no, specify reason:

E: ACCEPTABILITY OF TREATMENT

20. Please complete the items listed below by placing a checkmark on the box next to each question that best indicates how the client feels about the treatment she received.

	Strongly Disagree	Disagree,	Neutral,	Agree	strongly agree
(i) I find this treatment an acceptable way of dealing with cervical lesions	<input type="checkbox"/>				
(ii) I would be willing to use this procedure if I were to develop more lesions	<input type="checkbox"/>				
(iii) I would recommend this procedure to someone with cervical lesions	<input type="checkbox"/>				
(iv) overall, I have a positive reaction to this treatment	<input type="checkbox"/>				

F: OTHER

22.

—

23.

24. Comments:

PATIENT CONTACT

Study ID Number ___ ___

Interviewer Number ___ ___

Date of contact (day/month/year) ___ ___/___ ___/___ ___

1. Date of last Hope clinic visit (day/month/year)
2. Date of last study clinic visit (day/ month/year)
3. Was patient or patient contact reached by phone or home visit?
 Phone Home visit Other, specify

4. Did you talk to the patient or patient contact?

- Yes, talked to patient (*Go to 6*)
- Yes, talked to patient contact (*Go to 5*)
- No (*Go to 4*)

5. If talked to patient contact, who was the source of information (tick one)

- Clinician / clinic staff Mother or Father Neighbor
- Spouse or Partner Employer Caregiver
- Family member / Relative Friend Treatment supporter
- Other (specify)

5a. Did the source of information have credible knowledge for whether the patient was alive or dead?

- Yes, credible and patient confirmed alive (*Go to 4*)
- Yes, credible and patient confirmed dead (*Go to 7 and complete Verbal Autopsy form*)
- Source did not know whether patient was dead or alive (*Go to 4*)
- Source not credible (*Go to 7 and complete Verbal Autopsy form*)

6. Reasons for missed study appointment (tick all that apply)

- N/A, did not reach patient or patient contact
- Unable attend because health problem
- No longer willing to be in study
- Family problems
- Wait time too long in clinic
- Client will go to faith healer
- Conflict with work
- Unwilling to disclose
- Financial problems
- No longer willing attend Hope Clinic
- Client moved or relocated
- Other (specify)

7. If talked to patient, did the patient wish to remain in the study?

- Yes (*Go to 7a*)
- No (*Go to 8 and complete Exit Form*)
- N/A, did not talk to patient (*Go to 9*)

7a. Did the patient schedule a study appointment? Yes No (*If Yes go to 7b, If No go to 7c*)

7b. If yes, date of scheduled appointment (DD/MM/YYYY)

7c. If no, why did the patient not schedule an appointment (tick all that apply)

- Unable to attend because of health problems
- Family problems
- Conflict with work
- Financial problems
- Client moved or relocated
- Client lives too far away
- Other (specify)

8. If the patient does not wish to return to the study, specify why (tick all that apply)

F Not willing to attend Hope Clinic

F Not willing to be in study

F Attend clinic closer to home

F Wait time too long

F Conflict with work

F Financial problems

F Unwilling to attend because of health problems

F Family problems

F Client will go to a faith healer

F Not willing to disclose HIV status

F Referred else where

F Other (specify)

F Unknown

9. Comments

VERBAL AUTOPSY

Study ID Number _____

Interviewer Number _____

Date of interview (day/month/year) ____/____/____

1. Age at death FF years

2. Date of death (DD/MM/YYYY) FF/FF/FFFF

3. Place of death

4. The information source for the cause of death was (tick all that apply)

Partner/spouse

Mother or Father

Friend

Hospital records/staff

Other family member/relative

Care giver

Neighbor

Unknown

Other (specify)

5. Was the deceased seeking other medical treatment (other than Hope Center) during the last 3 months before his/her death? Yes No Unknown *If yes go to*

5a, if no go to 6

5a. If yes, where specifically was the deceased receiving other medical care?

Name of facility: _____

5b. What type of care was the deceased receiving at the these other facilities? (Tick all that apply)

General medical care

TB

STD

HIV/AIDS

Malaria

Other infectious disease(s)

Other care (specify all types of care received: _____)

6. Respondent’s detailed account of the illness of the deceased:

7. Did a health care worker tell you the cause of death? F Yes F No F Unknown
If yes go to 7a, if no go to 8

7a. What did the health care worker say was the cause of death? _____

8. Did s/he have any operation for the illness? F Yes F No F Unknown
If yes go to 8a, if no go to 9

8a. How long before the death was the operation? _____ days

8b. On what part of the body was the operation?

F Abdomen F Chest F Head F Other(Specify _____)

9. Has spouse or other sexual partner(s) of the deceased died in the past 5 years?

F Yes F No F Unknown
If yes go to 9a, if no go to 10

9a. If yes, what is the believed cause(s) of death of the partner(s)

9a1. Partner 1: _____

9a2. Partner 2: _____

Injury/accident/suicide

10. Did s/he suffer from any injury or accident that led to her death? F Yes F No F Unknown

If yes go to 10a, if no go to 11

10a. What kind of injury or accident did the deceased suffer?

Road traffic accident Fall Drowning Poisoning

Burns Violence/assault

Other: _____

Unknown

10b. Was the injury or accident intentionally inflicted by someone else? Yes No Unknown

10c. Do you think that s/he committed suicide? Yes No Unknown
Skip to Q.12

11. Did s/he suffer from any animal/insect bite that led to her/his death? Yes No Unknown

11a. If yes, what type of animal/insect? _____

History of previously known medical conditions

12. Did the deceased suffer from any of the following conditions?

- a. High blood pressure Yes No Unknown
- b. Diabetes Yes No Unknown
- c. Asthma Yes No Unknown
- d. Epilepsy Yes No Unknown
- e. Malnutrition Yes No Unknown
- f. Cancer Yes No Unknown

f1. If yes, specify type of cancer or site: _____

g. Tuberculosis Yes No Unknown

h. Any other medically diagnosed illness? Yes No Unknown

h1. If yes, specify: _____

13. Signs, symptoms, and their severity during the last illness:

Symptom/ Signs	Symptom present?	If present, duration of symptom
a. Fever	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
b. Loss of weight	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
c. Diarrhea	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
d. Vomiting/associated abdominal pain	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
e. Constipation/associated abdominal pain	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
f. Cough	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
g. Cough followed by vomiting	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
h. Breathing trouble (chest indrawing/difficult/rapid/wheezing)	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
i. Neck stiffness	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
j. Unconscious episodes	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
k. Fits	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
l. Jerking of individual limbs	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
m. History of epileptic illness in earlier years	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
n. Paralysis of limbs	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
o. Rigid body stiffness, unable to open mouth	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
p. Red and sore eyes	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
q. Skin rash and itching	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
r. Herpes Zoster (at any time in life)	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
s. Abscesses/body sores	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
t. White patches on the inside of mouth and tongue	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
u. Oedema	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
v. Hair changes	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
w. Yellowing of eyes or passing of brown urine	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
x. Chest pain	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
y. Other (Specify:)	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown

Unexpected vaginal bleeding or discharge	No F Yes	≤2 weeks F >2 weeks F Unknown
Pelvic or vaginal pain	No F Yes	≤2 weeks F >2 weeks F Unknown

14. Records available in home, e.g., death certificate (extract findings):
15. Comment

EXITStudy ID Number: ____ ____ ____

Interviewer Number: _____

Date of visit (day/month/year) ____ ____ / ____ ____ / ____ ____ ____ ____

1. Is the patient exiting the study because she has completed the study protocol and qualifies as per study guidelines to exit the study at this time? Yes No *(If Yes, go to 9, If No go to 2)*

2. Date of last study visit (DD/MM/YYYY) HH/HH/HHH3. Date last seen by study staff (DD/MM/YYYY) HH/HH/HHH4. Has the patient accessed care at the Hope Clinic in the past year? Yes No5. Did you talk to the patient? Yes No *(If Yes go to 6, If No go to 5a)*

5a. If no, who was the source of information (tick one)

 Clinician / clinic staff Employer Treatment supporter Spouse or Partner Friend Other (specify) Family member / Relative Neighbor Mother or Father Caregiver

6. Has the patient transferred HIV care to another program? Yes No *(If Yes go to 6a, If No go to 7)*

6a. If yes, where is the patient transferring care to (tick one)

 Transferred to another non-Hope clinic, specify Transferred to Industrial area clinic Transferred to Maseno clinic

7. Has Hope Clinic asked the patient to Exit or leave the program? Yes No (*If Yes go to 7a, If No go to 8*)

7a. Reasons for being Exited from the program (tick all that apply)

Client has not returned to clinic for 1 year

Poor Adherence

Poor Clinic Attendance

Not willing to disclose HIV status

Referred else where

Tested Negative

Other (specify)

Unknown

8. Has the patient asked to be Unenrolled from the study? Yes No (*If Yes go to 8a, If No go to 9*)

8a. Reasons patient is asking to Unenroll from the study (tick all that apply)

Not willing to attend

Waiting time too long

Conflict with work

Financial problems

Unwilling to attend because of health problems

Family Problems

Client will go to a faith healer

Not willing to disclose HIV status

Referred else where

Other (specify)

Unknown

9. At what point was the patient exited? After Pap smear After Biopsy Mortality Withdrawn from study Other (If other specify)

10. Comment

9. Any microbiological findings? Yes No
 If “Yes”, specify findings:

- Lactobacilli
- Mixed flora
- Bacterial vaginosis
- Candida
- Trichomonas vaginalis
- Actinomyces
- Schistosoma
- Herpes simplex
- Other, specify.....

10. Any reactive changes observed? Yes No
 If “Yes”, specify findings

- Metaplasia
- Inflammatory changes
- Follicular cervicitis
- Parakeratosis
- Atrophy
- Other reactive changes
- If “Yes”, specify.....

11. Squamous epithelium: Normal Abnormal
 If “Yes”, specify findings

- ASC-US Yes No
- ASC-H Yes No
- Low grade SIL Yes No
- With koilocytosis Yes No

- High grade SIL Yes No
- HSIL invasion not ruled out Yes No
- Squamous cell carcinoma Yes No
- Others, specify _____

12. Glandular epithelium: Normal
abnormal

If “Yes”, specify findings

- Abnormal presence of endometrial cells Yes No
- Atypical glandular cells, pref. neoplastic Yes No
- Adenocarcinoma in situ Yes No
- Adenocarcinoma Yes No
 - Endometrial Yes No
 - Endocervical Yes No

Others, specify _____

13. Other

remarks.....

Interviewer number of pathologist

Date: _____

COLPOSCOPIC BIOPSY HISTOLOGY REPORT

First read Re-read 1 Consensus reading Tie-break

Study ID Number _____

Hope ID Number _____

Test/ Visit code _____

Visit (tick one) Initial Visit Month 6 FU Month 12 FU Month 18 FU
 Month 24 FU Other, specify

Interviewer number --- ---- Patient Age _____

1. Date sample was collected in clinic (day/month/year) ____/____/____
2. Time sample was collected in clinic
3. Date sample was received in laboratory (day/month/year) ____/____/____
4. Time sample was received in laboratory
5. When was the biopsy processed? (day/month/year) ____/____/____
6. When was the biopsy reported? (day/month/year) ____/____/____
7. Was the amount of sample adequate for reading? F Yes F No Other, specify
8. What was the histology result of the cervical biopsy (tick one)?
 - No dysplasia (NIL)
 - CIN 1
 - CIN 2
 - CIN 3
 - Invasive carcinoma
 - Indeterminate/
 - CIS
 - Other, specify
9. Was there evidence of cervicitis? F Yes F No Other, specify
10. Who read the biopsy and gave this histology result?
Pathologist Interviewer number _____

11. Comments:

ENDOCERVICAL CURETTAGE HISTOLOGY REPORT

First read Re-read 1 Consensus reading Tie-break

Study ID Number _____

Hope ID Number _____

Test/ Visit code _____

Visit (tick one) Initial Visit Month 6 FU Month 12 FU Month 18 FU

Month 24 FU Other, specify

Interviewer number --- ----Patient Age _____

1. Date sample was collected in clinic (day/month/year) ____/____/____

2. Time sample was collected in clinic

3. Date sample was received in laboratory (day/month/year) ____/____/____

4. Time sample was received in laboratory

5. When was the biopsy processed? (day/month/year) ____/____/____

6. When was the biopsy reported? (day/month/year) ____/____/____

7. Was the amount of sample adequate for reading? F Yes F No Other, specify

8. What was the histology result of the Endocervical Curettage biopsy (tick one)?

No dysplasia (NIL)

CIN 1

CIN 2

CIN 3

Invasive carcinoma

Indeterminate/

CIS

Other, specify

9. Was there evidence of cervicitis? F Yes F No Other, specify

10. Who read the biopsy and gave this histology result?

Pathologist Interviewer number _____

11. Comments:

--

LEEP BIOPSY HISTOLOGY REPORT

First read Re-read 1 Consensus reading Tie-break

Study ID Number _____

Hope ID Number _____

Test/ Visit code _____

Visit (tick one) Initial Visit Month 6 FU Month 12 FU Month 18 FU
 Month 24 FU Other, specify

Patient Age _____ Interviewer number --- ----

1. Date sample was collected in clinic (day/month/year) ____/____/____

2. Time sample was collected in clinic

3. Notes:

4. Date sample was received in laboratory (day/month/year) ____/____/____

5. Time sample was received in laboratory

6. When was the LEEP specimen processed? (day/month/year) ____/____/____

7. When was the LEEP specimen reported? (day/month/year) ____/____/____

8. Was the amount of sample adequate for reading? F Yes F No Other, specify

9. What was the histology result of the LEEP cervical biopsy (tick one)?

No dysplasia (NIL)

CIN 1

CIN 2

CIN 3

Invasive carcinoma

Indeterminate/

CIS

Other, specify

10. Was there evidence of cervicitis? F Yes F No Other, specify

11. Who read the biopsy and gave this histology result?

Pathologist Interviewer number ____

12. Comments:

--

CD4 REPORT

Study ID Number

Hope ID Number

Visit (tick one) Randomization/LEEP Visit Month 6FU Month 12FU Month 18FU Month 24FU Other, specify

Patient Age

Interviewer number

1. Date sample was collected in clinic (day/month/year) / /
2. Time sample was collected in clinic
3. Date sample was received in laboratory (day/month/year) / /
4. Time sample was received in laboratory
5. When was the CD4 count run? (day/month/year) / /
6. What was the result of the CD4 count?
7. Who ran the CD4 count and gave this CD4 result? Coptic lab F Other, specify

8. Comments:

FOLLOW-UP FORM

Study ID Number _____ Hope ID Number _____ Interviewer Number _____

Date of interview (day/month/year) _____/_____/_____

Visit (tick one) Month 6 FU Month 12 FU Month 18 FU

Month 24 FU Other, specify _____

A: MEDICAL HISTORY

1. Did you consult a doctor or clinical officer between today and your previous study visit?

- Yes
- No (go to question 2)
- Don't know (go to question 2)
- Refused (go to question 2)

If yes:

1.1. How many consultations?

- _____ Don't know (go to question 2)
- _____ Refused (go to question 2)

1.2. Why did you consult?

1.2.1. Visit 1:

Reason for consultation		
Routine check-up visit not study related	<input type="checkbox"/>	yes
Sickness.....	<input type="checkbox"/>	yes
Other, specify	<input type="checkbox"/>	yes
Do not recall the reason	<input type="checkbox"/>	yes
Refused	<input type="checkbox"/>	yes

1.2.2. Visit 2:

Reason for consultation		
Routine check-up visit not study related	<input type="checkbox"/>	yes
Sickness.....	<input type="checkbox"/>	yes
Other, specify	<input type="checkbox"/>	yes
Do not recall the reason	<input type="checkbox"/>	yes
Refused	<input type="checkbox"/>	yes

1.2.3. Visit 3:

Reason for consultation		
Routine check-up visit not study related	<input type="checkbox"/>	yes
Sickness.....	<input type="checkbox"/>	yes
Other, specify	<input type="checkbox"/>	yes
Do not recall the reason	<input type="checkbox"/>	yes
Refused	<input type="checkbox"/>	yes

1.2.4 Visit 4

Reason for consultation		
Routine check-up visit not study related	<input type="checkbox"/>	yes
Sickness.....	<input type="checkbox"/>	yes
Other, specify	<input type="checkbox"/>	yes
Do not recall the reason	<input type="checkbox"/>	yes
Refused	<input type="checkbox"/>	yes

1.2.5 Visit 5

Reason for consultation		
Routine check-up visit not study related	<input type="checkbox"/>	yes
Sickness.....	<input type="checkbox"/>	yes
Other, specify	<input type="checkbox"/>	yes
Do not recall the reason	<input type="checkbox"/>	yes
Refused	<input type="checkbox"/>	yes

1.2.6 Visit 6

Reason for consultation		
Routine check-up visit not study related	<input type="checkbox"/>	yes
Sickness.....	<input type="checkbox"/>	yes
Other, specify	<input type="checkbox"/>	yes
Do not recall the reason	<input type="checkbox"/>	yes
Refused	<input type="checkbox"/>	yes

(For those previously not on ARVs- please refer to enrollment form)

2. Have you initiated anti-retroviral medication since the last visit?

- Yes
- No
- Don't know
- Refused

If yes:

2.1. Which date did you initiate ARV's --/ --/ ----

2.2. Specify current regimen

3. Are you still on ARVs? (For those previously on ARVs-please refer to enrollment form)

- Yes
 - No
 - NA
- (If NA go to Q4)

3.1 If no, reasons for not being on ARVs

- Poor adherence
- Side effect
- Stigma
- Concurrent illness
- Other (specify)

3.2 If yes, has your anti-retroviral medication changed since last cervical treatment visit?

- Yes
- No (go to question 4)
- Don't know (go to question 4)
- Refused (go to question 4)

If yes:

3.2.1. On which date was it changed: ____/____/____

3.2.2. What was the reason for changing your antiretroviral medication? (Check all that apply)

- a) Because of low CD4 counts: Yes
- b) Because of high viral load: Yes
- c) Because of clinical symptoms: Yes
- f) Other, specify: _____ Yes
- e) Don't know: Yes
- d) Refused: Yes

3.2.3 Specify current medications: _____

4. Did you receive any cervical treatment other than what we offered?

- Yes
- No (go to part B)
- Don't know (go to part B)
- Refused (go to part B)

If yes:

4.1. On which date was this: ____/____/____

4.2. Which treatment did you receive?

- a) Cryotherapy: Yes
- b) LEEP: Yes
- c) Cold knife exconisation: Yes
- d) Hysterectomy: Yes
- f) Other, specify: _____ Yes
- e) Don't know: Yes
- d) Refused: Yes

B. SEXUAL HISTORY

5. Have you had sex in the last 6 months?

- Yes
- No (go to Q6)
- Don't know (go to Q6)
- Refused (go to Q6)

If yes:

5.1. With your regular partner? Yes
 No (go to Q5.2)

If yes,

5.1.1. How often have you used condoms during sex with your regular partner in the last 6 months?

- Always (100%)
- Most of the time (75-99%)

- Half of the time (50%-74%)
- Sometimes (25-49%)
- Rarely (1-25%)
- Never (0%)
- Don't know
- Refused to answer

5.1.2. Do you suspect that your partner has had other sexual partners during the last 6 months?

- Yes
- No
- Don't know
- Refused

5.2. Did you have sex with any other partner(s) than your regular partner during the last 6 months?

- Yes
- No (Go to Q6)
- Don't know (Go to Q6)
- Refused (Go to Q6)

If yes:

5.2.1. How many other different sexual partners besides your regular partner did you have in the last 6 months? _____

- Don't know
- Refused

5.2.2. How often have you used condoms during sex with these other partner(s) in the last 6 months?

- Always (100%)
- Most of the time (75-99%)
- Half of the time (50%-74%)
- Sometimes (25-49%)
- Rarely (1-25%)
- Never (0%)
- Don't know

6. Refused to answer

7. Comment

ADVERSE EVENTS

Study ID Number _____

Interviewer Number _____

Date of visit (day/month/year) _____/_____/_____

- Visit (tick one) Shedding1 Shedding2 Shedding3
- Month 6 FU Month 12 FU Month 18 FU Month 24 FU
- Other, specify _____

	Adverse Event	Start Date (DD/MM/YYYY)	End Date (DD/MM/YYYY)	Severity 1= Mild 2= Moderate 3= Severe 4= Life Threatening	Relationship to Study procedures 1= Related 2= Notrelated	Outcome 1= Resolved without sequelae 2= Resolved with sequelae 3= Ongoing 4= Death 5= Unknown	Serious 0= No 1= Yes**
1							
2							
3							
4							

Comments: _____

**** If Serious Adverse Event, contact a PI immediately after filling this form.****

PI Contacts: Dr. Michael Chung – 020-271-2947, 0722-579-963, Dr. Nelly Mugo – 020-273-6744

UANDIKISHAJI

Study ID Number _____ Hope ID Number _____ Interviewer Number _____

Date of interview (day/month/year) _____/_____/_____

Ulikubali kuhifadhi sampuli : Ndio La Sijui Nyingine, eleza

A: SOCIODEMOGRAPHICS

1. Tarehe ya kuzaliwa (siku/mwezi/mwaka) _____/_____/_____

2. Miaka _____ years

3. Ulikamilisha miaka ngapi ya elimu? _____ years

4. Elimu ya juu zaidi uliyokamilisha? Hakuna
Msingi

Sekondari

Elimu ya juu/ Chuo Kikuu

Sijui

Amekataa

Nyingine, eleza

5. Hali ya ndoa (jibu moja) Ndoa (mke mmoja)

Ndoa (wake wengi)

Pekee

Talaka/Tenganishwa

Mjane

Amekataa

Kuishi pamoja

Nyingine eleza

6. Ajira (jibu moga): Kazi inayokupa mshahara

Kujiajiri

- Mama wa nyumbani
- Hujaajiriwa
- Mfanyikazi wa kawaida
- Amekataa
- Nyingine, eleza
7. Kipato cha kaya kwa mwezi (jibu moja): Hakuna
- < 5000 Ksh
- 5001 – 10000 Ksh
- 10001 – 15,000 Ksh
- >15,000 Ksh
- Sijui
- Amekataa

B: HISTORIA YA KUFANYA MAPENZI

8. Ulikuwa na miaka mingapi ulipofanya ngono ya uke mara ya kwanza? ___
Sijui Amekataa Hujawahi Nyingine, eleza
9. Umewahi kuwa na wapenzi wangapi wa ngono? ___
Sijui Amekataa
Nyingine, eleza _____
10. Umekuwa na wapenzi wangapi tofauti wa ngono mwaka wa mwisho? ___
___ Sijui Amekataa Nyingine, eleza
11. Umefanya ngono katika mwezi wa mwisho? (If no, don't know, refused, skip to 14) Ndio La Sijui Amekataa Nyingine, eleza
12. Ni mara ngapi umetumia mpira wakati wa ngono katika mwezi uliopita?
Kila wakati (100%) Wengi wa wakati huo (75-99%)
Nusu ya mudu (50%-74%) Wakati mwingine (25-49%)
Mara chache (1-25%) Kamwe (0%)
Amekataa Sijui
Nyingine, eleza
13. Je, unafikiri unaweza kuwa ma mimba sasa? Ndio La Sijui
Amekataa Nyingine, eleza

C: SARATANI YA UZAZI HISTORIA YA UCHUNGUZI

14. Hapo awali, umewahi kupimwa saratani ya uzazi? Ndio La
Sijui Alikataa Nyingine, eleza

If no, don't know or refused skip to 20.

15. Uchunguzi gani wa saratani ya uzazi uliofanyiwa hivi karibuni? (jibu moja)

- Pap smear
- Ukaguzi na asetiki aside
- HPV
- Nyingine, eleza : _____
- Sijui
- Amekataa

16. Uchunguzi huu wa hivi karibuni ulifanyiwa wapi?

- Coptic
- KNH
- Amekataa Sijui
- Nyingine, eleza

17. Uchunguzi huu ulifanyiwa wapi?

sijui

18. Majibu yalikuwa nini?

- Kawaida
- Usiokuwa ya kawaida, eleza _____
- Sijui
- Amekataa

19. Ulipokea matibabu yoyote ya kizazi au upasuaji kwasababu ya majibu ya hii uchunguzi?

Ndio La Sijui Amekataa Nyingine, eleza _____

Kama ni ndio, eleza _____

20. Pap Smear ilifanywa leo? Ndio La Amekataa Sijui

Kama ni la, fafana

21. Maoni

UBAHATISHAJI

Study ID Number ___ Hope ID Number _____

Randomization Number ___ Interviewer Number _

Date of interview (day/month/year) ___/___/_____

A: HISTORIA YA SASA YA MATIBABU

1. Je, unauchungu unapopitisha mkojo? Ndio La Nyingine, eleza Amekataa Sijui

2. Je, una uchungu sehemu ya chini ya tumbo? Ndio La Nyingine, eleza Amekataa Sijui

3. Je, unatokwa na uchafu yasiyo ya kawaida katika uzazi wako wa kike? Ndio La Nyingine, eleza Amekataa Sijui

4. Je, umeona uvimbe yoyote katika uzazi wako wa kike? Ndio La Nyingine, eleza Amekataa Sijui

B: AFYA YA UZAZI

5. Ulikuwa na miaka mingapi ulipopata damu ya mwezi? ___ Amekataa Sijui Nyingine, eleza

6. Tarehe ya mwisho ya kupata damu yako ya mwezi ? (dd/mm/yy) ___/___/___/___ Sijui Amekataa Nyingine eleza

7. Je, una historia ya kutokwa na damu isiyo ya kawaida sehemu ya uke? Ndio La Sijui Amekataa Nyingine, eleza

Kama ndio, eleza aina ya damu

Isiyo mara kwa mara

Nzito

Kutokwa na damu nzito sana wakati wako wa mwezi

Nyingine, eleza

Sijui

Amekataa

8. Je, umewahi kutumia njia yoyote ya kupanga uzazi? Ndio La
Sijui

Amekataa
Nyingine, eleza _____

Kama ndio, eleza (chagua yote yanayotumika)

Sindano IUCD Asili

Mpira Tembe Norplant/Implant

BTL Nyingine, eleza _____

9. Je, sasa hivi unatumia njia yoyote ya kupanga uzazi?

ndio la Sijui Amekataa

Nyingine, eleza _____

Kama ndio, eleza (chagua yote yanayotumika)

Sindano IUCD Asili

Mpira Tembe Norplant/Implant

BTL Nyingine, eleza _____

10. Je, umekuwa na mimba mara ngapi? ___ Amekataa Sijui Nyingine

11. Je, umekuwa na watoto walioishi mara ngapi? ___ Amekataa Sijui
Nyingine

12. Je, umetoa mimba ngapi au kupoteza mimba au mtoto kufia kabla kuzaliwa ?
___ Amekataa Sijui Nyingine

13. Je, umewahi kulazwa hospitalini na tatizo yoyote ya gynecologia?

Ndio La Sijui Amekataa Nyingine, eleza

14. Je, umewahi kupasuliwa tumbo? Ndio La Amekataa
Sijui Nyingine, eleza

15. Je, umewahi kupasuliwa uke? Ndio La Amekataa
Sijui Nyingine, eleza

16. Sasa hivi, unavuta sigara? Ndio La
Amekataa Sijui Nyingine, eleza

C: HISTORIA YA HIV

17. Ulijulikana uko na HIV lini? (dd/mm/yyyy) ____ / ____ / ____

18.1. HIV ilitambuliwa aje? (jibu moja pekee)

- Ulipotembelea kituo cha VCT: Ndio
- Kwa kliniki ya wajawazito : Ndio
- Kwasababu ya ugonjwa , eleza: _____ Ndio
- Nyingine, eleza: _____
- Sijui:
- Amekataa:

19. Je, sasa hivi unatumia madawa ya kurefusha maisha? Ndio La
Sijui

Amekataa
Nyingine, elezaKama

ndio,

a) Eleza madawa unayotumia sasa : _____

b) tarehe ya awali uliyoanza ____ / ____ / ____ sijui

c) Je, unajua sababu ulioanzishwa madawa ya kurefusha maisha?

- Kwasababu ya ugonjwa, eleza: _____ Ndio
- Kwasababu ya CD4 kuwa chini : Ndio
- Kwasababu ya viwango vya virusi kuwa juu: Ndio
- Nyingine, eleza: _____
- Sijui:
- Amekataa

20. Maoni

ANUANI NA ULAJI

Study ID number _____

Interviewer number _____

Date of interview (day/month/year) ____/____/____

Format(tick one)

New

Update (fill only updated info)

A. YA KIBINAFSI

19. Jina Lako Nani?

- a. Jina la kwanza
- b. Jina la katikati
- c. Jina la mwisho

20. Wajulikana kwa jina lipi eneo unako ishi?

21. Jina kamili ya eneo unako ishi.

- a. Wilaya
- b. Mji
- c. Kijiji
- d. Mtaa
- e. Nambari ya ploti
- f. Nambari ya mlango
- g. Jina la barabara

22. Usafiri wa umma hadi kwako nyumbani:

4a. Aina: F Bus F Boda boda F Matatu F Taxi F Mengine(Fafanua)

4b. Numbari ya gari iendako eneo unakoishi:

4c. Jina la kituo cha kuabiria magari?

4d. Jina inalojulikana eneo unakoishi.

23. Maelezo ya njia kutoka kwako hadi kituo cha kuabiria magari?

24. Alama ya kuwa msaidizi katika kuuweka kaya: (Majina ya shule, makanisa, biashara nk)

25. Je, unaweza kufikiwa kwa njia ya simu? Ndiyo La
Kama ndiyo,

(i) Nambari ya simu yako 1?

(ii) Nambari ya simu yako 2?

iii. Nani anaye beba simu hiyo (Weka alama kwa moja) Mimi Mwingine, fafanua

B. MAWASILIANO MUHIMU

26. Je, kuna mtu mwingine ambaye anjua hali yako ya HIV ambaye tunaweza kuwasiliana kwa njia ya simu kama hatuwezi kufikia wewe moja kwa moja? If no skip to 9.

27. Jina lake ni nani?
a. Jina la kwanza
b. Jina la katikati
c. Jina la mwisho
28. Uhusiano gani upo kati yako na mtu huyu ?
29. (a) Nambari yake ya simu1?
(b) Nambari yake ya simu2?

C. MAWASILIANO KUHUSU KWAKO MASHAMBANI

30. Je, una nyumbani bara? Ndiyo La
If no, skip to 19
31. Ni wapi kwako Bara?
a. Wilaya
b. Mji
c. Kijiji
d. Mtaa
e. Nambari ya ploti
f. Nambari ya mlango
32. Maelekezo maalum kwa makazi
33. Je, kuna mtu u ko bara ambaye tunaweza kuwasiliana kama hatuwezi kufikia wewe moja kwa moja? Ndiyo La
If no, skip to 19
34. Jina lake huyu mtu ni nani?
a. Jina la kwanza
b. Jina la katikati
c. Jina la mwisho
35. Uhusiano gani upo kati yako na mtu huyu ?
36. Je huyu mtu ako na simu? Ndiyo La
Iwapo ndiyo , fafanua

(a) Nambari yake ya simu1?
(b) Nambari yake ya simu2?

D. ANDRA

Maoni

SHEDDING

Study ID Number ___ ___ ___

Interviewer Number ___ ___

Date of interview (day/month/year) ___ ___ / ___ ___ / ___ ___ ___ ___

Visit (tick one) Week 1 FU Week 2 FU Week 3 FU Other, specify _____

A: MEDICAL HISTORY

1. Unasikia maumivu ya tumbo ya chini?

ndio la

(Ndio, fafanua)

- a. muda wa ___ ___ (siku)
- b. Ukali (kidogo 1-5, 5 kuwakali zaidi)

9. Umekuwa na historia ya kutokwa na damu ambaye si yakawaida? Fafanua

Ndio La

(Ndio, fafanua)

- g. muda wa ___ ___ siku
- h. kiasi
- doasuruali)

Inahitajipediya usafi

Mwingine, eleza

Unatokwa na majimaji ya uke yasiyo ya kawaida?

10.

Ndio la

(Ndio, fafanua)

- i. muda wa ___ siku)
- j. rangi ya
 - Majano

hudurungi

nyeupe

wazi

Nyingine ,fafanua

k. harufu

Harufu mbaya

Hakunaharufu

Mengine,eleza

l. kiasi

inadoasuruali

Inahitajipediya usafi

Mengine,eleza

11. Mgonjwa amekuwa nahoma yoyotebaada ya ziara yamwisho?

Ndio la

kama ndio, fafaua muda...siku

12. Je,mgonjwaametafutahuduma zamatibabu kwa ajili yahaya aumalalamiko mengine?

Ndio La

If yes, specify what complaint prompted the participant to seek care:

Uchungu wa tumbo ya chini

Homa

Damu kutoka uzazi wa kike

Uchafu unaotoka katika uzazi wa kike

e. Mengine, fafaua

If yes, specify where the participant sought care:

Study clinic

Hope Center

Coptic Hospital

KNH

f. Other, specify

13. Was the participant's condition possibly due to a study procedure? (Ask if any of questions 1-4 is yes)

Yes No

If yes, fill out a Complications questionnaire.

14. Damu yako ya mwezi ya mwisho ilikuwa lini? (dd/mm/yyyy)

15. Umewahi fanya ngono uke tangu utibiwe? Ndio la Kataa

Nyingine fafaua _____

8 (a). Ndio, mara ngapi? __

8(b) Kamawamefanya mapenziuketangumatibabu,je,ni mara ngapikutumia kondomu wakati wa ngono?

Kila wakati (100%) Wengi wa wakati huo (75-99%)
 Nusu ya mudu (50%-74%) Wakati mwingine (25-49%)
 Mara chache (1-25%) Kamwe (0%)
 Amekataa Sijui
 Nyingine, eleza

9. Unafikiri unaweza kuwa mjamzito

Ndio la

10. Kwa sasa unatumia madawa ya kupungua makali ya virusi

Ndio Kataa Nyingine,fafaua

La Sijui

Ndio, fafaua madawa unayo tumia sasa na tarehe uliyo yaanza

- d4t, 3tc, nvp
- d4t, 3tc, efv
- azt, 3tc, nvp
- azt, 3tc, efv
- zingine, fafanua

Tarehe ya kuanza ARV awali (dd /mm /yyyy)

11 Katika siku 7 ya mwisho, mgonjwa alikosa kumeza tembe ngapi za dawaza kurefusha maisha?

12 Katika siku 30 ya mwisho, mgonjwa alikosa kumeza tembe ngapi za dawaza kurefusha maisha?

(If Q11 and Q12 patient did not MISS taking any pills skip to Q14)

13. If the patient missed any doses, please specify reasons (check all that apply)

- Toxicity/ side effect
- Share with others
- Forgot
 - g. Felt better
- Too ill
- Stigma, disclosure or privacy issues
- Drug out of stock
- Patient lost or ran out of pills
- Delivery /travel problem
- Inability to pay
- Alcohol
- Depression
 - h. Other specify

B: DIAGNOSIS

14. Is there any new infection that was related to the procedure since the last visit?

Yes No

If yes, specify and fill a Complications and/or Adverse Event questionnaire:

15. Were there any new complications diagnosed today related to the treatment?

Yes No

If yes, specify and fill a Complications and/or Adverse Event questionnaire:

C: TREATMENT

16. Was any treatment provided today? Yes No

If yes, specify

17. Was the patient referred for further cervical treatment at another institution?

Yes No

17 (i) If yes, specify institution

17(ii) If yes, specify reason for referral

D: SPECIMEN COLLECTION

18. Did you collect cervical HIV swab? Yes No

If no, specify reason:

19. Did you collect blood? Yes No

If no, specify reason:

E: ACCEPTABILITY OF TREATMENT

20. Please complete the items listed below by placing a checkmark on the box next to each question that best indicates how the client feels about the treatment she received.

	SIKUBALI KAMWE	haukubaliani	kadri,	kukubaliana	sanakukubaliana
(i) Naona hii njia ni mwafaka kwa kukabiliana na magonjwa ya njia ya uzazi)	<input type="checkbox"/>				
(ii) Naweza tumia haya ya matibabu nikipatamagonjwa kama haya siku ya mbeli	<input type="checkbox"/>				
(iii) ningependekeza haya matibabu kwa mtu mwingine	<input type="checkbox"/>				
(iv) kwa jumla nakubaliana na haya matibabu)	<input type="checkbox"/>				

F: OTHER

24. Comments:

7. Je, mfanyakazi wahuduma za afya alikuambiasababu ya kifo? Ndio/ La/ sijui/
 If yes go to 7a, if no go to 8

7a. Ni mfanyakazi wahuduma za afya wanasemakilichosababisha kifo?

8. Alifanyiwa upasuaji wowote kwa sababu ya ugonjwa? Ndio/ La/ sijui/
 If yes go to 8a, if no go to 9

8a. Upasuaji ulikuwa muda gani kabla ya kifo? _____ siku

8b. Sehemu gani ya mwili alifanyiwa upasuaji?

_____ (Kifua/ Kichwa/ Mengine (fafanua) _____)

9. Mpenzi (wapeenzi) wa marehemu wowote wamekufakatika kipindi cha miaka 5 iliyopita?

Ndio/ La/ Haijulikani/
 If yes go to 9a, if no go to 10

9a. Kama ndiyo, ni nini waliamini kilichosababisha kifo cha mpenzi(s)

9a1. Partner 1: _____

9a2. Partner 2: _____

Injury/accident/suicide

10. Je, alikabiliwa na majeraha yote au ajali ambayo imesababisha kifo chake? Ndio/ La/ sijui/

If yes go to 10a, if no go to 11

10a. Ni aina gani ya majeraha au ajali marehemu aliteseka nayo?

_____ (Barabara ya ajalizi barabarani/ Kuanguka/ Kuzama/ Sumu)

_____ (Kuanguka/ Magombano/ Mengine/ Haijulikani)

10b. Majeruhi au ajali ilikuwa ya makusudi au ilifanywa na mtu mwingine) ndio/ La/ sijui/

10c. Je, unafikirikwambayeye alijiua?) ndio La sijui

11. Je, yeye aliumwa na mnyama/mdudu yeyote iliyosababishakifo chake? Ndio
La Sijui

11a. Kama ndiyo, niaina yamnyama/mdudu?

History of previously known medical conditions

12. Did the deceased suffer from any of the following conditions?

- a. High blood pressure Yes No Unknown
- b. Diabetes Yes No Unknown
- c. Asthma Yes No Unknown
- d. Epilepsy Yes No Unknown
- e. Malnutrition Yes No Unknown
- f. Cancer Yes No Unknown

f1. If yes, specify type of cancer or site: _____

- g. Tuberculosis Yes No Unknown
- h. Any other medically diagnosed illness? Yes No Unknown

h1. If yes, specify: _____

13. Signs, symptoms, and their severity during the last illness:

Symptom/ Signs	Symptom present?	If present, duration of symptom
a. Fever	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> ≤2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
b. Loss of weight	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> ≤2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
c. Diarrhea	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> ≤2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
d. Vomiting/associated abdominal pain	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> ≤2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
e. Constipation/associated abdominal pain	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> ≤2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
f. Cough	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> ≤2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown

g. Cough followed by vomiting	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
h. Breathing trouble (chest indrawing/difficult/rapid/wheezing)	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
i. Neck stiffness	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
j. Unconscious episodes	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
k. Fits	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
l. Jerking of individual limbs	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
m. History of epileptic illness in earlier years	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
n. Paralysis of limbs	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
o. Rigid body stiffness, unable to open mouth	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
p. Red and sore eyes	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
q. Skin rash and itching	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
r. Herpes Zoster (at any time in life)	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
s. Abscesses/body sores	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
t. White patches on the inside of mouth and tongue	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
u. Oedema	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
v. Hair changes	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
w. Yellowing of eyes or passing of brown urine	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
x. Chest pain	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown
y. Other (Specify:)	FNo F Yes	F ≤2 weeks F >2 weeks F Unknown

Unexpected vaginal bleeding or discharge	No F Yes	2 weeks F >2 weeks F Unknown
Pelvic or vaginal pain	No F Yes	2 weeks F >2 weeks F Unknown

14. Records available in home, e.g., death certificate (extract findings):

15. Comment

EXIT

Study ID Number: ___ ___ ___

Interviewer Number: _____

Date of visit (day/month/year) ___ ___ / ___ ___ / ___ ___ ___

1. Is the patient exiting the study because she has completed the study protocol and qualifies as per study guidelines to exit the study at this time? Yes No (*If Yes, go to 9, If No go to 2*)

2. Date of last study visit (DD/MM/YYYY) HH/H/HHH

3. Date last seen by study staff (DD/MM/YYYY) HH/H/HHH

4. Has the patient accessed care at the Hope Clinic in the past year? Yes No

5. Did you talk to the patient? Yes No (*If Yes go to 6, If No go to 5a*)

5a. If no, who was the source of information (tick one)

Clinician / clinic staff

Employer

Treatment supporter

Spouse or Partner

Friend

Other (specify)

Family member / Relative

Neighbor

Mother or Father

Caregiver

6. Je, mgonjwa anahamisha huduma ya HIV kwa mpangilio mwingine? Ndiyo La
(If Yes go to 6a, If No go to 7)

6a. If yes, where is the patient transferring care to (tick one)

Kuhamishiwa kliniki ingine ambayo siya HOPE, elezea

Kuhamishiwa kliniki ya Industrial area

Kuhamishiwa kliniki ya Maseno

7. Has Hope Clinic asked the patient to Exit or leave the program? Yes No (*If Yes go to 7a, If No go to 8*)

7a. Reasons for being Exited from the program (tick all that apply)

Client has not returned to clinic for 1 year

Poor Adherence

Poor Clinic Attendance

Not willing to disclose HIV status

Referred else where

Tested Negative

Other (specify)

Unknown

8. Has the patient asked to be Unenrolled from the study? Yes No (*If Yes go to 8a, If No go to 9*)

8a. Reasons patient is asking to Unenroll from the study (tick all that apply)

Kutokuwa na nia ya kuhudhuria

Muda wa kusubiri kuwa mrefu sana

Kutoambana na kazi

Matatizo ya fedha

Kukosa nia ya kuhudhuria kwa sababu ya matatizo ya kiafya

Matatizo ya familia

Mteja kwenda kwa mganga waimani

Kutokuwa tayari kusema hali yake ya Ukimwi

Kutumwa kwingine

Nyingine (taja)

Haijulikani

9. At what point was the patient exited? After Pap smear After Biopsy

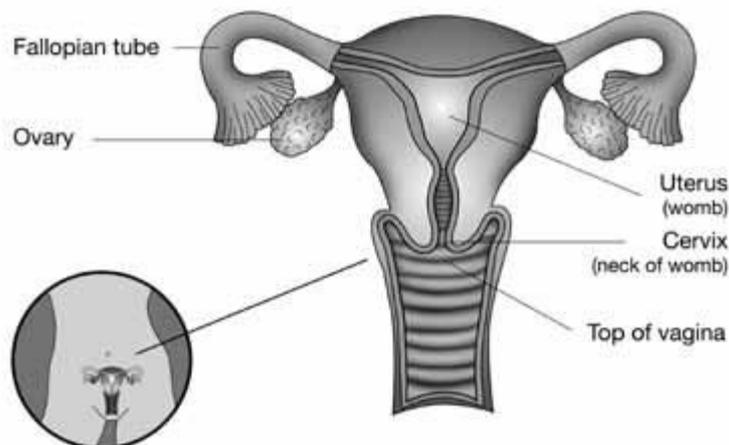
Mortality Withdrawn from study Other (If other specify)

10. Comment

CERVICAL TREATMENT STUDY

What is a Pap smear?

A Pap smear is a simple test to check your cervix to make sure it is healthy. Your cervix is the opening of the uterus, and is at the top of your vagina (see the diagram below). A Pap smear takes only a few minutes and is not painful. Having a Pap smear every two years is the best way to prevent cancer of the cervix.



Why have a Pap smear?

A Pap smear can show the early warning signs of cancer of the cervix. Sometimes the cells of the cervix change from healthy to unhealthy (abnormal). A Pap smear can find abnormal cells before cancer develops.

What causes cervical cancer?

An infection with a virus called HPV (human Papillomavirus) is the cause of almost all cervical cancers. There are over 100 different types of HPV. Two of these types are known to cause most of the cervical cancer cases. HPV is very common. Most people (four out of five) will have HPV at some time in their lives. Anyone who has ever had sex can have HPV.

In most cases, HPV clears up by itself in a few years. Sometimes the virus can stay in your body longer, and can lead to cervical cancer. This usually takes a long time – about 10 years. A Pap smear every two years can find cell changes caused by HPV before they turn into cancer. Your doctor, nurse or health worker can then make sure your health is monitored and that you get treatment if you need it, so you can stay healthy.

How is a Pap smear done?

First the doctor or nurse asks you to undress from the waist down and to lie on your back for the examination. You can ask for a female doctor or nurse. Next the doctor or nurse will use a speculum (medical instrument) to open your vagina so your cervix can be seen more clearly. Some cells are gently wiped from your cervix with a small brush or spatula (a small plastic or wooden stick). The cells are placed on a glass slide and sent to a laboratory where they are looked at under a microscope.

What does it feel like?

Sometimes having a Pap smear can be a little embarrassing. Remember, for the person doing the smear, this is just part of their everyday work and they are not embarrassed. The

procedure might be a bit uncomfortable, but it shouldn't hurt. If it hurts, tell your doctor, nurse or health worker straight away.

What if my results are not normal?

If your results are not normal this does not mean you have cancer. Very often it will be that you have something simple like an infection that will clear up naturally. Sometimes a woman may need to have a Pap smear more often. Some types of abnormal cells may need to be treated by a specialist. Make sure you talk to your doctor, nurse or health worker about what is best for you.

HIV and cervical cancer

HIV-positive women are more likely to be infected with human Papillomavirus (HPV), the primary cause of cervical cancer, and progress to invasive, life-threatening disease than those who are HIV-negative.

Note: Being on HIV medication does NOT reduce your risk of cervical cancer.

Cervical Treatment Study

Researchers from the University of Washington in the USA, Kenyatta National Hospital, WHO and Coptic Hospital are conducting a study to see how treatment can prevent abnormal cells from becoming cancer, and how treatment might affect HIV. Those patients who enroll in the study will be given free screening for cervical disease that may develop into cancer and will also receive free treatment if they are found to have abnormal cells.

For more information about the study, contact:

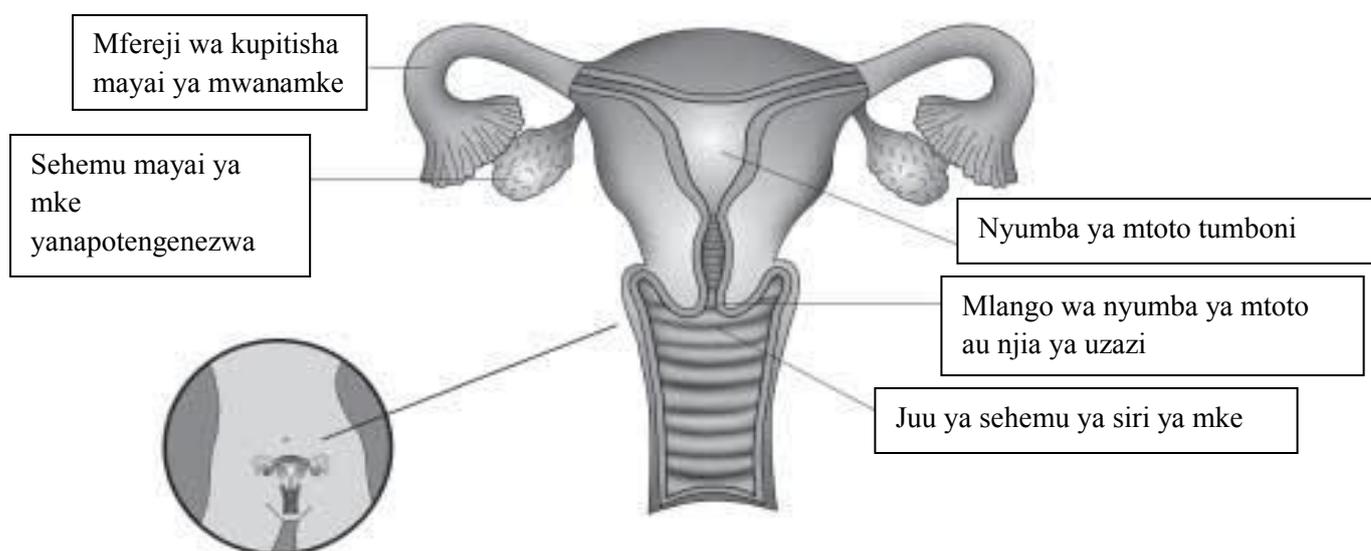
1. Peter Juma: 0721-898-785
2. Elizabeth Makena: 0728-456-540
3. Dr. Evans Malava: 0721-289-733

CERVICAL TREATMENT STUDY

Je Pap Smear ni nini?

Pap Smear ni kupimwa njia yako ya uzazi kwa njia rahisi kuhakikisha ni salama. Njia ya uzazi ni mlango wa nyumba ya mtoto tumboni na iko kwa ndani juu ya sehemu yako ya siri (angalia mchoro hapo chini). Kupimwa njia ya uzazi huchukuwa dakika chache tu na si uchungu.

Kupimwa njia ya uzazi kila baada ya miaka miwili ndio njia bora zaidi ya kuzuia saratani ya njia ya uzazi.



Kwa nini upimwe njia ya uzazi?

Kupimwa njia ya uzazi kunaweza kuonyesha dalili za saratani ya njia ya uzazi. Wakati mwingine hali ya njia ya uzazi hubadilika, na kuwa na hitilafu. Ukipimwa njia ya uzazi inaweza kujulikana kama ina hitilafu kabla haijabadilika kuwa saratani.

Nini kinachosababisha Saratani wa njia ya uzazi?

Kuambukizwa kwa virusi vinavyoitwa HPV (humanpapillomavirus) kunasababisha karibu saratani zote za njia ya uzazi. Kuna zaidi ya aina 100 tofauti za HPV. Aina mbili za HPV zinajulikana kusababisha saratani kwa karibu wote wanaouguu njia za uzazi. Virus vya HPV vinapatikana kwa wingi. Watu wengi (wanne kwa watano) watakuwa na HPV wakati mmoja maishani mwao. Yeyote ambaye ashawahi kufanya ngono anaweza kuwa na HPV.

Mara nyingi, virusi vya HPV hutoweka vyenyewe baada ya miaka michache. Mara nyingine hivi virusi vinaweza vikakaa mwilini na vinaweza kusababisha saratani ya njia ya uzazi. Hii huchukuwa mda mrefu – kama miaka 10. Kupimwa njia ya uzazi kila baada ya miaka miwili kunaweza kufanya mabadiliko ya njia ya uzazi yajulikane kabla hayajasababisha saratani. Daktari wako, muuguzi au mfanya kazi kutoka kituo cha afya anaweza kuhakikisha kuwa afya yako inafuatiliwa na unapata matibabu kama unayahitaji ili uwe na afya bora.

Je mtu hupimwaje njia ya uzazi?

Kwanza daktari au muuguzi anakuuliza uvue nguo kutoka kiunoni kwenda chini na ulale chali ili akupime. Unaweza kuuliza upimwe na daktari au muuguzi wa kike. Halafu daktari au

muuguzi atatumia chombo cha kupimia kufungua sehemu yako ya siri ili njia ya uzazi ione kane vyema zaidi. Atapangusa ukuta katika njia yako ya uzazi na brashi ndogo au chombo cha kupima (kijiti kidogo ama kipande cha mpira kidogo). Kilichopanguswa kutoka njia ya uzazi, wataweka kwenye kioo kidogo na kupelekwa maabara (lebu) watakakotazama wakitumia darubini.

Je mtu husikiaje?

Wakati mwingine kupimwa njia ya uzazi ni jambo la kutahayari au kuonea aibu. Kumbuka, kuwa

anayekupima njia ya uzazi, hii ni kazi yake ya kawaida na hatahayari au kuona aibu.

Unapopimwa, unaweza kusikia ovyo, lakini haina uchungu. Ukisikia uchungu, mwambie daktari wako, muuguzi au mfanyakazi kwenye kituo cha afya hapohapo.

Je kama majibu yangu sio sawa au salama?

Kama majibu yako sio sawa, haimaanishi uko na saratani. Mara nyingi itakuwa una jambo dogo kama kuambukizwa ambako huisha kwenyewe bila kutibiwa. Mara nyingine mwanamke anahitaji kupimwa njia ya uzazi mara kwa mara. Hali nyingine zisizo za kawaida zahitaji matibabu maalum. Hakikisha umeshauriana na daktari wako, muuguzi au mfanya kazi wa kituo cha afya ili akufahamisha kilicho bora kwako.

Uhusiano wa UKIMWI na saratani ya njia ya uzazi

Wanawake wanaouguwa ugonjwa wa UKIMWI wanauwezekano mkubwa kuliko wale ambao hawana UKIMWI kuambukizwa virusi vya human Papillomavirus (HPV), vinavyosababisha saratani ya njia ya uzazi, na iendelee hadi ivaamie mwili na iwe, kitisho kwa maisha.

Kumbuka: Kutumia dawa ya UKIMWI haiwezi kupunguza hatari ya saratani ya njia ya kizazi.

Utafiti wa matibabu ya njia ya kizazi (Cervical Treatment Study)

Watafiti kutoka Chuo Kikuu cha Washington huko Marekani, Hospitali Kuu ya Kenyatta, Shirika la Afya Duniani na Hospitali ya Coptic wanafanya utafiti kuona jinsi gani tiba inaweza kuzuia chembe chembe zisizo za kawaida isiwe saratani ya njia ya kizazi, na jinsi tiba inaweza kuathiri ugonjwa wa UKIMWI. Wagonjwa ambao watahusika katika utafiti huu watapewa uchunguzi (Pap smear) wa bure na pia watapata matibabu ya bure wakipatikana na chembe chembe zisizo za kawaida.

Kwa habari zaidi kuhusu utafiti huu, wasiliana na:

1. Peter Juma: 0721-898-785
2. Elizabeth Makena: 0728-456-540
3. Dr. Evans Malava: 0721-289-733

COPTIC HOPE MEDICAL RECORD FORMS

HOPE CLINIC ADDRESS AND INTAKE

HOPE ID Number

Site Code

Today's Date (DD.MM.YYYY)

Interviewer number

Name (First, Middle,Last)

1. Gender Male Female

2. Age (Years) Month 3. Date Of Birth (DD/MM/YYYY) / /

4. In which region does the client reside(stay)?

- Nairobi province (see 4a) Rift Valley Province
- Nyanza Province (see 4b) North Eastern Province
- Western Province (see 4b) Eastern Province
- Central Province Coast Province
- Other country (specify)

4a. If Nairobi province, specify area: (tick one)

- Langata/Kibera Westlands
- Starehe Kasarani
- Dagoretti Kamukunji
- Makadara Eastland/ Industrial Area
- Embakasi Other (specify)

4b. If western or Nyanza, specify area: (tick one)

- Kisumu Rural Maseno
- Siaya Chulaimbo
- Vihiga Lela
- Kakamega Other (Specify)
- Luanda

5. Email

6. Physical Address

HOPE CLINIC ADDRESS AND INTAKE

7. How long has the client stayed at this residence?

- Less than a year Greater than a year

8. How long does the client plan to stay at this residence?

- Less than a year Greater than a year

9. Phone number (Cell)

9a. Relationship to phone owner (tick one)

- Self Employer

- Relative Other (specify)

- Friend

10. Phone number (Landline)

10a. Relationship to phone owner (tick one)

- Self Employer

- Relative Other (specify)

- Friend

Emergency contact (in case we cannot reach the patient)

11. Name (Last, First, Middle)

11a. Relationship to client (tick one)

- Self Employer

- Relative Other (specify)

- Friend

- Husband/Wife

11b. Phone No.

Clients transferring from Pediatric Clinic

12. Pediatric Hopeid

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

HOPE CLINIC COUNSELING ADHERENCE #1

HOPE ID Number

Site Code

Today's date (DD.MM.YYYY)

 /
 /

Interviewer number

Checklist	Tick
Explain about HIV and how it affects the body	<input type="radio"/>
Explain about CD4 cells and why it is necessary to measure the CD4 count	<input type="radio"/>
Explain the difference between HIV and AIDS.	<input type="radio"/>
Explain about ARV.	<input type="radio"/>
Explain ARV is not a cure.	<input type="radio"/>
Explain the cause of resistance.	<input type="radio"/>
Explain treatment failure	<input type="radio"/>
Explain importance of adherence.	<input type="radio"/>
Explain problem of side effects.	<input type="radio"/>
Have patient think about life long commitment of therapy.	<input type="radio"/>
Have patient think about ability to follow up care	<input type="radio"/>
Explore patient support system	<input type="radio"/>
Discuss adherence promotion strategies e.g. treatment buddy, pill diary e.t.c	<input type="radio"/>

2. Identify barriers to adherence (tick all that apply)

- | | |
|---|--|
| <input type="radio"/> Poor Communication | <input type="radio"/> Lack of social support |
| <input type="radio"/> Inadequate understanding about HIV/AIDS | <input type="radio"/> Alcohol/ Drug use |
| <input type="radio"/> Failure to disclose status | <input type="radio"/> None |
| <input type="radio"/> Mental State | <input type="radio"/> Others (specify) |
| <input type="radio"/> Stigma | <div style="border: 1px solid black; width: 150px; height: 20px;"></div> |
| <input type="radio"/> Low literacy | |

3. Does the patient need to move forward with the protocol or to repeat counselling adherence #1 ?

- Move forward Repeat Counselling Adherence # 1

Notes/Remarks

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

HOPE CLINIC COUNSELING ADHERENCE #2

HOPE ID Number

Site Code

Today's date (DD.MM.YYYY)

Interviewer number

--	--	--	--

--

--

--	--	--	--	--

--	--	--

Score scale: 1-Poor 2-Fair 3-Good

Section A: Knowledge Assessment

Question	Rationale	Score
1. What do you know about ARVs?	Assess whether information given in Counseling Adherence#1 has been understood.	<input type="checkbox"/>
2. What are the names of any ARVs?	Assess the client knows that AZT, NVP, etc. are ARVs, but septrin is not	<input type="checkbox"/>
3. How do ARVs work?	Assess the client's knowledge of basic ARV action (especially that ARV is not a cure for HIV infection.)	<input type="checkbox"/>
4. What side effects are associated with ARVs? What do you know about them?	Assess client's knowledge of side effects related to his/her ARV regimen and the appropriate response to deal with side effects.	<input type="checkbox"/>
5. How long should you normally take ARVs?	Assess whether client knows ARV is life long treatment.	<input type="checkbox"/>
6. What happens if you dont take ARVs consistently?	Assess whether client understands the problem of resistance given ARV interruptions.	<input type="checkbox"/>
7. What is the purpose of CD4 counts?	Assess whether client knows that CD4 count is a laboratory indicator for monitoring the effect of ARV	<input type="checkbox"/>
8. What are your expectations from ARVs?	Assess whether client has realistic expectations, e.g., prolonging life, keeping them well enough from their family, e.t.c. Assess for false expectations, e.g., a cure for HIV, e.t.c	<input type="checkbox"/>
9. Can someone still transmit HIV while taking ARVs?	Assess/review need for continued prevention e.g. condom use.	<input type="checkbox"/>
If total score is less than 18, patient has failed section A If total score is 18 or above, the patient passes section A		Total Score: <input style="width: 20px; height: 20px;" type="text"/> <input style="width: 20px; height: 20px;" type="text"/>

Section B: Counselor Assessment

Question	Rationale	Score
10. Assess for barriers that help determine capability for followup	Assess whether client can attend HIV clinic for follow up medical and counselling care	<input type="checkbox"/>
11. Ask the client whether s/he has a relative/friend whom s/he can rely on to support her/him taking ARV	Assess availability of support from home	<input type="checkbox"/>
12. On a scale of 1 to 5 (5 being most ready, 1 being least), please rate the client's ability to adhere to medications	Score 1: Patient will adhere to the ARVs very poorly (Misses more than half the doses) Score 2: Patient will likely miss doses of ARVs on a regular basis (misses up to 50% of doses) Score 3: Patient will only miss some doses of ARVs (1 dose a month at most) Score 4: Patient will rarely miss a dose of ARVs (1 dose every 6 months at most) Score 5: Patient will almost never miss any doses (1 dose every year at most)	<input type="checkbox"/>
If total score is 2 or below on question 12, patient fails section B If total score is 3 or above question 12, patient passes section B		

HOPE CLINIC COUNSELING ADHERENCE #2

Section C: Participant Assessment

Question	Rationale	Score
13. Ask the client if s/he can come to HIV clinic for regular follow-up?	YES or NO answer	<input type="checkbox"/> Yes <input type="checkbox"/> No
14. Do you want to start ARV treatment now?	YES or NO answer	<input type="checkbox"/> Yes <input type="checkbox"/> No
If any answers in this section are "NO" then patient fails Section C		

Section D: Final Assessment

15. a) Did the patient pass section A, B and C?

Yes (Patient moves forward) No

b) If NO, is the patient scheduled to repeat counseling?

Yes No

Notes/Remarks

For the data use only (tick after scanning the form)

Scanned Date ___/___/_____ Name of data person

HOPE CLINIC COUNSELING ADHERENCE #3

To be performed 1 month after starting ARVs

HOPE ID Number

Site Code

Today's date (DD.MM.YYYY)

Interviewer number

Score scale: 1- poor 2-Fair 3-Good

Section A: Knowledge Assessment

Question	Rationale	Score
1. What do you know about ARVs?	Assess whether information given in Counseling Adherence#1 has been understood.	<input type="checkbox"/>
2. What are the names of your medications and dosage?	Assess whether the client knows his/her medication and dosage.	<input type="checkbox"/>
3. How do ARVs work?	Assess the client's knowledge of basic ARV action (especially that ARV is not a cure for HIV infection.)	<input type="checkbox"/>
4. What side effects are associated with ARVs and what do you do if you have side effects?	Assess client's knowledge of side effects related to his/her ARV regimen and the appropriate response to deal with side effects.	<input type="checkbox"/>
5. How long should you normally take ARVs?	Assess whether client knows ARV is life long treatment.	<input type="checkbox"/>
6. What happens if you don't take your ARVs consistently?	Assess whether client understands the problem of resistance given ARV interruptions.	<input type="checkbox"/>
7. What is the purpose of CD4 counts?	Assess whether client knows that CD4 count is a laboratory indicator for monitoring the effect of ARV	<input type="checkbox"/>
8. What are your expectations from ARVs?	Assess whether client has realistic expectations, e.g. prolonged life, keeping them well enough from thier family, etc. Assess for false expectations, e.g., a cure for HIV, etc.	<input type="checkbox"/>
9. How can someone still transmit HIV while taking ARVs?	Assess review need for continued prevention e.g. condom use	<input type="checkbox"/>
Total Score		<input style="width: 20px;" type="text"/>

Section B: Counselor Assessment

Question	Rationale	Score
10. During the last 7 days how many pills did the patient MISS taking?	Get exact number of pills missed	<input type="checkbox"/>
11. On a scale of 1 to 5 (5 being most ready, 1 being least), please rate the client's ability to adhere to medications?	Score 1: Patient will adhere to the ARVs very poorly (Misses more than half the doses) Score 2: Patient will likely miss doses of ARVs ona regular basis (Misses up to 50% of doses) Score 3: Patient will only miss some doses of ARVs (1 dose a month at most) Score 4: Patient will rarely miss a dose of ARV (1 dose every 6 months at most) Score 5: Patient will almost never miss any doses (1 dose every year at most)	<input type="checkbox"/>

Notes/Remarks

For the data use only (tick after scanning the form)

Scanned Date ___/___/_____ Name of data person

HOPE CLINIC PHONE NUMBER AND ADDRESS UPDATE FORM

HOPE ID Number

Site Code	Today's Date (DD.MM.YYYY)	Interviewer number
<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="text"/> / <input type="text"/> / <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="text"/> <input type="text"/> <input type="text"/>

Name (First, Middle,last)

1. Has the client's phone number changed? Yes No *(If Yes go to 2, If No go to 4)*

2. Phone number (Cell)

2a. Relationship to phone owner (tick one)

- | | |
|--|---|
| <input type="checkbox"/> Self | <input type="checkbox"/> Friend |
| <input type="checkbox"/> Husband or wife | <input type="checkbox"/> Employer |
| <input type="checkbox"/> Relative | <input type="checkbox"/> Other (specify) <input style="width: 150px;" type="text"/> |

3. Phone number (Landline)

3a. Relationship to phone owner (tick one)

- | | |
|--|---|
| <input type="checkbox"/> Self | <input type="checkbox"/> Friend |
| <input type="checkbox"/> Husband or wife | <input type="checkbox"/> Employer |
| <input type="checkbox"/> Relative | <input type="checkbox"/> Other (specify) <input style="width: 150px;" type="text"/> |

4. Has the client moved? Yes No *(If Yes go to 5, If No go to 7)*

5. Physical Address

6. In which region does the client now reside (stay) *(For Maseno, check 2nd, 3rd and where applicable 4th column)*

- | | | |
|--|---|---|
| <input type="checkbox"/> Nairobi province (See 6a) | <input type="checkbox"/> Central province | <input type="checkbox"/> Eastern province |
| <input type="checkbox"/> Nyanza province (See 6b) | <input type="checkbox"/> Rift Valley province | <input type="checkbox"/> Coast province |
| <input type="checkbox"/> Western province (See 6b) | <input type="checkbox"/> North Eastern province | |

6a. If Nairobi province, specify area: (tick one)

- | | |
|---|--|
| <input type="checkbox"/> Langata/Kibera | <input type="checkbox"/> Westlands |
| <input type="checkbox"/> Starehe | <input type="checkbox"/> Kasarani |
| <input type="checkbox"/> Dagoretti | <input type="checkbox"/> Kamukunji |
| <input type="checkbox"/> Makadara | <input type="checkbox"/> Eastlands/Industrial Area |
| <input type="checkbox"/> Embakasi | <input type="checkbox"/> Other(Specify) <input style="width: 150px;" type="text"/> |

6b. If western or Nyanza, specify area: (tick one)

- | | |
|---------------------------------------|--|
| <input type="checkbox"/> Kisumu Rural | <input type="checkbox"/> Maseno |
| <input type="checkbox"/> Siaya | <input type="checkbox"/> Chulaimbo |
| <input type="checkbox"/> Vihiga | <input type="checkbox"/> Lela |
| <input type="checkbox"/> Kakamega | <input type="checkbox"/> Other(specify) <input style="width: 150px;" type="text"/> |
| <input type="checkbox"/> Luanda | |

HOPE CLINIC

PHONE NUMBER AND ADDRESS UPDATE FORM

7. Has the client's email changed? Yes No *(If Yes go to 8, If No go to 9)*

8. Email

Emergency contact (in case we cannot reach the patient)

9. Has the client's emergency contact changed? Yes No *(If Yes go to 10)*

10. Name (Last, First, Middle)

10a. Relationship to client (tick one)

Self Employer

Relative Other (specify)

Friend

Husband/Wife

10b. Phone No.

For the data use only (tick after scanning the form)

Scanned Date ___/___/_____ Name of data person

HOPE CLINIC PATIENT PHONE CONTACT FORM

HOPE ID Number

Site Code **Today's Date (DD.MM.YYYY)** **Interviewer number**
 / /

1. Date of phone contact (DD/MM/YYYY) / /
2. Did you talk to the patient or patient's contact?
 - Yes, talked to patient *(Go to 4)*
 - Yes, talked to patient's contact *(Go to 3)*
 - No *(Go to 4)*
3. If talked to patient's contact, who was the source of information *(tick one)*
 - Clinician/clinic staff Employer Caregiver
 - Spouse or partner Friend Treatment supporter
 - Family member/Relative Neighbour Other (specify)
 - Mother/Father
- 3a. Did the source of information have credible knowledge for whether the patient was alive or dead?
 - Yes, credible and patient confirmed alive *(Go to 4)*
 - Yes, credible and patient confirmed dead *(Go to 9 and complete Mortality form)*
 - Source did not know whether patient was dead or alive *(Go to 4)*
4. What was the reason for calling or contacting the patient or patient's contact?
 - Patient missed clinic appointment *(Go to 5)*
 - Patient missed pharmacy pickup *(Go to 6)*
 - Other (specify) *(Go to 7)*
5. Reasons for missed clinic appointment *(tick all that apply)*
 - N/A, did not reach patient or patient's contact Unable to attend because of health problems
 - No longer willing to attend Family problems
 - Wait time too long Client will go to faith healer
 - Conflict with work Unwilling to disclose
 - Financial problems Other (specify)
 - Client moved or relocated
6. Reasons for missed pharmacy pick up *(tick all that apply)*
 - Unable to attend because of health problems Got medication somewhere else
 - Family problems Client will go to faith healer
 - Conflict with work Unwilling to disclose
 - Financial problems N/A, did not reach patient
 - Client moved or relocated Other (specify)

7. If talked to patient, did the patient wish to remain in the clinic?

- Yes (Go to 7a)
- No (Go to 8 and complete Exit form)
- N/A, did not talk to patient (Go to 9)

7a. Did the patient schedule a clinic appointment? Yes No (If YES go to 7b, If NO go to 7d)

7b. If YES, date of scheduled appointment (DD/MM/YYYY) / /

7d. If NO, why did the patient not schedule an appointment (tick all that apply)

- Unable to attend because of health problems
- Family problems
- Conflict with work
- Financial problems
- Client moved or relocated
- Client lives too far away
- Other (specify)

8. If the patient does not wish to return to the clinic, specify why (tick all that apply)

- Not willing to attend
- Attend clinic closer to home
- Wait time too long
- Conflict with work
- Financial problems
- Unwilling to attend because of health problems
- Family problems
- Client will go to faith healer
- Not willing to disclose HIV status
- Referred elsewhere
- Unknown
- Other (specify)

9. Did you refer the client to any of the following (tick all that apply)

- Clinic
- Counselor
- Nutritionist
- HBC
- None

Comments

For the data use only (tick after scanning the form)

Scanned Date ___/___/_____ Name of data person

HOPE CLINIC

Counseling General Session

HOPE ID Number

Site Code

Today's Date (DD/MM/YYYY)

Interviewer number

Notes

What was the focus of the session? (tick all that apply)

- Pretest
- Hygiene
- Family planning
- Crisis
- Discordance
- Sex and sexuality

- Post-test
- PMTCT
- Child(ren)
- Opportunistic infection
- Welfare
- Drug therapy

- HIV/STD Prevention
- Nutrition
- Bereavement
- Spirituality
- Disclosure
- Child transfer to Adult Clinic
- Other (specify)

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

HOPE CLINIC

COUNSELOR SCREENING

HOPE ID Number

Site Code

Today's date (DD.MM.YYYY)

Interviewer number

1. How many children do you have or care for? If = 0 go to 2

1a. Describe HIV test results and HIV care for each child.

	Test Result	Receiving HIV care?	Receiving HAART?	Receiving HAART and/or HIV care at Hope Clinic?
1	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> Not tested	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A
2	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> Not tested	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A
3	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> Not tested	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A
4	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> not tested	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> Unknown <input type="checkbox"/> No <input type="checkbox"/> N/A

2. Have you revealed your serostatus to:

- a. Spouse(s) or steady partner(s) All Some None Has no spouse
- b. Casual or non-casual partner(s) All Some None Has no partner
- c. Parents All Some None Has no parents
- d. Siblings All Some None Has no siblings
- e. Children All Some None Has no children
- f. Friends All Some None Has no friends
- g. Others All Some No one else Specify

3. Who forms your closest social support (tick one)

- Spouse/steady partner Sibling
- Mother Other (specify)
- Father No social supporter
- Friend

3a. Have you informed this person of your serostatus?

- Yes No N/A

4. Have you ever had sex? Yes No (If YES, go to 5, If NO go to 13)

5. How many spouse(s) or steady partner(s) do you have? (If = 0 go to 6)

5a. Describe HIV test results and HIV care for each spouse(s) or steady partner(s)

	Relationship	Test Result	Receiving HIV care?	Receiving HAART?	Receiving HAART and/or HIV care at Hope Clinic?
1	<input type="checkbox"/> Spouse <input type="checkbox"/> Steady partner	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> not tested	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A
2	<input type="checkbox"/> Spouse <input type="checkbox"/> Steady partner	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> Not tested	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A
3	<input type="checkbox"/> Spouse <input type="checkbox"/> Steady partner	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> Not tested	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A
4	<input type="checkbox"/> Spouse <input type="checkbox"/> Steady partner	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> not tested	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A
5	<input type="checkbox"/> Spouse <input type="checkbox"/> Steady partner	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> Not tested	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A
6	<input type="checkbox"/> Spouse <input type="checkbox"/> Steady partner	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Unknown <input type="checkbox"/> Not tested	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> N/A

6. How many different casual or non-steady partner(s) did you have in the past year?
7. In the past year, how many different sexual partners have you had, including your spouse(s) and steady partner(s)?
8. During your lifetime have you had sex with
 Men Only Women Only Both None Refused to answer
9. Describe your condom use in the past 12 months todate:-
 a.Spouse(s) or steady partiners *(Tick one)*
 Never Sometimes Always No sex in past 12 months No spouse/steady partner
 b.Non- steady partiners. *(Tick one)*
 Never Sometimes Always No sex in past 12 months No Non-steady partner
10. Did you use a condom during your last sexual encounter
 No Yes Refused to answer
11. Are you able to talk about using condoms with your spouse(s) or steady partner(s)?
 No Yes Refused to answer No spouse/steady partner Dont know
- 11a. Are you able to say NO to sex if your spouse or steady partner will not use a condom?
 No Yes No spouse/steady partner Refused to answer Don't know
12. Are you able to talk about condoms with your casual or non-steady partner(s)?
 Yes No No non-steady partner(s) Refused to answer Don't know
- 12a. Are you able to say NO to sex if your casual or non-steady partner(s) will not use a condom?
 No Yes No non-steady partner(s) Refused to answer Don't know
13. Are you Circumcised ?*(Ask Male client only)*
 Yes No Refused to answer Female client
14. Do you feel neglected by anyone *(tick all that apply)*
 Family Friends Health care workers None Others (specify)
15. Counselor, does the client need a treatment supporter? Yes No *(If yes go to 15a, if no go 16)*
- 15a. If YES why? specify (Tick all that apply)
 Client requests treatment supporter
 Physically disabled
 Mentally disabled
 Other (specify)

16. Counselor is this client recommended for homecare? Yes No

16a. If YES, specify why? (Tick all that apply)

- Physically or mentally disabled adult
- Adult dependent on care-giver
- Client requests a home visit
- Disclosure (patient would like help disclosing status to family members)
- Family testing (client would like other family members to be tested)
- Other (specify)

17. Have you explained/discussed or checked the following with the Client?

Checklist	Tick (if Yes)	Tick (if No)
1. Overview of HOPE Center Program and Services	<input type="radio"/>	<input type="radio"/>
2. Importance of commitment to the program	<input type="radio"/>	<input type="radio"/>
3. Policy for adherence and clinic attendance	<input type="radio"/>	<input type="radio"/>
4. Patients enrollment status in other programs or facilities	<input type="radio"/>	<input type="radio"/>
5. Patients long term goals for health management at our program	<input type="radio"/>	<input type="radio"/>

Notes / Assessment

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

HOPE CLINIC ADULT LOCATOR FORM

HOPE ID Number

Site Code

Today's Date (DD/MM/YYYY)

//

Interviewer number

1. Name (Last, First, Middle)

2. Gender Male Female

3. Age (Years)

CLIENT'S RESIDENTIAL AND TELEPHONE CONTACT INFORMATION

4. Public Transportation to the house:

4a. Type

Bus Citi Hoppa Matatu Taxi Other(specify)

4b. Number

4c. Stage Name

4d. General name of the area

5. Walking directions to house from the stage?

6. Landmarks that aide in locating the household: (Schools, churches, businesses etc.)

7. How is the Client or Caregiver called or referred to in home area:

8. Primary Telephone Contact:

Mobile Landline Unknown

8a. Line belongs to:

- | | |
|---|---|
| <input type="checkbox"/> Client | <input type="checkbox"/> Friend |
| <input type="checkbox"/> Parent | <input type="checkbox"/> Neighbour |
| <input type="checkbox"/> Caregiver | <input type="checkbox"/> Nerby Simu ya Jamii |
| <input type="checkbox"/> Other household member | <input type="checkbox"/> Guardian Institution or organisation |
| <input type="checkbox"/> Relative | <input type="checkbox"/> Other(specify) |

8b. If the phone is not the client's does the owner know of the client's status?

- Yes
 No
 Unknown

9. Secondary Telephone Contact:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

- Mobile
 Landline
 Unknown

9a. Line belongs to:

- | | |
|---|---|
| <input type="checkbox"/> Client | <input type="checkbox"/> Friend |
| <input type="checkbox"/> Parent | <input type="checkbox"/> Neighbour |
| <input type="checkbox"/> Caregiver | <input type="checkbox"/> Nearby Simu ya Jamii |
| <input type="checkbox"/> Other household member | <input type="checkbox"/> Guardian Institution or organisation |
| <input type="checkbox"/> Relative | |
| <input type="checkbox"/> Other(specify) | |

9b. If the phone is not the client's does the owner know of the client's status?

- Yes
 No
 Unknown

10. How long has the client been living at this residence:

Years

 Months

10a. This residence is:

- Permanent
 Temporary
 Unknown

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

HOPE CLINIC

ADULT MEDICAL FOLLOW-UP

HOPE ID Number

Site Code

Today's Date (DD.MM.YYYY)

/

/

Interviewer number

HISTORY OF PRESENT ILLNESS

MEDICAL REVIEW

1. Does the patient currently have Extra pulmonary TB? Yes No

2. Does the Patient have Pulmonary TB? Yes No *(If YES go to 2a, If NO go to 4)*

2a. If YES, what was the diagnosis based on (tick all that apply)

- Chest X-ray
- Sypmtoms (persistent cough >2 weeks, fever, night sweats, etc)
- Failure to respond to empirical antibiotics
- Sputum
- Other (specify)
- Unknown

3. Is the patient currently on treatment for TB(PTB & ETB) *(If YES go to 3a, If NO or Unknown go to 4)*
 Yes No Unknown

3a. If yes, specify treatment start date (DD.MM.YYYY) / /

3b. If YES, specify treatment

- EH SHRZE
- RHZ unknown
- RHZE Other (specify)
- RH

S=streptomycine
 H=Isoniazid
 R=Rifampicin
 Z=Pyrazinamid
 E=Ethambutol

3c. Is the patient currently being treated for TB at coptic? Yes No *(If YES go to 4, If NO go to 3d)*

3d. If NO, specify where:

- Private hospital
- Public hospital
- Other government facility
- Other (specify)

4. In the past month, has the patient experienced any of the following? (tick all that apply)

- Dysparenuia Testicular pain or swelling
- Genital sores or ulcers Urethral discharge
- Lower Abdominal pain Vaginal discharge
- Painful micturation(Dysuria) Vaginal itching/burning
- None

MEDICATIONS

5. Is the patient currently taking HAART or ARVs, excluding PMTCT & PEP? Yes No (If YES go to 5a, If NO go to 7)

5a. If yes, specify (tick one)

- | | |
|--|---|
| <input type="checkbox"/> AZT-3TC-EFV | <input type="checkbox"/> d4T(40mg)-DDI-LPV/rit |
| <input type="checkbox"/> AZT-3TC-NVP | <input type="checkbox"/> TDF 3TC EFV |
| <input type="checkbox"/> AZT-3TC-LPV/rit | <input type="checkbox"/> TDF 3TC NVP |
| <input type="checkbox"/> d4T(30mg)-3TC-EFV | <input type="checkbox"/> TDF-ABC-LPV/rit |
| <input type="checkbox"/> d4T(30mg)-3TC-NVP | <input type="checkbox"/> Unknown |
| <input type="checkbox"/> d4T(30mg)-DDI-LPV/rit | <input type="checkbox"/> Other (specify) <input type="text"/> |

5b. Has the patient had any recent side effects due to HAART or ARV medications? Yes No (If Yes go to 5c, If NO go to 6)

5c. If YES, describe the symptoms and severity of possible side effects (Tick for each symptom)

Symptom	Frequency of Symptom	If YES, severity of symptom
a. Nausea or vomiting	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
b. Rash	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
c. Fat changes	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
d. Diarrhea	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
e. Anemia	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
f. Cough	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
g. Fatigue	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
h. Abdominal pain	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
i. CNS - dizziness, anxiety, nightmares	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
j. Headache	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
k. Jaundice	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
l. Difficulty breathing	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
m. Burning/numbness/tingling	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
n. Fever	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
o. Heartburn	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe
p. Other (specify) <input type="text"/>	<input type="checkbox"/> Sometimes <input type="checkbox"/> Often	<input type="checkbox"/> Mild <input type="checkbox"/> Moderate <input type="checkbox"/> Severe

6. During the last 7 days how many ARV pills did the patient MISS taking? (tick one)

- None Very Few Half Most All

6a. If patient MISSED doses, please specify reasons (tick all that apply).

- Alcohol Forgot Too ill
 Depression Inability to pay Toxicity/side effect
 Drug stock out - dispensary Patient lost or ran out of pills Other (specify)
 Delivery/travel problems Stigma, disclosure or privacy issues
 Felt better Share with others

7. Is patient taking any of the following medications? (tick all that apply)

- Cotrimoxazole Antimalarial medications None of the above
 Dapsone Herbal traditional medications
 Fluconazole Multivitamin supplements

8. What other medications is patient currently taking?

Medication	Dose (mg)	Frequency (per day)	Start date	Stop date
1.				
2.				
3.				

9. PHYSICAL EXAMINATION

Nurse number

Temp (F) HR BP / RR Wt (Kg) HGT Sa O2

BMI BMI > 18.5 BMI < 18.5 BMI Unknown Gender MALE Female

10. System (tick one)	Normal	Abnormal	Not done	Findings (if abnormal)
General	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Skin	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Lymph nodes	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
HEENT	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Lungs	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Cardiovascular	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Abdomen	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Genitourinary	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Extremities	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Musculoskeletal	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Neurological	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Psychological	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

ASSESSMENT AND PLAN

11. What diagnoses does the patient have on today's visit (tick all that apply)?

- | | | | |
|---|--|--|--|
| <input type="checkbox"/> Anaemia | <input type="checkbox"/> Dermatitis | <input type="checkbox"/> Myalgia | <input type="checkbox"/> URTI |
| <input type="checkbox"/> Asthma | <input type="checkbox"/> Extra Pulmonary TB | <input type="checkbox"/> Neuropathy | <input type="checkbox"/> UTI |
| <input type="checkbox"/> Candidiasis (thrush) - oral | <input type="checkbox"/> Gonorrhoea | <input type="checkbox"/> Peptic Ulcer disease | <input type="checkbox"/> Zoster |
| <input type="checkbox"/> Candidiasis (thrush) - vaginal | <input type="checkbox"/> Genital Ulcer disease | <input type="checkbox"/> Pneumonia | <input type="checkbox"/> None |
| <input type="checkbox"/> Chancroid | <input type="checkbox"/> Hypertension | <input type="checkbox"/> Side effects due to ARV | <input type="checkbox"/> Other (specify) |
| <input type="checkbox"/> Conjunctivitis | <input type="checkbox"/> HSV - genital | <input type="checkbox"/> Soft tissue infection | <input type="text"/> |
| <input type="checkbox"/> Diarrhea | <input type="checkbox"/> IRIS | <input type="checkbox"/> Syphilis | |
| <input type="checkbox"/> Dementia | <input type="checkbox"/> Malaria | <input type="checkbox"/> Ulcers - oral | |

12. What is the patients current Pulmonary TB diagnosis?

- Pulmonary TB suspected *(If suspected go to 13)*
- Pulmonary TB diagnosed today *(If diagnosed go to 14)*
- Currently on pulmonary TB treatment *(If currently on TB treatment go to 15)*
- Previously diagnosed with TB, not on Treatment *(skip to 14a)*
- No Pulmonary TB *(If NO TB go to 15)*
- Other(specify)

13. If pulmonary TB suspected, what is the suspected pulmonary TB based on (tick all that apply)?

- Abnormal X-ray
- Symptoms (persistent cough >2 weeks, fever, night sweats, etc)
- Failure to respond to empirical antibiotics
- Recent contact with people with Pulmonary TB
- Other (specify)

13a. Will the patient be sent for pulmonary TB testing? Yes NO *(If yes, go to 13b, if no go to 15)*

13b. If YES, which of the following tests will the patient be sent for:

- Sent for sputum
- Sent for X-ray
- Other (specify)
- (Go to 15)*

14. If Pulmonary TB is diagnosed today, what is it based on (tick all that apply)?

- Abnormal X-ray Failure to respond to empirical antibiotics
- Symptoms (persistent cough >2 weeks, fever, night sweats, etc) Sputum
- Other (specify)

14a. Specify treatment to be started: S=streptomycine

- | | | |
|--|--------------------------------|---------------|
| <input type="checkbox"/> RHZE | <input type="checkbox"/> SHRZE | H=Isoniazid |
| | | R=Rifampicin |
| <input type="checkbox"/> RHZ | <input type="checkbox"/> EH | Z=Pyrazinamid |
| | | E=Ethambutol |
| <input type="checkbox"/> Other (specify) | <input type="text"/> | |

Section B: Female Patients Only (If Male go to 19)

15. Is the patient currently pregnant?
 Yes No Unknown *(If YES, complete Pregnancy Monitoring Form)*
16. Is the patient currently breastfeeding? Yes No *(If YES, go to 16a, if no go to 17)*
 16a. If yes, specify what type of breastfeeding (tick one)
 Exclusive breastfeeding *(Child given ONLY mother's milk and NO water, tea, formula, cow's milk or food of any kind)*
 Mixed feeding *(Child given mother's milk and water, tea formula, cow's milk or food)*
- 16b. If YES, has she had a session with the nutritionist since she started breast feeding?
 Yes No ***(If NO, refer patient to Nutritionist)***
17. Has the patient delivered in the past 18 Months? Yes No *(If yes go to 17a, If No go to 18)*
 17a. If yes, is this the first time the patient has returned to the Hope Center since delivery?
 Yes ***(If yes, complete pregnancy Close -Out form)***
 No
- 17b. Age of the child Months Days Child not Alive
- 17c. If not currently breastfeeding, at what age did the child stop breastfeeding?
 Months Days Unknown NA, Child never breastfed
- 17d. Has the child ever had a PCR test? Yes No
- 17e. If Yes, what was the result of the last PCR test?
 HIV Positive *(Go to 18)*
 HIV Negative
 Indeterminate
 Result not yet available
(Complete Infant PCR form; If Q17e is HIV Negative, Indeterminate or Result not yet available)
- 17f. Will an Infant PCR test be ordered today? Yes No
18. Is the patient being referred for Cervical Cancer screening at Hope? Yes No *(If yes go to 19, If NO go to 18a)*
 18a. If NO, reasons why patient NOT referred for Cervical Cancer Screening *(tick all that apply)*
 Patient is younger than 18 years
 Patient has had total hysterectomy, LEEP or cryotherapy
 Patient is currently pregnant
 Patient has had a screening test in the last year
 No service available at this time
 Patient wishes to defer until a later time Specify reason
 Patient does not accept screening Specify reason
 Other(specify)

19. Is the Client currently on HAART, excluding PMTCT and PEP? Yes No *(If YES go to 19a, If No go to 19b)*

19a. If yes, did you continue current HAART? Yes No *(If YES go to 20, If No go to 19a1)*

19a1. If no, did you change or stop HAART today? Changed Stopped

19a2. Specify why HAART was changed or stopped *(Tick all that apply)*

- Toxicity / Side effects
- Clinical treatment failure
- Pregnancy
- Immunologic treatment failure
- Risk of pregnancy
- Virologic treatment failure
- Newly diagnosed TB
- Poor adherence
- New drug available
- Illness, hospitalization
- Drug not available
- Other (specify)
- Patient lacks finances

(Skip to 20)

19b. If no, is patient ELIGIBLE for therapy? Yes No Not yet determined *(If Eligible go to 19b1; if NOT eligible or NYD go to 22)*

19b1. If ELIGIBLE for therapy then by what criteria *(tick all that apply)*?

- CD4 count CD4 % Date (DD/MM/YYYY) / /
- WHO Clinical Stage 1 2 3 4 Viral Load
- Patient on HAART/ARV in past

19c. Did you initiate HAART/ARV treatment at this clinic visit, excluding PMTCT and PEP? Yes No *(If YES go to 19c1 If No go to 19c2)*

19c1. If yes, what was the WHO stage 1 2 3 4

19c2. If NO, specify:

- Patient has not completed HAART protocol
- Patient preference
- Patient currently on drugs which may interact with HAART/ARV
- Patient too ill to begin HAART today
- Other (specify)

20. Has the patient completed HAART protocol? Yes No

21. What ARV medications were prescribed or continued today

- AZT-3TC-EFV
- TDF-ABC -LPV/rit
- AZT-3TC-NVP
- TDF 3TC EFV
- AZT-3TC- LPV/rit
- TDF 3TC NVP
- d4T(30mg)-3TC-EFV
- None
- d4T(30mg)-DDI-LPV/rit
- Other (specify)
- d4T(30mg)-3TC-NVP

22. Did you initiate or continue Cotrimoxazole today? Yes No *(If YES go to 23, If NO go to 22a)*

22a. If no, why?

- Side effects/ toxicity Patient preference
 Stockout/drug supply interruption Other (Specify)

23. Did you initiate or continue any of the following medications today (tick all that apply)?

- Dapsone Fluconazole Multivitamin supplement None of the above

24. Other medications prescribed during this visit

Medication	Dose (mg)	Frequency (per day)	Start date	
1.				
2.				
3.				
4.				

25. What laboratory tests were ordered today (tick all that apply)

- ALT Hgb Viral Load
 cd4 HIV ELISA Confirmatory test Widal Test
 Chest X-ray LFT None
 Creatinine Lactic Test Other (specify)
 Complete Blood Count Urinalysis

Comments

For the data use only (tick after scanning the form)

Scanned Date ___/___/_____ Name of data person

HOPE CLINIC ADULT MEDICAL SCREENING

HOPE ID Number

Site Code Today's date (DD.MM.YYYY) Interviewer number

 /

 /

HISTORY OF PRESENT ILLNESS

PAST MEDICAL HISTORY

1. Has the patient ever had pulmonary TB in the past?

Yes No Unknown *(If YES go to 1a, If NO or unknown go to 2)?*

1a. If YES, what was the diagnosis based on *(tick all that apply)*

- Chest X-ray
- Sypmtoms (persistent cough >2 weeks, fever, night sweats, etc)
- Failure to respond to empirical antibiotics
- Sputum
- Other (specify)
- Unknown

1b. Was the patient treated

Yes No Unknown *(If YES go to 1c, If NO or unknown go to 2)*

1c. If YES, specify treatment

- EH SHRZE
- RHZ unknown
- RHZE Other (specify)
- RH

S=streptomycine
H=Isoniazid
R=Rifampicin
Z=Pyrazinamid
E=Ethambutol

1d. If YES, did the patient (tick one)?

- Never completed full treatment
- Doesn't know if received full treatment
- Completed Full Treatment

Date started (DD/MM/YYYY) / /

Date stopped (DD/MM/YYYY) / /

2. Does the patient currently have Extra pulmonary TB? Yes No

3. Does the Patient have Pulmonary TB? Yes No *(If YES go to 3a, If NO go to 4)*

3a. If YES, what was the diagnosis based on (tick all that apply)

- Chest X-ray
- Symptoms (persistent cough >2 weeks, fever, night sweats, etc)
- Failure to respond to empirical antibiotics
- Sputum
- Other (specify)
- Unknown

4. Is the patient currently on treatment for TB (either PTB or EPTB) *(If YES go to 4a If NO or Unknown go to 5)*

Yes No Unknown

4a. If yes, specify treatment start date (DD.MM.YYYY) / /

4b. If YES, specify treatment

- EH SHRZE
- RHZ unknown
- RHZE Other (specify)
- RH

S=streptomycine
H=Isoniazid
R=Rifampicin
Z=Pyrazinamid
E=Ethambutol

4c. Is the patient currently being treated for TB at coptic? Yes No *(If YES go to 5, If NO go to 4d)*

4d. If NO, specify where:

- Private hospital
- Public hospital
- Other government facility
- Other (specify)

5. In the past month, has the patient experienced any of the following? (tick all that apply)

- Dyspareunia Vaginal itching/burning
- Genital sores or ulcers Vaginal discharge
- Lower Abdominal pain Urethral discharge
- Painful micturation(Dysuria) None
- Testicular pain or swelling

6. Has the patient ever had or been told he/she had a sexually transmitted infection? Yes No Unknown

6a. If yes, specify (tick all that apply)

- Chlamydia Trichomonas vaginalis
- Chancroid Syphilis
- Herpes Unknown
- Neisseria gonorrhoea Other specify

7. Does the patient currently have, or has the patient ever had, any of the following conditions (tick all that apply)

WHO stage 1	
Asymptomatic HIV infection	<input type="checkbox"/>
Persistent generalized lymphadenopathy (PGL)	<input type="checkbox"/>
WHO stage 2	
Herpes Zoster (within last 5 years)	<input type="checkbox"/>
Minor Mucocutaneous Manifestations	<input type="checkbox"/>
Recurrent Upper Respiratory Infections	<input type="checkbox"/>
Weight loss < 10% of Body weight	<input type="checkbox"/>
WHO stage 3	
Severe Bacterial infections (i.e Pneumonia, pyomyositis)	<input type="checkbox"/>
Oral Candidiasis (Thrush)	<input type="checkbox"/>
Unexplained chronic diarrhea (> 1 month)	<input type="checkbox"/>
Unexplained Prolonged Fever (intermittent or constant, > 1 month)	<input type="checkbox"/>
Oral Hairy Leukoplakia	<input type="checkbox"/>
Tuberculosis, Pulmonary (within last 12 months from today)	<input type="checkbox"/>
Weight loss > 10% of body weight	<input type="checkbox"/>
WHO stage 4	
Candidiasis (Esophageal, Bronchi, Trachea, or lungs)	<input type="checkbox"/>
Cryptococcosis, Extrapulmonary	<input type="checkbox"/>
Cryptosporidiosis with Diarrhea (> 1 month duration)	<input type="checkbox"/>
Herpes Simplex (mucocutaneous > 1 months, or visceral or any duration)	<input type="checkbox"/>
HIV Encephalopathy	<input type="checkbox"/>
HIV Wasting Syndrom	<input type="checkbox"/>
Kaposi's Sarcoma (KS)	<input type="checkbox"/>
Lymphoma	<input type="checkbox"/>
Atypical Mycobacteriosis, Disseminated	<input type="checkbox"/>
Tuberculosis, Extrapulmonary	<input type="checkbox"/>
Progressive Multifocal Leukoencephalopathy (PML)	<input type="checkbox"/>
Mycosis, disseminated endemic (i.e., histoplasmosis, coccidiomycosis)	<input type="checkbox"/>
Pneumocystis Carinii Pneumonia (PCP)	<input type="checkbox"/>
Salmonella Septicemia, Non-typhoid	<input type="checkbox"/>
Toxoplasmosis, CNS	<input type="checkbox"/>

8. What is the WHO Clinical Stage of the patient? (tick one) 1 2 3 4

9. Last CD4 count Unknown Not tested
- 9a. Last CD4 count date (DD.MM.YYYY) / / Unknown
- 9b. Last Viral load undetectable Unknown Not tested
- 9c. Last viral load date (DD.MM.YYYY) / / Unknown

MEDICATIONS

10. Has the patient ever taken or is the patient presently on HAART excluding for the purpose of PMTCT and PEP?

- Yes No *(If YES go to 10a, if NO go to 11)*

10a. If YES, specify history below

	Medication (tick all that apply)										
	d4T	3TC	AZT	DDI	ABC	NVP	EFV	LPV/rit	TDF	Truvada	Other (Specify)
First regimen	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="text"/>
Second regimen	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="text"/>
Third regimen	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="text"/>

	Generic or brand (tick one)	CD4 count at time of regimen initiation (if known)	Adherence (1=V. poor, 2=poor, 3=Fair, 4= Good, 5=Excellent)	Date started (DD/MM/YYYY)	Did patient stop?	If yes, Date stopped (DD/MM/YYYY)
First regimen	<input type="checkbox"/> Generic <input type="checkbox"/> Brand <input type="checkbox"/> Unknown	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="checkbox"/> Unknown	<input type="checkbox"/>	<input type="text"/> / <input type="text"/> / <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="text"/> / <input type="text"/> / <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>
Second regimen	<input type="checkbox"/> Generic <input type="checkbox"/> Brand <input type="checkbox"/> Unknown	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="checkbox"/> Unknown	<input type="checkbox"/>	<input type="text"/> / <input type="text"/> / <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="text"/> / <input type="text"/> / <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>
Third regimen	<input type="checkbox"/> Generic <input type="checkbox"/> Brand <input type="checkbox"/> Unknown	<input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="checkbox"/> Unknown	<input type="checkbox"/>	<input type="text"/> / <input type="text"/> / <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>	<input type="checkbox"/> Yes <input type="checkbox"/> No	<input type="text"/> / <input type="text"/> / <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/>

If stopped why? (tick all that apply)

	Costs	Side effects	Failure of therapy	New diagnosis TB	Drug out of stock	Doctor orders	Unknown	Other (Specify)
First regimen	<input type="checkbox"/>	<input type="text"/>						
Second regimen	<input type="checkbox"/>	<input type="text"/>						
Third regimen	<input type="checkbox"/>	<input type="text"/>						

11. Is patient taking any of the following medications? (tick all that apply)

- Antimalarial medications
 Dapsone
 Herbal traditional medications
 None of the above
 Cotrimoxazole
 Fluconazole
 Multivitamin supplements

12. What other medications is patient currently taking?

Medication	Dose (mg)	Frequency (per day)	Start date	Stop date
1.				
2.				
3.				

13. PHYSICAL EXAMINATION

Nurse number

Temp (F) .
 HR
 BP /
 RR
 Wt (Kg) .
 Height (cm)
 Sa O2

BMI BMI > 18.5
 BMI < 18.5
 BMI Unknown
 Gender MALE
 Female

14.

System (tick one)	Normal	Abnormal	Not done	Findings (if abnormal)
General	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Skin	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Lymph nodes	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
HEENT	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Lungs	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Cardiovascular	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Abdomen	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Genitourinary	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Extremities	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Musculoskeletal	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Neurological	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	
Psychological	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	

ASSESSMENT AND PLAN

15. What new diagnoses does the patient have on today's visit (tick all that apply)?

- | | | | |
|---|--|--|--|
| <input type="checkbox"/> Anaemia | <input type="checkbox"/> Dermatitis | <input type="checkbox"/> Malaria | <input type="checkbox"/> Syphilis |
| <input type="checkbox"/> Asthma | <input type="checkbox"/> Extra Pulmonary TB | <input type="checkbox"/> Myalgia | <input type="checkbox"/> Ulcers genital |
| <input type="checkbox"/> Candidiasis (thrush) - oral | <input type="checkbox"/> Gonorrhoea | <input type="checkbox"/> Neuropathy | <input type="checkbox"/> Ulcers - oral |
| <input type="checkbox"/> Candidiasis (thrush) - vaginal | <input type="checkbox"/> Genital Ulcer disease | <input type="checkbox"/> Peptic Ulcer disease | <input type="checkbox"/> URTI |
| <input type="checkbox"/> Chancroid | <input type="checkbox"/> Hypertension | <input type="checkbox"/> Pneumonia | <input type="checkbox"/> UTI |
| <input type="checkbox"/> Conjunctivitis | <input type="checkbox"/> HSV - genital | <input type="checkbox"/> Side effects due to ARV | <input type="checkbox"/> Zoster |
| <input type="checkbox"/> Diarrhea | <input type="checkbox"/> IRIS | <input type="checkbox"/> Soft tissue infection | <input type="checkbox"/> None |
| <input type="checkbox"/> Dementia | | | <input type="checkbox"/> Other (specify) |

16. What is the patients current TB diagnosis?

- TB suspected *(If suspected go to 17)*
- TB diagnosed today *(If diagnosed go to 18)*
- Currently on pulmonary TB treatment *(If currently on TB treatment go to 19)*
- No TB *(If NO TB go to 19)*
- Other(specify)

17. If TB suspected, what is the suspected TB based on (tick all that apply)?

- Abnormal X-ray
- Symptoms (persistent cough >2 weeks, fever, night sweats, etc)
- Failure to respond to empirical antibiotics
- Recent contact with people with TB
- Other (specify)

17a. Will the patient be sent for TB testing? Yes NO *(If YES, go to 17b, If NO go to 19)*

17b. If YES. which of the following tests will the patient be sent for:

- Sent for sputum
- Sent for X-ray
- Other (specify)

(Go to 19)

18. If TB is diagnosed today, what is it based on(tick all that apply)?

- Abnormal X-ray
- Symptoms (persistent cough >2 weeks, fever, night sweats, etc)
- Failure to respond to empirical antibiotics
- Sputum
- Other (specify)

18a. Specify treatment to be started:

- EH SHRZE
- RHZE Other (specify)
- RHZ

SECTION B: FEMALE PATIENTS ONLY

19. In the past, has the patient ever taken any antiretroviral drugs for PMTCT?

Yes No Unknown (If YES go to 19a, If NO or Unknown go to 20)

19a. If YES, how many times did the patient take PMTCT?

19b. Specify history for each pregnancy in which the patient had PMTCT, beginning with the youngest child:

	Date of delivery (DD/MM/YYYY)	Regimen	Specify when the drug was taken (Tick all that apply)	Location of delivery	Mode of delivery
1.	<input type="text"/> / <input type="text"/> / <input type="text"/>	<input type="checkbox"/> AZT	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum	<input type="checkbox"/> Hospital <input type="checkbox"/> Home <input type="checkbox"/> Other (specify) <input type="text"/>	<input type="checkbox"/> C-section <input type="checkbox"/> Unassisted vaginal <input type="checkbox"/> Assisted vaginal (use of forceps or vacuum)
		<input type="checkbox"/> 3TC	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> NVP	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> HAART	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> Other (Specify) <input type="text"/>	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> Unknown			
2.	<input type="text"/> / <input type="text"/> / <input type="text"/>	<input type="checkbox"/> AZT	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum	<input type="checkbox"/> Hospital <input type="checkbox"/> Home <input type="checkbox"/> Other (specify) <input type="text"/>	<input type="checkbox"/> C-section <input type="checkbox"/> Unassisted vaginal <input type="checkbox"/> Assisted vaginal (use of forceps or vacuum)
		<input type="checkbox"/> 3TC	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> NVP	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> HAART	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> Other (Specify) <input type="text"/>	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> Unknown			

	Date of delivery (DD/MM/YYYY)	Regimen	Specify when the drug was taken (Tick all that apply)	Location of delivery	Mode of delivery
3.	<input type="text"/> / <input type="text"/> / <input type="text"/>	<input type="checkbox"/> AZT	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum	<input type="checkbox"/> Hospital <input type="checkbox"/> Home <input type="checkbox"/> Other (specify) <input type="text"/>	<input type="checkbox"/> C-section <input type="checkbox"/> Unassisted vaginal <input type="checkbox"/> Assisted vaginal (use of forceps or vacuum)
		<input type="checkbox"/> 3TC	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> NVP	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> HAART	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> Other (Specify) <input type="text"/>	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> Unknown			
4.	<input type="text"/> / <input type="text"/> / <input type="text"/>	<input type="checkbox"/> AZT	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum	<input type="checkbox"/> Hospital <input type="checkbox"/> Home <input type="checkbox"/> Other (specify) <input type="text"/>	<input type="checkbox"/> C-section <input type="checkbox"/> Unassisted vaginal <input type="checkbox"/> Assisted vaginal (use of forceps or vacuum)
		<input type="checkbox"/> 3TC	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> NVP	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> HAART	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> Other (Specify) <input type="text"/>	<input type="checkbox"/> Antepartum <input type="checkbox"/> Labor & Delivery <input type="checkbox"/> Postpartum		
		<input type="checkbox"/> Unknown			

20. Is the patient currently pregnant?

Yes No Unknown *(If YES also complete Pregnancy Monitoring Form)*

21. Is the patient currently breastfeeding? Yes No *(If YES go to 21a, if NO go to 22)*

21a. If yes, specify what type of breastfeeding (tick one)

Exclusive breastfeeding *(Child given ONLY mother's milk and NO water, tea, formula, cow's milk or food of any kind)*

Mixed feeding *(Child given mother's milk and water, tea formula, cow's milk or food)*

21b. If YES, has she had a session with the nutritionist since she started breastfeeding.

Yes No ***(If NO refer to Nutritionist)***

22. Is the patient currently on HAART, excluding PMTCT and PEP? Yes No *(If YES, go to 22a, If NO go to 22b)*

22a. If yes, did you continue current HAART today? Yes No *(If YES, go to 23, If NO go to 22a1)*

22a1. If no, did you change or stop HAART today? Changed Stopped

22a2. Specify why HAART was changed or stopped (Tick all that apply) :

Toxicity / Side effects

Clinical treatment failure

Pregnancy

Immunologic treatment failure

Risk of pregnancy

Virologic treatment failure

Newly diagnosed TB

Poor adherence

New drug available

Planned treatment interruption

Drug not available

Illness, hospitalization

Patient lacks finances

(Go To 23)

Other (specify)

22b. If NO, is patient ELIGIBLE for therapy? Yes No Not yet determined *(If Eligible=NO or NYD >>24)*

22b1. If ELIGIBLE for therapy then by what criteria (tick all that apply)?

CD4 count

Date (DD/MM/YYYY) / /

WHO Clinical Stage 1 2 3 4

Viral Load copies

Patient on HAART/ARV in past

22c. Did you initiate HAART/ARV treatment at this clinic visit, excluding PMTCT and PEP? Yes No

(If YES go to 23, If NO go to 22c1)

22c1. If NO, specify:

Patient has not completed HAART protocol

Patient preference

Patient currently on drugs which may interact with HAART/ARV

Patient pregnant

Patient too ill to begin HAART today

Other (specify)

23. What ARV medications were prescribed or continued today

AZT-3TC-EFV

TDF-ABC -LPV/rit

AZT-3TC-NVP

TDF 3TC EFV

AZT-3TC- LPV/rit

TDF 3TC NVP

d4T(30mg)-3TC-EFV

None

d4T(30mg)-DDL-LPV/rit

Other (specify)

d4T(30mg)-3TC-NVP

24. Did you initiate or continue Cotrimoxazole today? Yes No *(If yes go to 25, if no go to 24a)*

24a. If no, why?

Side effects/ toxicity

Patient preference

Stockout/drug supply interruption

Other (Specify)

25. Did you initiate or continue any of the following medications today (tick all that apply)?

Dapsone

Fluconazole

Multivitamin supplement

None of the above

26. Other medications prescribed during this visit

	Medication	Dose (mg)	Frequency (per day)	Start date	Stop date
1.					
2.					
3.					
4.					
5.					

27. What laboratory tests were ordered today (tick all that apply)

ALT

Hgb

Viral Load

cd4

HIV ELISA Confirmatory test

Widal Test

Chest X-ray

LFT

None

Creatinine

Lactic Test

Other (specify)

Complete Blood Count

Urinalysis

Additional Comments

For the data use only (tick after scanning the form)

Scanned

Date ___/___/_____

Name of data person

HOPE CLINIC VERBAL AUTOPSY FORM

HOPE ID Number

Site Code

Today's Date (DD.MM.YYYY)

Interviewer number

1. Gender Female Male

2. Age at death Years

3. Date of death (DD/MM/YYYY) / /

4. The information source for the cause of death was (Tick all that apply)

- | | |
|---|---|
| <input type="checkbox"/> Hospital records/Staff | <input type="checkbox"/> Mother or Father |
| <input type="checkbox"/> Partner/Spouse | <input type="checkbox"/> Other family member/relative |
| <input type="checkbox"/> Neighbour | <input type="checkbox"/> Caregiver |
| <input type="checkbox"/> Friend | <input type="checkbox"/> Unknown |
| <input type="checkbox"/> Other (Specify) <input style="width: 250px; height: 20px;" type="text"/> | |

5. Was the deceased seeking other medical treatment (other than Hope Center) during that last 3 months before his/her death?

- Yes No Unknown *(If YES go to 5a, If NO go to 6)*

5a. If YES, where specifically was the deceased receiving other medical care?

Name of facility:

5b. What type of care was the deceased receiving at these other facilities? (Tick all that apply)

- | | | |
|---|----------------------------------|--|
| <input type="checkbox"/> General medical care | <input type="checkbox"/> TB | <input type="checkbox"/> STD |
| <input type="checkbox"/> HIV/AIDS | <input type="checkbox"/> Malaria | <input type="checkbox"/> Other infectious disease(s) |
| <input type="checkbox"/> Other (Specify) <input style="width: 240px; height: 20px;" type="text"/> | | |

6. Respondent's detailed account of the illness of the deceased:

7. Did a health care worker tell you the cause of death? *(If YES go to 7a, If NO go to 8)*

- Yes No Unknown

7a. What did the health care worker say was the cause of death?

8. Did s/he have any operation for the illness? (If YES go to 8a, If NO go to 9)

- Yes No Unknown

8a. How long before the death was the operation? : Months

8b. On what part of the body was the operation?

- Abdomen Chest Head

Other(specify)

9. Has the deceased's spouse or partner died in the past in the past 5 years? (If YES go to 9a, If NO go to 10)

- Yes No Unknown Had no spouse

9a. If YES, what is the perceived cause(s) of death of the partner(s)

9a1. Partner 1:

9a2. Partner 2:

Injury/accident/suicide

10. Did s/he suffer from any injury or accident that led to her/his death? (If YES go to 10a, If NO go to 11)

- Yes No Unknown

10a. What kind of injury or accident did the deceased suffer?

- Road traffic accident Fall
 Burns Violence/assault
 Unknown Poisoning
 Drowning Other:

10b. Was the injury or accident intentionally inflicted by someone else?

- Yes No Unknown

10c. Do you think that s/he committed suicide?

- Yes No Unknown

History of previously known conditions

11. Did the deceased suffer from any of the following conditions?

- a. High blood pressure Yes No Unknown
- b. Diabetes Yes No Unknown
- c. Asthma Yes No Unknown
- d. Epilepsy Yes No Unknown
- e. Malnutrition Yes No Unknown
- f. Cancer Yes No Unknown

f1. If YES, specify type or site

g. Tuberculosis Yes No Unknown

h. Any other medically diagnosed illness? Yes No Unknown

h1. If YES, specify

12. Signs, symptoms and their severity during the last illness:

Symptom	Symptom present?	If present, duration of symptom
a. Fever	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
b. Loss of weight	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
c. Diarrhea	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
d. Vomiting/associated abdominal pain	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
e. Constipation/associate abdominal pain	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
f. Cough	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
g. Cough followed by vomiting	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
h. Breathing trouble (chest indrawing/difficult /rapid/wheezing)	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
i. Neck stiffness	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
j. Unconscious episodes	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
k. Fits	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
l. Jerking of individual limbs	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
m. History of epileptic illness in earlier years	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
n. Paralysis of limbs	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
o. Rigid body stiffness, unable to open mouth	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
p. Red and sore eyes	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
q. Skin rash and itching	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
r. Herpes zoster (at any time in life)	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
s. Abscesses/body sores	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
t. White patches on the inside of mouth and tongue	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
u. Oedema	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
v. Hair changes	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
w. Yellowing of eyes or passing of brown urine	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
x. Chest pain	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
y. Other (specify) <input style="width: 100px; height: 15px;" type="text"/>	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown

Women only:

z. unexpected vaginal bleeding or discharge	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown
aa.Pelvic or vaginal pain	<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown	<input type="checkbox"/> =<2 weeks <input type="checkbox"/> >2 weeks <input type="checkbox"/> Unknown

14. Records available in home e.g. death certificate (extract findings):

Comments (if the form is incomplete or any other comments)

For the data use only (tick after scanning the form)

Scanned Date ___/___/_____ Name of data person

HOPE CLINIC ADULT NURSING SCREENING

HOPE ID Number

Site Code

Today's date (DD.MM.YYYY)

/
/

Interviewer number

Section A: All Patients

1. Who referred patient here? (tick one)

- | | |
|--|--|
| <input type="radio"/> Hope VCT
<input type="radio"/> Other VCT
<input type="radio"/> PMTCT
<input type="radio"/> CCC Clinic or HIV Clinic
<input type="radio"/> Hope TB clinic
<input type="radio"/> Other TB clinic
<input type="radio"/> Self-referral
<input type="radio"/> Child welfare Clinic | <input type="radio"/> Family member, spouse or friend
<input type="radio"/> Other patients
<input type="radio"/> NGO
<input type="radio"/> Coptic ward
<input type="radio"/> Other hospital ward (specify) <input style="width: 150px; height: 15px;" type="text"/>
<input type="radio"/> Coptic Pharmacy
<input type="radio"/> Private doctor
<input type="radio"/> Other (specify) <input style="width: 250px; height: 15px;" type="text"/> |
|--|--|

2. Does the client have a NASCOP referral number? Yes No *(If YES go to 2a, If NO go to 3)*

2a. If Yes, specify the client's NASCOP referral number with date

NASCOP Referral Number

Referral Date (DD/MM/YYYY)

NASCOP referral number unknown

3. Has client ever been on antiretroviral drugs, excluding for PMTCT and PEP? Yes No *(If YES go to 3a, If NO go to 4)*

3a. If YES, where did the client receive antiretroviral drugs ?

- | | |
|---|---|
| <input type="radio"/> CCC Clinic or HIV Clinic
<input type="radio"/> Private Doctor
<input type="radio"/> Hospital ward | <input type="radio"/> NGO
<input type="radio"/> Other (specify) <input style="width: 250px; height: 15px;" type="text"/> |
|---|---|

3b. What is the reason for transfer of care (tick all that apply)

- | | |
|--|--|
| <input type="radio"/> Financial
<input type="radio"/> Client's preference
<input type="radio"/> Distance to clinic
<input type="radio"/> Doctors advice | <input type="radio"/> Poor management
<input type="radio"/> Client was asked to leave
<input type="radio"/> Facility unable to <input style="width: 150px; height: 15px;" type="text"/>
<input type="radio"/> Other (specify) |
|--|--|

4. In the past, has the patient ever taken any antiretroviral drugs for PMTCT?

- Yes No Unknown N/A, Male Client

4a. If YES, how many times did the patient take PMTCT?

5. Has the patient been tested for HIV? Yes No (If Yes go to 5a, If NO go to 6)

5a. If yes, where was the test performed? (Tick one) (If **PITC**, go to 5ai, else go to 5b)

- PITC (ProviderInitiated HIV Testing and Counseling)
- VCT
- PMTCT
- Postnatal Clinic
- CWC(Child welfare Clinic)
- TB clinic
- Other (Specify)

5ai. Where was the PITC done ?

- Coptic Hospital-Outpatient Coptic Hospital-inpatient
- Hope TB clinic Private Hospital
- Hope Home-base care program Public Hospital
- Muangalizi Program Other(specify)

5b. When was the test done (DD/MM/YYYY) / /

5c. What were the test results? Positive Negative Unknown

6. Has the patient been hospitalized in the last 1 year? Yes No

6a. If yes, how many times?

7. Does patient have Penicillin allergy? Yes No Dont Know

8. Does patient have Sulfa allergy? Yes No Dont Know (Medical and non-medical allergies)

9. Does the patient have anv other Yes No Dont Know

9a. If yes, please specify

(If Patient ticked any "YES" in Q7, 8 or 9 highlight allergy in chart)

Section B: Female patients only [If Male go to 12]

10. How many times has the patient been pregnant? IF = 0 go to 13

11. How many children has patient given birth to? IF = 0 go to 13

12. How many of the children the patient has given birth to are alive? IF = 0 go to 13

12a) What is the age of the first child? Years Month Weeks

12b) What is the age of the last child? Years Month Weeks

13. Is the patient or partner using any form of family planning? Yes No

13a. If YES (tick all that apply)

Condoms

Oral contraceptive pills

Injectable/implantable hormones

Diaphragm/Cervical cap

Intrauterine device

Vastectomy/tubal ligation/hysterectomy

Natural method (specify)

Other (specify)

Other Comments

Scanned Date ___/___/___ Name of data person

(Q7. applies to those initiated/or already on food supplement)

7. How would you Classify today's case? Severe Cases Moderate/Mild Cases

8. Assessment and Recommendations

For the data use only (tick after scanning the form)

Scanned Date ___/___/_____ Name of data person

HOPE CLINIC NUTRITION SCREENING

HOPE ID Number

Site Code	Today's date (DD.MM.YYYY)	Interviewer number																
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	/		/															

1. Anthropometric Assessment

Weight <table border="1" style="width: 40px; height: 20px;"></table> <table border="1" style="width: 20px; height: 20px;"></table> Kgs	Height <table border="1" style="width: 40px; height: 20px;"></table> cm	Hip Circ. <table border="1" style="width: 40px; height: 20px;"></table> cm	BMI <table border="1" style="width: 40px; height: 20px;"></table> <table border="1" style="width: 20px; height: 20px;"></table>
Waist circ. <table border="1" style="width: 40px; height: 20px;"></table> cm	Waist:Hip ratio <table border="1" style="width: 20px; height: 20px;"></table> <table border="1" style="width: 20px; height: 20px;"></table> cm	MUAC <table border="1" style="width: 40px; height: 20px;"></table> <table border="1" style="width: 20px; height: 20px;"></table> cm	

2. Medical

Nausea	<input type="radio"/> No	<input type="radio"/> Yes	Vomiting	<input type="radio"/> No	<input type="radio"/> Yes
Diarrhoea	<input type="radio"/> No	<input type="radio"/> Yes	Constipation	<input type="radio"/> No	<input type="radio"/> Yes
Weight loss greater than 10%	<input type="radio"/> No	<input type="radio"/> Yes	Chewing/teeth problem	<input type="radio"/> No	<input type="radio"/> Yes
Swallow difficulty	<input type="radio"/> No	<input type="radio"/> Yes	Taste changes	<input type="radio"/> No	<input type="radio"/> Yes
Poor appetite	<input type="radio"/> No	<input type="radio"/> Yes			
Other medical conditions	<input type="radio"/> No	<input type="radio"/> Yes			

If patient has other medical conditions, specify

1	
2	
3	
4	

Medications? Yes No (specify)

Medication	Time	with food?	Time	with food?	Time	with food?

3. Social History

Permanent housing	<input type="radio"/> No	<input type="radio"/> Yes	
Adequate food resources	<input type="radio"/> No	<input type="radio"/> Yes	
Adequate cooking facilities	<input type="radio"/> No	<input type="radio"/> Yes	
Activity/exercise	<input type="radio"/> No	<input type="radio"/> Yes	
Smoking	<input type="radio"/> No	<input type="radio"/> Yes	specify <table border="1" style="width: 150px; height: 20px;"></table>
Alcohol	<input type="radio"/> No	<input type="radio"/> Yes	specify <table border="1" style="width: 150px; height: 20px;"></table>
Drugs	<input type="radio"/> No	<input type="radio"/> Yes	specify <table border="1" style="width: 150px; height: 20px;"></table>

4. Dietary History

4a. Number of meals or snacks per day

4b. Times per week eat out

4c. Who prepares food?

- Self Relative
 Spouse Caregiver
 Child/children Other (specify)
 Neighbour

- 4d. Special or alternative diets No Yes specify
 4e. Food intolerances or allergies No Yes specify
 4f. Food likes No Yes specify
 4g. Food dislikes No Yes specify

5. Is the client currently on multivitamins? Yes NO (If YES go to 5a, If NO go to 6)

5a. If YES, did you continue the multivitamins today? Yes NO

6. Is the client on food supplement? Yes NO (If YES go to 6a, If NO go to 7)

6a. If YES, which type of food supplement?

- First Food Advantaged Foundation Other (specify)

6b. If YES, what is the qualifying criteria

- BMI < 18.5 Pregnant Breastfeeding mother Other (specify)

7. 24 hour recall /usual diet

B/Fast	M. Morning	Lunch	M. Afternoon	Dinner

8. Is the client initiating food supplements today? Yes No (If YES go to 8a, If NO go to 10)

8a. If yes, why?

- BMI < 18.5 Pregnant Breastfeeding mother Other (specify)

8b. If YES, which type of food supplement is client initiating today?

- First food Advantaged Foundation Other (specify)

8c. If yes, has the patient been on food supplement before? Yes No

(Q9. applies to those initiated/or already on food supplement)

9. How would you Classify today's case? Severe Cases Moderate/Mild Cases

10. Assessment and Recommendations

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

HOPE CLINIC
SOCIAL WORK SCREENING

HOPE ID Number

Site Code	Today's date (DD/MM/YYYY)	Interviewer number															
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	/		/														

Client's Residential and Telephone Contact Information

1. Public Transport to the House:

First trip

Ia. Type

Bus
 Citi Hoppa
 Matatu
 Taxi
 Other(specify)

Ib. Number

Ic. Stage Name

Id. General name of the area

Second trip

IIa. Type

Bus
 Citi Hoppa
 Matatu
 Taxi
 Other(specify)

IIb. Number

IIc. Stage Name

IId. General name of the area

2. Walking directions to house from the stage?

3. Landmarks that aid in locating the household: (Names of schools, Churches, businesses etc.)

4. How is the client called or referred to in the home area:

4a. How is the caregiver called or referred to in the home area: N/A

5. Does the patient have a treatment supporter? Yes No *(If yes, go to 5a, if no go to 6)*

5a. Treatment supporter name (Last, First, Middle)

5b. Treatment supporter home address

5c. Treatment supporter postal address (P.O Box)

5d. Treatment supporter number (Cell) .

5e. Treatment supporter number (Landline) .

6. Upcountry name for the client? N/A

6a. Upcountry contact home address? N/A

6b. Upcountry contact postal address? N/A

6c. Upcountry contact phone number (Cell) . N/A

6d. Upcountry contact phone number (Landline) . N/A

7. During the last year from today, have you been hit, slapped, Kicked, or hurt by someone?

Yes No Refused to answer

7a. If yes Who?

Spouse Steady Partner

Casual Partner Sibling

Parent Other(specify)

8. During the last year from today, have you ever hit, slapped, Kicked, or hurt someone?

Yes No Refused to answer

8a. If yes Who?

Spouse Steady Partner

Casual Partner Sibling

Parent Other(specify)

9. In the past 12 months have you:

9a1. Smoked? Yes No Refused to answer

9a2. If yes, number of cigarretes per day

9b1. Chewed Miraa Yes No Refused to answer

9b2. If yes, number of times per month

9c1. Smoked marijuana ? Yes No Refused to answer

9c2. If yes, number of times per month

9d1. Used cocaine? Yes No Refused to answer

9d2. If yes, number of times per month

9d3. If yes, how? Sniff Inject

smoke Other (specify)

9e1. Used intravenous drugs? Yes No Refused to answer

9e2. If yes, number of times per month

9e3. If yes, do you share needles? yes No Refused to answer

10. Do you drink alcohol? Yes No

10a. If yes, number of drinks per week

10b. In the last month, how often did you get drunk?

Never Daily Weekly 1-3 times a month

10c. In the last month, have you experienced any of the following after drinking alcohol?

Gotten in to a fight No Yes Refused to answer

Had accident/ Injured No Yes Refused to answer

Been arrested No Yes Refused to answer

Been raped (sex was forced on you) No Yes Refused to answer

Sexually assaulted someone No Yes Refused to answer

11. Can you use a condom during sex after you have been drinking or taking drugs?

Yes No Do not drink or use drugs Refused to answer

12. What is your Current Marital Status (tick one)

Married (monogamous) Separated

Married (Polygamous) Widowed

Cohabiting (come we stay) Single

Divorced

13. What is your occupation?

Unemployed Housewife

Employed Casual labourer

Self-employed Student

14. Housing roof type?

Corrugated iron sheet Makuti

Tiles Asbestos

Concrete Tin

Grass Other (specify)

15. How many adults live in the home?

16. How many children live in the home?

17. How much do you or your spouse earn in one month? (Ksh)

- 0 - 2,000 Ksh 20,001 - 30,000 Ksh
 2,001 - 5000 Ksh 30,001 - 50,000 Ksh
 5,001 - 10,000 Ksh > 50,001 Ksh
 10,001 - 20,000 Ksh Refused to answer

18. What is your highest level of education?(Tick one)

- No education
 Lower primary education (< 5years education)
 Five to eight years of primary education
 Some secondary education
 Beyond secondary education

19. How long does it take for you to travel to the clinic from home one way (Hrs/ Minutes)

Hours	Minutes
<input type="text"/>	<input type="text"/>

20. Do you have piped water in your home? Yes No

21. Do you have electricity at home? Yes No

22. What is your main source of cooking at home?

- Electricity Paraffin Firewood Solar Energy Other (Specify)

23. Social worker, does the client require home assessment Yes No

23a. If YES, why?

- Physically or mentally disabled adult
 Adult dependent on care-giver
 Client requests a home visit
 Disclosure (patient would like help disclosing status to family members)
 Family testing (patient would like other family members to be tested)
 Other (specify)

For the data use only (tick after scanning the form)

Scanned Date ___/___/_____ Name of data person

HOPE CLINIC ADULT TRACING AND HOME CARE FOLLOW-UP FORM

HOPE ID Number

Site Code

Today's Date (DD.MM.YYYY)

Interviewer number

REASONS FOR FOLLOW-UP VISIT

1. What is the reason for today's follow-up visit, as specified by previous visit (Check all that apply)

- | | |
|--|---|
| <input type="checkbox"/> Clinic attendance and/or ART adherence monitoring | <input type="checkbox"/> PMTCT Client |
| <input type="checkbox"/> Follow-up on physical health, HBC or first aid | <input type="checkbox"/> Hospital Admission |
| <input type="checkbox"/> Monitoring consistency of caregiver | <input type="checkbox"/> Lost to follow up client |
| <input type="checkbox"/> Follow-up on psycho-social issue or referral made(specify referral) | |
| <input type="checkbox"/> Counseling <input type="checkbox"/> Social work <input type="checkbox"/> Clinic <input type="checkbox"/> Pharmacy | |

Homebased counseling for patient or caregiver/household members (*Specify all that apply*):

<input type="checkbox"/> Patient	<input type="checkbox"/> Counseling for the Caregiver/Household
<input type="checkbox"/> ART and Adherence	<input type="checkbox"/> Care and support for PLWHA
<input type="checkbox"/> Management of side effects	<input type="checkbox"/> General HIV: Prevention, Transmission and management
<input type="checkbox"/> Disclosure	<input type="checkbox"/> knowing HIV status: Testing and care options
<input type="checkbox"/> General counseling	<input type="checkbox"/> Management of side effects for client needs
<input type="checkbox"/> Hygiene	<input type="checkbox"/> ART and Adherence
<input type="checkbox"/> Client basic care	<input type="checkbox"/> Disclosure
<input type="checkbox"/> Nutrition	<input type="checkbox"/> General counseling
<input type="checkbox"/> HIV prevention	<input type="checkbox"/> Hygiene
<input type="checkbox"/> PMTCT and Family Planning	<input type="checkbox"/> Nutrition
<input type="checkbox"/> Other (specify) <input style="width: 100px;" type="text"/>	<input type="checkbox"/> Other (specify) <input style="width: 100px;" type="text"/>

1a. This visit is taking place at the client's

- Place of residence
- Place of work or school
- Central market, shop or other public meeting place
- Hospital due to client's admission

HOPE CLINIC ADULT TRACING AND HOME CARE FOLLOW-UP FORM

PHYSICAL HEALTH MONITORING

2. During this visit how would you describe the client's physical condition/health?

- Stable and self dependent
- Immobile
- Sought medical attention for health complaints from previous visit *(Specify outcome)*
 - Condition improving No change Condition worsening
- Did not seek medical attention for health complaints from previous visit *(Specify outcome)*
 - Condition improving No change Condition worsening
- Presenting new health complaints at this visit *(Specify in question 3)*

3. Current symptoms or complaints assessed during this visit:

<input type="checkbox"/> None	<input type="checkbox"/> Minor Health Complaints	<input type="checkbox"/> Severe Health Complaints
	<input type="checkbox"/> Low grade fever (below 38)	<input type="checkbox"/> Bedridden/immobile
	<input type="checkbox"/> Headaches	<input type="checkbox"/> Severe coughing (2 weeks or more) with difficulty breathing
	<input type="checkbox"/> Fatigue	<input type="checkbox"/> Severe Burning/tingling in extremities
	<input type="checkbox"/> Nausea and or occasional vomiting	<input type="checkbox"/> Poor feeding
	<input type="checkbox"/> Mild diarrhea (occasional and loose stool)	<input type="checkbox"/> Severe Diarrhea (frequent and watery)
	<input type="checkbox"/> Abdominal pain	<input type="checkbox"/> Severe vomiting
	<input type="checkbox"/> Cough	<input type="checkbox"/> Persistent or high grade fever (above 39)
	<input type="checkbox"/> Fat changes	<input type="checkbox"/> Jaundice
	<input type="checkbox"/> Burning tingling in extremities	<input type="checkbox"/> Sores or skin lesions
	<input type="checkbox"/> Skin rash	<input type="checkbox"/> Mental confusion/Dementia
	<input type="checkbox"/> Other (specify) <input style="width: 100px;" type="text"/>	<input type="checkbox"/> Other (specify) <input style="width: 100px;" type="text"/>

4. How many meals does the client eat per day?

- One
 Two
 Three
 More than three
 None

HOPE CLINIC ADULT TRACING AND HOME CARE FOLLOW-UP FORM

CAREGIVER/HOUSEHOLD MONITORING

5. Has the caregiver changed since the last visit? Yes No (If NO, go to Q6)

5a. If YES, specify reason

- | | |
|---|---|
| <input type="checkbox"/> No longer able or willing to care for the patient | <input type="checkbox"/> Patient's condition now requires the help of a caregiver |
| <input type="checkbox"/> Found more suitable long term caregiver | <input type="checkbox"/> Other (Specify) <input style="width: 150px;" type="text"/> |
| <input type="checkbox"/> Patient refused care from this person | |
| <input type="checkbox"/> Patient's condition no longer requires the need of a caregiver | |

5b. If YES, who is the new caregiver?

- | | |
|---|---|
| <input type="checkbox"/> Self (No caregiver needed) | <input type="checkbox"/> Neighbour |
| <input type="checkbox"/> Parent | <input type="checkbox"/> Friend |
| <input type="checkbox"/> Spouse | <input type="checkbox"/> Social worker/Institution |
| <input type="checkbox"/> Relative | <input type="checkbox"/> Other (Specify) <input style="width: 150px;" type="text"/> |

5c. If YES, is the new caregiver aware of the clients HIV status? Yes No

6. At this visit are there areas of counseling or education which are needed in the household? (Tick all that apply)

<input type="checkbox"/> Patient	<input type="checkbox"/> Counselling for the Caregiver/Household
<input type="checkbox"/> ART and Adherence	<input type="checkbox"/> Care and support for PLWHA
<input type="checkbox"/> Management of side effects	<input type="checkbox"/> General HIV: Prevention, Transmission and management
<input type="checkbox"/> Disclosure	<input type="checkbox"/> Knowing HIV status: Testing and care options
<input type="checkbox"/> General counselling	<input type="checkbox"/> Management of side effects for client needs
<input type="checkbox"/> Hygiene	<input type="checkbox"/> ART and Adherence
<input type="checkbox"/> Client basic care	<input type="checkbox"/> Disclosure
<input type="checkbox"/> Nutrition	<input type="checkbox"/> General counseling
<input type="checkbox"/> HIV prevention	<input type="checkbox"/> Hygiene
<input type="checkbox"/> PMTCT and Family Planning	<input type="checkbox"/> Nutrition
<input type="checkbox"/> Opportunistic Infections and STIs	<input type="checkbox"/> Opportunistic Infections and STIs
<input type="checkbox"/> Other (specify) <input style="width: 100px;" type="text"/>	<input type="checkbox"/> Other (specify) <input style="width: 100px;" type="text"/>

HOPE CLINIC ADULT TRACING AND HOME CARE FOLLOW-UP FORM

PSYCO-SOCIAL MONITORING

7. Was referral made at the previous visit? Yes No *(If NO go to Q8)*

7a. If YES, specify

- | | |
|--|---|
| <input type="checkbox"/> Domestic violence | <input type="checkbox"/> Food Insecurity |
| <input type="checkbox"/> Drug or Alcohol abuse | <input type="checkbox"/> Legal aid |
| <input type="checkbox"/> Sexual violence | <input type="checkbox"/> Other (Specify) <input style="width: 200px; height: 15px;" type="text"/> |

7b. If YES did the client seek referral services? Yes No

- | | |
|---|--|
| <input type="checkbox"/> Transportation costs | <input type="checkbox"/> Client went but was turned away |
| <input type="checkbox"/> Client refused to go | <input type="checkbox"/> Forgot or lost referral slip |
| <input type="checkbox"/> Other (specify) <input style="width: 250px; height: 15px;" type="text"/> | |

8. Have there been any noticeable changes from the last visit in the psycho-social issues that were identified?

- No noticeable change Improvement Issues are worsening Cannot assess

9. During this visit were there any concerns in the household regarding any of the following? (Tick all that apply)

Physical abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Sexual abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Emotional abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Alcohol abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Drug abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Potential to self-inflict harm	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Depression	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Stigma and isolation	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Food insecurity	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Neglect by caregiver	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
NO CONCERNS	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported

HOPE CLINIC ADULT TRACING AND HOME CARE FOLLOW-UP FORM

CLINIC ATTENDANCE AND ADHERENCE MONITORING

10. Is the client on ARVs? Yes No

11. Since the last visit has the client refilled their prescription? Yes No (If YES go to Q 12)

11a. If NO specify reason:

- Client was not due for a refill since the last home visit
- Unable to get to clinic due to transport costs
- Unable to get to the clinic due to illness or weakness
- Travelled

- Lacked someone to assist to clinic
- Forgot
- Refused to go
- Other (specify)

12. Since the last visit, has the client missed any doses?

- One Two Three More than three None

12a. If doses were missed specify reason:

- Refused to take medications
- Forgot
- Doses were administered by caregiver who did not give
- Ran out of medication and could not refill prescription
- Felt better and decided to stop
- Drug or alcohol use affecting adherence
- Stigma, disclosure or privacy issues

- Was stopped by physician
- Side effects
- Felt too ill or too weak to take
- Medications lost or stolen
- Sharing medications with others
- Other (specify)

13. Is the client/caregiver able to recall critical information related to thier ARV regime?

Drug names	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None
Doses	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None
Times taken	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None
Side effects	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None
Food related indications	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None

14. Client adherence since the last visit has

- No change Improved Worsened Refer to clinician for review

HOPE CLINIC ADULT TRACING AND HOME CARE FOLLOW-UP FORM

15. Since the last visit, has the client attended all their scheduled clinic visits? Yes No Not due for visit

15a. If NO, Specify

- | | |
|---|--|
| <input type="checkbox"/> Too ill to come
<input type="checkbox"/> Forgot
<input type="checkbox"/> Lost appointment card
<input type="checkbox"/> Stigma within the household related to disclosure or privacy
<input type="checkbox"/> Need assistance from care-giver and none was available
<input type="checkbox"/> Traveled
<input type="checkbox"/> Depression | <input type="checkbox"/> Alcohol or drug use affected ability to come
<input type="checkbox"/> Refused to come or continue with program
<input type="checkbox"/> Could not afford transport
<input type="checkbox"/> Was seen at another clinic
<input type="checkbox"/> Work
<input type="checkbox"/> Other (specify) <input style="width: 200px; height: 15px;" type="text"/> |
|---|--|

FOLLOW - UP PLAN

16. Is the client eligible for discharge from the tracer and home care services? Yes No

16a. If NO, Next visit scheduled for:

- One week
 Two weeks
 One month
 Two months

Specify date of next visit((DD/MM/YYYY) / /)

16b. What actions will be taken as follow-up to this visit?

- Follow-up counseling in the home for client or caregiver/household
- Follow-up on physical health
- Adherence monitoring
- Referral for household member for site based VCT services
- Referral for suspected TB of client or household member
- Follow-up and monitoring of social conditions and actions previously taken
- Alert authorities for further investigation of neglect, sexual or domestic violence
- PMTCT
- Organise Hospital ambulatory services for immediate and urgent medical needs
- HIV positive Family /Household member for HIV/AIDS management
- Home based VCT for household member
- Organization referral or accompaniment
- Accompany client to clinic
- Counselor assisted disclosure
- No Action

HOPE CLINIC
ADULT TRACING AND HOME CARE FOLLOW-UP FORM

17. During this visit, was an appointment booked for the client at the clinic? Yes No
- If YES, Specify date of next appointment((DD/MM/YYYY)) / /
- If YES, specify the appointment type:
- Doctor Social worker
- Counselor Nutritionist
- Pharmacy

Comments

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

HOPE CLINIC ADULT TRACING AND HOME CARE SCREENING FORM

HOPE ID Number

Page 1

Site Code

Today's Date (DD.MM.YYYY)

Interviewer number

1. The reason for this client to be monitored through the tracer and Home Care program is

- Adult on ARVs not adhering to medications or clinic appointments Lost to follow up client
- Adult dependent on care-giver
- Post-partum mother
- Staff referred for other reason: (Specify)

1a. This visit is taking place at the client's

- Place of residence Central market, shop or other public meeting place
- Place of work or school Other(specify)
- Hospital due to client's admission

PHYSICAL HEALTH ASSESSMENT

2. At this initial screening in what physical condition did you find the client?

- Stable , mobile and able to take care of self
- Weak but mobile and able to take care of self
- Immobile or needs assistance to walk or move, reliant on caregiver

2a. Current symptoms or complaints assessed during this visit:

<input type="checkbox"/> None	<input type="checkbox"/> Minor Health Complaints	<input type="checkbox"/> Severe Health Complaints
	<input type="checkbox"/> Low grade fever (below 38)	<input type="checkbox"/> Bedridden/immobile
	<input type="checkbox"/> Headaches	<input type="checkbox"/> Severe coughing (2 weeks or more) with difficulty breathing
	<input type="checkbox"/> Fatigue	<input type="checkbox"/> Severe Burning/tingling in extremities
	<input type="checkbox"/> Nausea and or occasional vomiting	<input type="checkbox"/> Poor feeding
	<input type="checkbox"/> Mild diarrhea (occasional and loose stool)	<input type="checkbox"/> Severe Diarrhea (frequent and watery)
	<input type="checkbox"/> Abdominal pain	<input type="checkbox"/> Severe vomiting
	<input type="checkbox"/> Cough	<input type="checkbox"/> Persistent or high grade fever (above 39)
	<input type="checkbox"/> Fat changes	<input type="checkbox"/> Jaundice
	<input type="checkbox"/> Burning tingling in extremities	<input type="checkbox"/> Sores or skin lesions
	<input type="checkbox"/> Skin rash	<input type="checkbox"/> Mental confusion/Dementia

Skip to Question 4 if the residence is an institution/organisation

HOUSEHOLD ASSESSMENT

3. Describe the type of residence this is: (Skip to Question 4 if the residence is an institution/organisation)

Private residential home

Institution/organization (specify name):

3a. Who is the head of the household

Patient (Self)

Patient's Father

Patient's mother

Both parents

Patient's sibling (Specify age) years

Neighbour

Patient's relative (Aunt, Uncle, grandparent, cousin)

Friend

Other (Specify)

3b. How many individuals living in the household?

3b1. Adults (Age 15 and over) :

3b2. Children (Age 14 and below) :

3c. Number of rooms in the house:

One

Two

Three

More than three

3d. Does the house have electricity? Yes No

3e. What is the household's water source?

Piped to the house

Communal water tank within the vicinity

Water tank and piping to the house

Water from river, pool or open water source

Communal water tap within the vicinity

No water source in the vicinity

3f. What is the household's sanitation system?

Own flush toilet in the house

Shared pit latrine in vicinity of house

Private pit latrine

None

Shared toilet in vicinity of house

3g. What is the source of Energy for cooking?

Gas

Paraffin

Wood

Electricity

Charcoal

None

4. Does the patient have any special needs? Yes No Unknown

4a. If YES specify:

4a. If YES specify:

- Advanced illness
- Physically handicapped or disabled
- Mentally handicapped or disabled
- Emotionally/psychologically unstable or unwell
- Other(specify)

4b. If YES does the client currently have a caregiver? Yes No Unknown

4b1. If YES specify

- Spouse Friend
- Partner Neighbour
- Parent Social worker/local community health worker
- Relative Other(specify)

5. Is the client living in an environment where hygiene is neglected? Yes No

5a. If YES specify:

- Foul smell in the room where the client is staying Client has not had a bath for more than 2 days
- Client is sleeping in soiled beddings Other(Specify)

6. How many meals is the client eating per day?

- One Two Three More than three None

7. Does the client have access to food? Always Sometimes Never

NO CONCERNS

8. Are there there any concerns in the household regarding any of the following? (Tick all that apply)

Physical abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Sexual abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Emotional abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Alcohol abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Drug abuse	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Potential of self-inflict harm	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Depression	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Stigma and isolation	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Food insecurity	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
Neglect by caregiver	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported
NO CONCERNS	<input type="checkbox"/> Client Reported	<input type="checkbox"/> Staff Assessed	<input type="checkbox"/> Household Reported

COUNSELLING AND DISCLOSURE ASSESSMENT

9. Is the client aware of his or her status? Yes No Unknown

9a. If NO, is the caregiver willing to begin the disclosure process with the patient? Yes No Unknown

10. If the client has a caregiver, is the current caregiver aware of the patient's status?

Yes No Unknown There is no caregiver

10a. If NO, is the patient willing to begin the disclosure process with the caregiver?

Yes No Unknown

11. Are other household members aware of the client's status? All Some None

11a. If NO, is the patient/caregiver willing to begin the disclosure process with other household members?

Yes No

12. Is there a need for HIV counseling and testing of other household members? (Tick all that apply)

Spouse/Partner

Siblings

Mother

Other household members of unknown status in need of testing

Father

None

13. What areas of counselling and education are necessary for the Patient and household?

<input type="checkbox"/> Patient	<input type="checkbox"/> Counselling for the Caregiver/Household
<input type="checkbox"/> ART and Adherence	<input type="checkbox"/> Care and support for PLWHA
<input type="checkbox"/> Management of side effects	<input type="checkbox"/> General HIV: Prevention, Transmission and management
<input type="checkbox"/> Disclosure	<input type="checkbox"/> Knowing HIV status: Testing and care options
<input type="checkbox"/> General counselling	<input type="checkbox"/> Management of side effects for client needs
<input type="checkbox"/> Hygiene	<input type="checkbox"/> ART and Adherence
<input type="checkbox"/> Client basic care	<input type="checkbox"/> Disclosure
<input type="checkbox"/> Nutrition	<input type="checkbox"/> General counseling
<input type="checkbox"/> HIV prevention	<input type="checkbox"/> Hygiene
<input type="checkbox"/> PMTCT and family planning	<input type="checkbox"/> Nutrition
<input type="checkbox"/> Other (specify) <input type="text"/>	<input type="checkbox"/> Other (specify) <input type="text"/>

CLINIC ATTENDANCE AND ARV ADHERENCE ASSESSMENT

14. Is the client on ARVs? Yes No (If NO, skip to question 16)

14a. If yes, is the client able to tell you the following information?

Drug names	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None
Doses	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None
Times taken	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None
Side effects	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None
Food related indications	<input type="checkbox"/> All	<input type="checkbox"/> Some	<input type="checkbox"/> None

14b. During the last 7 days, how many doses did the client miss?

- One Two Three More than three None

14c. During the last 30 days, how many doses did the client miss?

- One Two Three More than three None

14c1. If the patient missed any doses, please specify reasons (Check all that apply)

<input type="checkbox"/> Refused to take medication	<input type="checkbox"/> Medications lost or stolen
<input type="checkbox"/> Forgot	<input type="checkbox"/> Sharing medications with others
<input type="checkbox"/> Doses are administered by caregiver who did not give	<input type="checkbox"/> Felt better and decided to stop
<input type="checkbox"/> Ran out of medication and could not refill prescription	<input type="checkbox"/> Stigma, disclosure or privacy issues
<input type="checkbox"/> Was stopped by physician	<input type="checkbox"/> Drug or alcohol use affecting adherence
<input type="checkbox"/> Side effects	<input type="checkbox"/> Other (specify)
<input type="checkbox"/> Felt too ill or too weak to take	<input type="text"/>

14d. When is the client due for a refill (check prescription)

- Date passed in a previous month
 Within the current month
 Next month
 Last month
 2 months or more
 No prescription available to confirm

15. Has the client/caregiver missed the client's last clinic visit? Yes No

15a. If YES, what reason did the client/caregiver fail to return to the clinic for your appointments?

- | | |
|---|---|
| <input type="checkbox"/> Too ill to come | <input type="checkbox"/> Alcohol or drug use affected ability to come |
| <input type="checkbox"/> Forgot | <input type="checkbox"/> Refused to come or continue with program |
| <input type="checkbox"/> Lost appointment card | <input type="checkbox"/> Was seen at another clinic |
| <input type="checkbox"/> Stigma within the household related to disclosure or privacy | <input type="checkbox"/> Depression |
| <input type="checkbox"/> Could not afford transport | <input type="checkbox"/> work |
| <input type="checkbox"/> Need assistance from care-giver and none was available | <input type="checkbox"/> Travel |
| <input type="checkbox"/> Other (specify) <input type="text"/> | |

FINAL ASSESSMENT AND FOLLOW-UP PLAN

16. Based on this initial assessment, what actions will be taken as follow-up to this visit? (check all that apply)

- | | |
|--|---|
| <input type="checkbox"/> Follow-up counseling for client or caregiver/household (refer to Question 13) | <input type="checkbox"/> Home based VCT for household member |
| <input type="checkbox"/> Referral for household member for site based VCT services | <input type="checkbox"/> Counselor assisted disclosure |
| <input type="checkbox"/> HIV+ family/household member for HIV/AIDS management | <input type="checkbox"/> Organization referral or accompaniment |
| <input type="checkbox"/> Referral for suspected TB of client/household member | <input type="checkbox"/> Accompany client to clinic |
| <input type="checkbox"/> Follow-up on physical health | <input type="checkbox"/> Adherence monitoring |
| <input type="checkbox"/> PMTCT | <input type="checkbox"/> No Action |
| <input type="checkbox"/> Organize hospital ambulatory services for immediate and urgent medical needs | |
| <input type="checkbox"/> Alert authorities for further investigation of neglect, sexual or domestic violence | |
| <input type="checkbox"/> Follow-up and monitoring of social conditions and actions previously taken | |

17. Next home visit scheduled for:

- One week
 Two weeks
 One month
 Two months

Specify date of next home visit((DD/MM/YYYY)) / /

18. During this visit, was an appointment booked for the client at the clinic? Yes No

If YES, Specify date of next appointment((DD/MM/YYYY)) / /

If YES, specify the appointment type:

- Doctor
 Counselor
 Pharmacy
 Social worker
 Nutritionist

Comments

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

For the data use only (tick after scanning the form)

Scanned Date ___ / ___ / _____ Name of data person

Modified March 2008

**Hope
Clinic**

Version

References

1. Ahdieh-Grant L, Li R, Levine AM, et al. Highly active antiretroviral therapy and cervical squamous intraepithelial lesions in human immunodeficiency virus-positive women. *J Natl Cancer Inst* 2004;96:1070-6.
2. Mandelblatt JS, Kanetsky P, Eggert L, Gold K. Is HIV infection a cofactor for cervical squamous cell neoplasia? *Cancer Epidemiol Biomarkers Prev* 1999;8:97-106.
3. Parkin D, Ferlay J, Hamdi-Cherif M, et al. Cervix Cancer. In: *Cancer in Africa: Epidemiology and Prevention*. Lyon: IARC Press; 2003:268-76.
4. Duerr A, Paramsothy P, Jamieson DJ, et al. Effect of HIV infection on atypical squamous cells of undetermined significance. *Clin Infect Dis* 2006;42:855-61.
5. Rowhani-Rahbar A, Hawes SE, Sow PS, et al. The impact of HIV status and type on the clearance of human papillomavirus infection among Senegalese women. *J Infect Dis* 2007;196:887-94.
6. Gichangi PB, Bwayo J, Estambale B, et al. Impact of HIV infection on invasive cervical cancer in Kenyan women. *AIDS* 2003;17:1963-8.
7. Franceschi S, Jaffe H. Cervical cancer screening of women living with HIV infection: a must in the era of antiretroviral therapy. *Clin Infect Dis* 2007;45:510-3.
8. Parham GP, Sahasrabuddhe VV, Mwanahamuntu MH, et al. Prevalence and predictors of squamous intraepithelial lesions of the cervix in HIV-infected women in Lusaka, Zambia. *Gynecol Oncol* 2006;103:1017-22.
9. Visual inspection with acetic acid for cervical-cancer screening: test qualities in a primary-care setting. University of Zimbabwe/JHPIEGO Cervical Cancer Project. *Lancet* 1999;353:869-73.
10. Doh AS, Nkele NN, Achu P, Essimbi F, Essame O, Nkegoum B. Visual inspection with acetic acid and cytology as screening methods for cervical lesions in Cameroon. *Int J Gynaecol Obstet* 2005;89:167-73.
11. Gaffikin L, Blumenthal PD, Emerson M, Limpaphayom K. Safety, acceptability, and feasibility of a single-visit approach to cervical-cancer prevention in rural Thailand: a demonstration project. *Lancet* 2003;361:814-20.
12. Denny L, Kuhn L, De Souza M, Pollack AE, Dupree W, Wright TC, Jr. Screen-and-treat approaches for cervical cancer prevention in low-resource settings: a randomized controlled trial. *Jama* 2005;294:2173-81.
13. Spinillo A, Zara F, Gardella B, Preti E, Gaia G, Maserati R. Cervical intraepithelial neoplasia and cervicovaginal shedding of human immunodeficiency virus. *Obstet Gynecol* 2006;107:314-20.
14. Anderson JR, Paramsothy P, Heilig C, Jamieson DJ, Shah K, Duerr A. Accuracy of Papanicolaou test among HIV-infected women. *Clin Infect Dis* 2006;42:562-8.
15. Schuman P, Ohmit SE, Klein RS, et al. Longitudinal study of cervical squamous intraepithelial lesions in human immunodeficiency virus (HIV)-seropositive and at-risk HIV-seronegative women. *J Infect Dis* 2003;188:128-36.
16. Wright TC, Jr., Subbarao S, Ellerbrock TV, et al. Human immunodeficiency virus 1 expression in the female genital tract in association with cervical inflammation and ulceration. *Am J Obstet Gynecol* 2001;184:279-85.

17. Wawer M, Kigozie G, Serwadda D, et al. Trial of Male Circumcision in HIV+ Men, Rakai, Uganda: Effects in HIV+ Men and in Women Partners. In: 15th Conference on Retroviruses and Opportunistic Infections; 2008; Boston; 2008.
18. Wright TC, Jr., Massad LS, Dunton CJ, Spitzer M, Wilkinson EJ, Solomon D. 2006 consensus guidelines for the management of women with cervical intraepithelial neoplasia or adenocarcinoma in situ. *Am J Obstet Gynecol* 2007;197:340-5.
19. National Cervical Cancer Prevention Program: Action Plan 2005-2009. In: Health DoR, ed.: Government of Kenya; 2005.
20. Pfaendler KS, Mwanahamuntu MH, Sahasrabuddhe VV, Mudenda V, Stringer JS, Parham GP. Management of cryotherapy-ineligible women in a "screen-and-treat" cervical cancer prevention program targeting HIV-infected women in Zambia: lessons from the field. *Gynecol Oncol* 2008;110:402-7.
21. Nene BM, Hiremath PS, Kane S, Fayette JM, Shastri SS, Sankaranarayanan R. Effectiveness, safety, and acceptability of cryotherapy by midwives for cervical intraepithelial neoplasia in Maharashtra, India. *Int J Gynaecol Obstet* 2008;103:232-6.
22. Luciani S, Gonzales M, Munoz S, Jeronimo J, Robles S. Effectiveness of cryotherapy treatment for cervical intraepithelial neoplasia. *Int J Gynaecol Obstet* 2008;101:172-7.
23. Mitchell MF, Tortolero-Luna G, Cook E, Whittaker L, Rhodes-Morris H, Silva E. A randomized clinical trial of cryotherapy, laser vaporization, and loop electrosurgical excision for treatment of squamous intraepithelial lesions of the cervix. *Obstet Gynecol* 1998;92:737-44.
24. Sadler L, Saftlas A, Wang W, Exeter M, Whittaker J, McCowan L. Treatment for cervical intraepithelial neoplasia and risk of preterm delivery. *JAMA* 2004;291:2100-6.
25. Acharya G, Kjeldberg I, Hansen SM, Sorheim N, Jacobsen BK, Maltau JM. Pregnancy outcome after loop electrosurgical excision procedure for the management of cervical intraepithelial neoplasia. *Arch Gynecol Obstet* 2005;272:109-12.
26. Chirenje ZM, Rusakaniko S, Akino V, Munjoma M, Mlingo M. Effect of HIV Disease in Treatment Outcome of Cervical Squamous Intraepithelial Lesions Among Zimbabwean Women. *J Low Genit Tract Dis* 2003;7:16-21.
27. Martin-Hirsch PL, Paraskevaidis E, Kitchener H. Surgery for cervical intraepithelial neoplasia. *Cochrane Database Syst Rev* 2000:CD001318.
28. Gichangi P, De Vuyst H, Estambale B, Rogo K, Bwayo J, Temmerman M. HIV and cervical cancer in Kenya. *Int J Gynaecol Obstet* 2002;76:55-63.
29. Parkin DM, Sitas F, Chirenje M, Stein L, Abratt R, Wabinga H. Part I: Cancer in Indigenous Africans--burden, distribution, and trends. *Lancet Oncol* 2008;9:683-92.
30. Gichangi P, Estambale B, Bwayo J, et al. Knowledge and practice about cervical cancer and Pap smear testing among patients at Kenyatta National Hospital, Nairobi, Kenya. *Int J Gynecol Cancer* 2003;13:827-33.
31. Yamada R, Sasagawa T, Kirumbi LW, et al. Human papillomavirus infection and cervical abnormalities in Nairobi, Kenya, an area with a high prevalence of human immunodeficiency virus infection. *J Med Virol* 2008;80:847-55.
32. Kamali A, Wagner HU, Nakiyingi J, Sibiiti I, Kengeya-Kayondo JF, Mulder DW. Verbal autopsy as a tool for diagnosing HIV-related adult deaths in rural Uganda. *Int J Epidemiol* 1996;25:679-84.
33. Wang CC, Reilly M, Kreiss JK. Risk of HIV infection in oral contraceptive pill users: a meta-analysis. *J Acquir Immune Defic Syndr* 1999;21:51-8.

34. Mostad SB, Overbaugh J, DeVange DM, et al. Hormonal contraception, vitamin A deficiency, and other risk factors for shedding of HIV-1 infected cells from the cervix and vagina. *Lancet* 1997;350:922-7.
35. Wang CC, McClelland RS, Overbaugh J, et al. The effect of hormonal contraception on genital tract shedding of HIV-1. *AIDS* 2004;18:205-9.
36. Morrison CS, Richardson BA, Mmiro F, et al. Hormonal contraception and the risk of HIV acquisition. *AIDS* 2007;21:85-95.
37. Del Mistro A, Bertorelle R, Franzetti M, et al. Antiretroviral therapy and the clinical evolution of human papillomavirus-associated genital lesions in HIV-positive women. *Clin Infect Dis* 2004;38:737-42.
38. Lillo FB, Ferrari D, Veglia F, et al. Human papillomavirus infection and associated cervical disease in human immunodeficiency virus-infected women: effect of highly active antiretroviral therapy. *J Infect Dis* 2001;184:547-51.
39. Graham SM, Holte SE, Peshu NM, et al. Initiation of antiretroviral therapy leads to a rapid decline in cervical and vaginal HIV-1 shedding. *AIDS* 2007;21:501-7.
40. Emery S, Bodrug S, Richardson BA, et al. Evaluation of performance of the Gen-Probe human immunodeficiency virus type 1 viral load assay using primary subtype A, C, and D isolates from Kenya. *J Clin Microbiol* 2000;38:2688-95.
41. Wang CC, McClelland RS, Reilly M, et al. The effect of treatment of vaginal infections on shedding of human immunodeficiency virus type 1. *J Infect Dis* 2001;183:1017-22.
42. Hawes SE, Critchlow CW, Faye Niang MA, et al. Increased risk of high-grade cervical squamous intraepithelial lesions and invasive cervical cancer among African women with human immunodeficiency virus type 1 and 2 infections. *J Infect Dis* 2003;188:555-63.
43. Mbizvo EM, Msuya SE, Stray-Pedersen B, Chirenje MZ, Hussain A. Cervical dyskaryosis among women with and without HIV: prevalence and risk factors. *Int J STD AIDS* 2005;16:789-93.
44. McKenzie K, Rogers R, Pamnani R, et al. Free cervical cancer screening among HIV-positive women receiving antiretroviral treatment in Kenya: acceptance and findings. In: 5th IAS Conference on HIV Pathogenesis, Treatment and Prevention; 2009 July 19-22; Cape Tno; 2009.