

Counselling and entry into HIV care

Submission date 01/06/2010	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 13/07/2010	Overall study status Completed	☐ Protocol
Last Edited 17/06/2015	Condition category Infections and Infestations	☐ Statistical analysis plan
		☒ Results
		☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Fred Wabwire-Mangeni

Contact details

Makerere University
School of Public Health
PO BOX 7072
Kampala
Uganda
256

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fwabwire@musph.ac.ug

Additional identifiers

Protocol serial number

MUSPH 06012009

Study information

Scientific Title

Assessing the effect of extended counselling on uptake and adherence to pre- antiretroviral care: A longitudinal, randomised controlled trial

Acronym

pre-ARV

Study objectives

Extended counselling is associated with increased uptake and adherence to pre-antiretroviral therapy (ART)

Ethics approval required

Old ethics approval format

Ethics approval(s)

1. Makerere University School of Public Health Higher Degrees Research and Ethics Committee, 06/01/2009
2. Uganda National Council for Science and Technology

Primary study design

Interventional

Study design

Longitudinal double-blind multicentre randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Pre-ARV care adherence

Interventions

Newly screened HIV positive persons will be followed and monitored for uptake and adherence to the pre-ART care for a period of 5 months depending on the frequency of enrolment of eligible subjects. The HIV positive persons from three centres will be classified into two arms.

1. Control arm: HIV positive persons will receive the current format and text of counselling
2. Intervention arm: HIV positive persons will receive a specially packaged continuum of extended counselling

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Proportion of HIV positive persons with extended counselling who adhere to pre-ART care

Key secondary outcome(s)

Proportion of HIV positive persons not lost to follow up during subsequent HIV care

Completion date

02/06/2010

Eligibility

Key inclusion criteria

Adult patients (>18 years) who test HIV positive at the ART centres at the start of the study (November 2009)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 Years

Sex

All

Key exclusion criteria

1. Patients undergoing Prevention of Mother-To-Child Transmission (PMTCT) therapy
2. Patients who are not of sound mental status
3. Those who are expected to leave the district during the period of follow-up

Date of first enrolment

02/12/2009

Date of final enrolment

02/06/2010

Locations**Countries of recruitment**

Uganda

Study participating centre

Makerere University

Kampala

Uganda

256

Sponsor information**Organisation**

Swedish International Development Authority - Department for Research Cooperation (SIDA /SAREC) (Sweden)

ROR

<https://ror.org/01fn7me06>

Funder(s)

Funder type

Government

Funder Name

Swedish International Development Authority - Department for Research Cooperation (SIDA /SAREC) (Sweden)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/05/2010		Yes	No
Results article	results	27/07/2011		Yes	No
Results article	results	01/10/2015		Yes	No