Urban community randomised trial for prevention of sexually transmitted diseases

Recruitment status	Prospectively registered	
No longer recruiting	Protocol	
Overall study status	Statistical analysis plan	
Completed	[X] Results	
Condition category	Individual participant data	
	No longer recruiting Overall study status Completed	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number 059131

Study information

Scientific Title

An urban community randomised trial of a multicomponent intervention for the prevention of sexually transmitted disease in female sex workers and the general population

Acronym

PREVEN

Study objectives

We aim to:

- 1. Determine the impact of a hybrid intervention on:
- 1.1. Prevalence of gonococcal, chlamydial, and trichomonal infection and syphilis seroreactivity in high-risk Female Sex Workers (FSW), bridging groups of clients of FSW, and the lower-risk young adult general population
- 1.2. The antimicrobial susceptibility and virulence-associated phenotypes of N. gonorrhoeae 2. Develop and employ mathematical models of gonorrhoea, chlamydial infection, syphilis and trichomoniasis to explain the observed impact of the interventions, and facilitate transfer of the intervention to other locations

Ethics approval required

Old ethics approval format

Ethics approval(s)

- 1. Universidad Peruana Cayetano Heredia, 07/12/1999, ref: #99084
- 2. University of Washington, 25/07/2000, ref: #99-2288
- 3. Naval Medical Research Center Detachment, 19/09/2002, ref: DOD#31556

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Sexually transmitted disease prevention

Interventions

Using data from a baseline survey in 24 Peruvian cities with populations more than 50,000, 20 cities were selected for the trial and placed into matched pairs based on region (coastal, jungle or Andean), size and STD prevalence. The pairs were then randomised into control or intervention groups, with the latter receiving a two-year hybrid intervention of:

- 1. Strengthened syndromic management through a network of physicians and pharmacists
- 2. STD screening and treatment of FSW
- 3. Social marketing of condoms

At the end of the two-year intervention, FSW and the young adult general population in all 20 cities will be surveyed again to evaluate its impact.

The anticipated end date of this trial has been extended to the 31st December 2006. The previous anticipated end date was 31/12/2005.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

- 1. Gonorrhoea, chlamydia, trichomonas, bacterial vaginosis, syphilis and human immunodeficiency virus (HIV) infection
- 2. Demographic and behavioural information regarding sexual history and risk behaviours

Key secondary outcome(s))

No secondary outcome measures

Completion date

31/12/2006

Eligibility

Key inclusion criteria

- 1. General population survey: one adult per household ages 18 to 29 years, either sex
- 2. FSW: willing to participate
- 3. Clients of FSW: sex with a commercial sex worker

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Key exclusion criteria

- 1. General population survey: over 29 years old
- 2. All surveys: under 18 years old

Date of first enrolment

24/08/2002

Date of final enrolment

31/12/2006

Locations

Countries of recruitment

Peru

United States of America

Study participating centre University of Washington Seattle, WA United States of America 98104

Sponsor information

Organisation

University of Washington (USA)

ROR

https://ror.org/00cvxb145

Funder(s)

Funder type

Charity

Funder Name

Wellcome Trust

Alternative Name(s)

Funding Body Type

Private sector organisation

Funding Body Subtype

International organizations

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration

Study outputs

Output type	Details	Date created Date add	ded Peer reviewed?	Patient-facing?
Results article	results	24/03/2012	Yes	No
Results article	results	01/10/2012	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/20	025 No	Yes
Study website	Study website	11/11/2025 11/11/20	025 No	Yes