# A feasibility study of an internet-based sexually transmitted infection (STI)/human immunodeficiency virus (HIV) behaviour change intervention

<b>Submission date</b> 03/05/2005	Recruitment status No longer recruiting	[X] Prospectively registered
03/03/2003	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
21/06/2005	Completed	☐ Results
Last Edited	Condition category	Individual participant data
28/04/2016	Infections and Infestations	Record updated in last year

# Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

Dr Helen Ward

#### Contact details

Dept Infectious Disease Epidemiology Imperial College London Norfolk Place London United Kingdom W2 1PG

h.ward@imperial.ac.uk

# Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

# Secondary identifying numbers

G0300712

# Study information

## Scientific Title

A feasibility study of an internet-based sexually transmitted infection (STI)/human immunodeficiency virus (HIV) behaviour change intervention

## Study objectives

To develop and test the feasibility of a novel computer-aided behavioural intervention for STI /HIV prevention, targeting people at high risk, with a view to testing the efficacy of this intervention in a future randomised controlled trial

## Ethics approval required

Old ethics approval format

## Ethics approval(s)

Not provided at time of registration.

## Study design

Randomised controlled trial

# Primary study design

Interventional

## Secondary study design

Randomised controlled trial

## Study setting(s)

Not specified

## Study type(s)

Prevention

## Participant information sheet

## Health condition(s) or problem(s) studied

Sexual Health

#### **Interventions**

Intervention: introduction and access to an internet-based intervention to reduce sexual behaviour risk

Controls: treatment and advice as usual

## Intervention Type

Other

#### Phase

## **Not Specified**

## Primary outcome measure

- 1. Attrition rate after 6 months
- 2. Reported sexual risk behaviours
- 3. Reinfection with an STI by 6 month follow-up point

## Secondary outcome measures

Not provided at time of registration.

## Overall study start date

01/09/2005

## Completion date

31/08/2007

# **Eligibility**

## Key inclusion criteria

Patients with an incident sexually transmitted infection (gonorrhoea, chlamydia, infectious syphilis, first clinical episode genital warts, primary genital herpes).

## Participant type(s)

**Patient** 

## Age group

Adult

## Sex

Both

# Target number of participants

360

## Key exclusion criteria

- 1. Non-English speaker
- 2. Functional illiteracy for internet use

## Date of first enrolment

01/09/2005

## Date of final enrolment

31/08/2007

# Locations

## Countries of recruitment

England

**United Kingdom** 

# Study participating centre Imperial College London

Dept Infectious Disease Epidemiology London United Kingdom W2 1PG

# Sponsor information

# Organisation

Sponsor to be confirmed - Record provided by the Medical Research Council (UK)

## Sponsor details

20 Park Crescent London United Kingdom W1B 1AL

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h.ward@imperial.ac.uk

## Sponsor type

Research council

# Funder(s)

# Funder type

Research council

## **Funder Name**

Medical Research Council (MRC) (UK)

## Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

## **Funding Body Type**

Government organisation

## **Funding Body Subtype**

National government

#### Location

# **Results and Publications**

**Publication and dissemination plan**Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

**IPD sharing plan summary**Not provided at time of registration