Bridging from emergency contraceptive pills (ECPs) to regular contraception

Submission date	Recruitment status	Prospectively registered
19/03/2009	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/03/2009	Completed	Results
Last Edited	Condition category	Individual participant data
30/03/2009	Pregnancy and Childbirth	Record updated in last year

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Protection of Human Subject Committee, study #9978

Study information

Scientific Title

Building bridges from emergency contraceptive to regular contraceptive use in pharmacies: a multicentre participant-randomised unblinded trial

Study objectives

Can non-users of a regular contraceptive method (a regular contraceptive method for the purposes of this study is defined as either an oral contraceptive pill, an injectable contraceptive method or an intra-uterine devide [IUD]) in Jamaica who intended to purchase an emergency contraceptive product in a pharmacy be encouraged to move toward adopting regular contraception by providing a discount coupon for one cycle of a regular oral contraceptive pill product?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Family Health International Protection of Human Subjects Committee gave approval on the 28th November 2006 (ref: 9978)

Study design

Multicentre participant-randomised unblinded trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Prevention

Participant information sheet

Health condition(s) or problem(s) studied

Family planning

Interventions

The intervention is a discount coupon given to randomly designated clients who arrive at a pharmacy to purchase an emergency contraceptive product.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Adoption of a regular contraceptive method. Adoption is defined as:

- 1. Use of oral contraceptive pills for at least two months
- 2. Receiving one injection of an injectable contraceptive method and either having received a second injection or intending to receive a second injection
- 3. Use of an IUD for at least one month

Measured at three and six months after participant enrolment (i.e., the pharmacy intercept interview).

Secondary outcome measures

Longest length of continuous use by women who adopt a regular contraceptive method. Measured at three and six months after participant enrolment (i.e., the pharmacy intercept interview).

Overall study start date

01/12/2006

Completion date

01/02/2008

Eligibility

Key inclusion criteria

Any woman (aged 16 - 46 years) who arrives at study pharmacy intending to purchase a dedicated emergency contraceptive product.

Participant type(s)

Patient

Age group

Adult

Sex

Female

Target number of participants

800

Key exclusion criteria

- 1. Males
- 2. Women under the age of 16
- 3. Women currently using a regular contraceptive method (a regular contraceptive method for the purposes of this study is defined as either an oral contraceptive pill, an injectable contraceptive method or an IUD)

Date of first enrolment

01/12/2006

Date of final enrolment

01/02/2008

Locations

Countries of recruitment

Jamaica

Study participating centre 25 Burlington Ave

Kingston Jamaica Kingston 10

Sponsor information

Organisation

The William and Flora Hewlett Foundation (USA)

Sponsor details

2121 Sand Hill Road Menlo Park, CA United States of America 94025 info@hewlett.org

Sponsor type

Charity

Website

http://www.hewlett.org/Default.htm

ROR

https://ror.org/04hd1y677

Funder(s)

Funder type

Charity

Funder Name

The William and Flora Hewlett Foundation (USA)

Results and Publications

Publication and dissemination planNot provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration