

# Reproductive health program for young women

<b>Submission date</b> 19/10/2020	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 21/10/2020	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 17/11/2023	<b>Condition category</b> Pregnancy and Childbirth	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

### Background and study aims

Young adult women might face reproductive health issues without being aware of them and tend to lack preventive health behavior. Therefore, in order to promote the reproductive health of young adult women, it is important to identify the stage of their health behavior and to develop a program that can induce behavior change. The aim of this study is to develop a reproductive health promotion program based on the Precaution Adoption Process Model (PAPM) for young adult women and to evaluate its effects.

### Who can participate?

Healthy young adult women volunteers aged 18-25, women who had coitus in the last 3 months and had partners, and women in Stages 1, 2, and 3 of the PAPM for reproductive health via an online survey

### What does the study involve?

The reproductive health promotion program developed in this study is strategically applied to each stage of the PAPM (Stages 1, 2, and 3). For Stage 1 participants in the intervention group, the program consists of strategies such as lectures, providing periodic information, small group discussions, Q&A sessions, and practical training. For Stage 2 participants in the intervention group, the program includes the strategies of stage 1 and additionally consists of strategies such as case experience using videos and individual counselling using the 5 Rs. For Stage 3 participants in the intervention group, the program includes the strategies of stage 1 and additionally consists of strategies such as individual counselling using the 5 As and providing tailored information according to the individual's needs. The program is replaced with educational material for the control group. In both groups, the total duration of the intervention is 8 weeks and the outcomes are measured at three times: baseline, 1 week after the seminar and 1 week after completion of the interventions.

### What are the possible benefits and risks of participating?

As possible benefits of participating, participants will play an active role in their own health care. Also, by contributing to this research, they can help in many other ways. However, the participants should also consider the following risks: participating in the program may cause unpleasant feelings or the intervention may not be effective for participants.

Where is the study run from?  
Korea University (Korea, South)

When is the study starting and how long is it expected to run for?  
May 2017 to November 2018

Who is funding the study?  
The Health Fellowship Foundation (Korea, South)

Who is the main contact?  
Inhae Cho  
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## Contact information

**Type(s)**  
Public

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## Additional identifiers

**EudraCT/CTIS number**  
Nil known

**IRAS number**

**ClinicalTrials.gov number**  
Nil known

**Secondary identifying numbers**  
1

## Study information

**Scientific Title**

Development and evaluation of a reproductive health promotion program based on the Precaution Adoption Process Model for young adult women

**Study objectives**

1. There will be differences in variables of the primary outcome (reproductive health-promoting behavior, rate of contraception use) between the intervention group who were provided with the reproductive health promotion program and the control group.
2. There will be differences in variables of the secondary outcome (reproductive health knowledge, reproductive health belief, reproductive health motivation, reproductive health self-efficacy) between the intervention group who were provided with the reproductive health promotion program and the control group.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Approved 18/12/2017, Korea University Institutional Review Board (145 Anam-ro, Seongbuk-gu, Seoul, 02841, Korea; +82 (0)2 3290 1137; [kuirb@korea.ac.kr](mailto:kuirb@korea.ac.kr)), ref: 1040548-KU-IRB-17-169-A-3

**Study design**

Randomized controlled trial

**Primary study design**

Interventional

**Secondary study design**

Randomised controlled trial

**Study setting(s)**

Community

**Study type(s)**

Prevention

**Participant information sheet**

Not available in web format, please use the contact details to request a participant information sheet

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

Reproductive health

**Interventions**

After conducting a preliminary, experimental investigation procedure, the program for promoting reproductive health based on the PAPM was applied step by step to the intervention group, and the program was replaced with educational material for the control group. The random allocation sequence was conducted with a 1:1 ratio using a computer-generated program (<https://www.randomizer.org/>) after the pretest.

The program is a tailored program whereby each stage of the PAPM (Stages 1, 2, and 3) is outlined, using components of the Information-Motivation-Behavioral skills model. The program

comprises intervention strategies, such as lectures; periodic information and information tailored to individuals' needs; small group discussions; question and answer sessions (Q&A); individual counselling; case experience using videos; and practical training.

The total duration of the intervention is 8 weeks. Outcomes are measured at three timepoints: T0 indicates the baseline, T1 indicates 1 week after the seminar (2 weeks after baseline) and T2 indicates within 1 week after completion of the interventions (8 weeks after baseline).

### **Intervention Type**

Behavioural

### **Primary outcome measure**

1. Reproductive health-promoting behavior measured using self-reported questionnaire via online survey at T0 (baseline), T1 (2 weeks after baseline), and T3 (after completion of intervention/8 weeks after baseline)
2. Rate of contraception use measured using self-reported questionnaire via online survey at T0, T3

Outcomes are measured at three timepoints: T0 indicates the baseline, T1 indicates 1 week after the seminar (2 weeks after baseline) and T2 indicates within 1 week after completion of the interventions (8 weeks after baseline).

### **Secondary outcome measures**

1. Reproductive health knowledge measured using self-reported questionnaire via online survey at T0, T1, T3
2. Reproductive health belief measured using self-reported questionnaire via online survey at T0, T1, T3
3. Reproductive health motivation measured using self-reported questionnaire via online survey at T0, T1, T3
4. Reproductive health self-efficacy measured using self-reported questionnaire via online survey at T0, T1, T3

Outcomes are measured at three timepoints: T0 indicates the baseline, T1 indicates 1 week after the seminar (2 weeks after baseline) and T2 indicates within 1 week after completion of the interventions (8 weeks after baseline).

### **Overall study start date**

24/05/2017

### **Completion date**

05/11/2018

## **Eligibility**

### **Key inclusion criteria**

1. Young adult women aged between 18 and 25
2. Women who had coitus in the last 3 months and had partners

Phase 2 of the study:

3. Women in Stages 1, 2, and 3 of the PAPM for reproductive health via an online survey

**Participant type(s)**

Healthy volunteer

**Age group**

Adult

**Lower age limit**

18 Years

**Sex**

Female

**Target number of participants**

The desired sample size for repeated ANOVA was 26 per study group. Considering the expected dropout rate, 30 participants were needed per group.

**Total final enrolment**

60

**Key exclusion criteria**

Women in Stages 4, 5, and 6 of the PAPM for reproductive health

**Date of first enrolment**

26/12/2017

**Date of final enrolment**

13/04/2018

**Locations****Countries of recruitment**

Korea, South

**Study participating centre****Korea University**

145 Anam-ro, Seongbuk-gu

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**Sponsor information****Organisation**

Korea University

### Sponsor details

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### Sponsor type

University/education

## Funder(s)

### Funder type

Charity

### Funder Name

The Health Fellowship Foundation (Korea, South)

## Results and Publications

### Publication and dissemination plan

Planned publication in a high-impact peer-reviewed journal.

### Intention to publish date

31/12/2022

### Individual participant data (IPD) sharing plan

The data-sharing plans for the current study are unknown and will be made available at a later date.

### IPD sharing plan summary

Data sharing statement to be made available at a later date

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>		27/10/2022	17/11/2023	Yes	No