

A randomised, controlled clinical trial involving women who have aborted spontaneously; the health, social and operational costs, outcomes of conservative and routine management

Submission date 10/10/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 10/10/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 02/07/2009	Condition category Pregnancy and Childbirth	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr TKH Chung

Contact details

Department of Obstetrics & Gynaecology

Prince of Wales Hospital

Chinese University of Hong Kong

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Hong Kong

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Additional identifiers

Protocol serial number

511007

Study information

Scientific Title

Study objectives

Not provided at time of registration

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Spontaneous abortion

Interventions

Routine surgical evacuation or medical evacuation of the uterus using misoprostol

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Misoprostol

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/01/1998

Eligibility**Key inclusion criteria**

All women admitted to the gynaecological unit of the Prince of Wales Hospital from October 1995 to January 1998 with:

1. A clinical diagnosis of spontaneous abortion

2. A positive urinary pregnancy test, and
3. Transvaginal Sonography (TVS) evidence of retained Products Of Conception (POCs)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/10/1995

Date of final enrolment

01/01/1998

Locations

Countries of recruitment

Hong Kong

Study participating centre

Department of Obstetrics & Gynaecology

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Hong Kong

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Sponsor information

Organisation

Hong Kong Health Services Research Fund (Hong Kong)

ROR

<https://ror.org/03qh32912>

Funder(s)

Funder type

Government

Funder Name

Hong Kong Health Services Research Fund (Hong Kong)

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	01/06/1999		Yes	No