ISRCTN43067169 https://doi.org/10.1186/ISRCTN43067169

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

Registration date
10/10/2002

Overall study status
Completed

Last Edited
02/07/2009

Condition category
Pregnancy and Childbirth

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] 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
-
Hong Kong
-

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

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 measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/10/1995

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

Age group
Adult

Sex
Female

Target number of participants
Not provided at time of registration

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
-

Hong Kong
-

Sponsor information

Organisation
Hong Kong Health Services Research Fund (Hong Kong)

Sponsor details
Health Welfare and Food Bureau
Government Secretariat, HKSAR
20th floor Murray Building
Garden Road
-
Hong Kong
-
+852 (0)2973 8288
hsrf@hwfb.gov.hk

Sponsor type
Government

Website
http://www.fhb.gov.hk/grants/english/funds/funds_hhsrf/funds_hhsrf_abt/funds_hhsrf_abt.html

ROR
https://ror.org/03qh32912

Funder(s)

Funder type
Government

Funder Name
Hong Kong Health Services Research Fund (Hong Kong)

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/1999		Yes	No