

# Non-surgical management of delayed miscarriage: a randomised trial

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|--|---|--|
| <b>Submission date</b><br>30/09/2004   | <b>Recruitment status</b><br>No longer recruiting     | <input type="checkbox"/> Prospectively registered    |
|  |   | <input type="checkbox"/> Protocol                    |
| <b>Registration date</b><br>30/09/2004 | <b>Overall study status</b><br>Completed              | <input type="checkbox"/> Statistical analysis plan   |
|  |   | <input type="checkbox"/> Results                     |
| <b>Last Edited</b><br>31/03/2020       | <b>Condition category</b><br>Pregnancy and Childbirth | <input type="checkbox"/> Individual participant data |
|  |   | <input type="checkbox"/> Record updated in last year |

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Mr D Economides

### Contact details

University Department Of Obstetrics and Gynaecology  
The Royal Free & University Medical School  
Pond Street  
Hampstead  
London  
United Kingdom  
NW3 2QG  
+44 (0)20 7830 2561 ext 3864  
abc@email.com

## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0256126964

# Study information

## Scientific Title

Non-surgical management of delayed miscarriage: a randomised trial

## Study objectives

Is outpatient medical management of delayed miscarriage effective, safe and acceptable as compared to expectant management?

## 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

Pregnancy and Childbirth: Miscarriage

## Interventions

1. Vaginal and oral misoprostol
2. Women managed expectantly

## Intervention Type

Other

## Phase

Not Specified

## Primary outcome measure

Complete miscarriage rates and number requiring surgical evacuation

## Secondary outcome measures

Not provided at time of registration

**Overall study start date**

09/07/2003

**Completion date**

31/12/2003

## **Eligibility**

**Key inclusion criteria**

100 women in total

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Female

**Target number of participants**

100 women in total

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

09/07/2003

**Date of final enrolment**

31/12/2003

## **Locations**

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

The Royal Free & University Medical School

London

United Kingdom

NW3 2QG

# Sponsor information

## Organisation

Department of Health

## Sponsor details

Richmond House  
79 Whitehall  
London  
United Kingdom  
SW1A 2NL

## Sponsor type

Government

## Website

<http://www.dh.gov.uk/Home/fs/en>

# Funder(s)

## Funder type

Hospital/treatment centre

## Funder Name

The Royal Free Hampstead NHS Trust (UK)

# 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