# A Comparison of Information Letters Sent to Patients Prior to Outpatient Clinic Appointments

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
30/09/2004	Stopped	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2004	Stopped	Results
Last Edited	Condition category	Individual participant data
25/10/2011	Other	<ul><li>Record updated in last year</li></ul>

# Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

### Contact name

Mrs V Cook

### Contact details

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# Additional identifiers

Protocol serial number N0558125486

# Study information

Scientific Title

# **Study objectives**

Does patient information sent prior to an appointment make patients better informed or increase the DNA (did not arrive) rate?

# 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)

**Not Specified** 

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

Not Applicable: Service delivery

### **Interventions**

Patient information sent prior to an appointment vs patient information not sent prior to an appointment

Added August 2008: trial stopped.

### Intervention Type

Other

### Phase

**Not Specified** 

# Primary outcome(s)

- 1. Reduction in DNA rate
- 2. Better informed patients

# Key secondary outcome(s))

Not provided at time of registration

### Completion date

01/08/2003

# **Eligibility**

# Key inclusion criteria

Initial pilot of 50 patients to test methodology. Sample convenience of 200 consecutive new patients to clinic.

# Participant type(s)

**Patient** 

# Healthy volunteers allowed

No

# Age group

**Not Specified** 

# Sex

**Not Specified** 

# Key exclusion criteria

Not provided at time of registration

# Date of first enrolment

01/01/2003

# Date of final enrolment

01/08/2003

# Locations

# Countries of recruitment

**United Kingdom** 

England

# Study participating centre Milton Keynes General NHS Trust Milton Keynes United Kingdom MK6 5LD

# Sponsor information

# Organisation

Department of Health

# Funder(s)

# Funder type

Government

# Funder Name

Milton Keynes General NHS Trust (UK)

# **Results and Publications**

Individual participant data (IPD) sharing plan

# IPD sharing plan summary

Not provided at time of registration