

A randomised pilot study of symptom-driven versus routine follow-up in asymptomatic breast cancer patients receiving regular mammography

Submission date 19/08/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 19/08/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 21/11/2019	Condition category Cancer	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

Dr - -

Contact details

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Additional identifiers

Protocol serial number

HOTLINE

Study information

Scientific Title

A randomised pilot study of symptom-driven versus routine follow-up in asymptomatic breast cancer patients receiving regular mammography

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)

Not Specified

Health condition(s) or problem(s) studied

Breast cancer

Interventions

1. Group A: Conventional follow-up
2. Group B: Mammogram only ("Hotline")

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/12/1997

Eligibility**Key inclusion criteria**

1. Breast cancer stage I or II
2. No known metastases
3. No prior or concurrent treatment other than tamoxifen

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

01/01/1995

Date of final enrolment

31/12/1997

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

UKCCCR Register Co-ordinator

London

United Kingdom

NW1 2DA

Sponsor information**Organisation**

International Collaborative Cancer Group (ICGG) (UK)

Funder(s)**Funder type**

Research organisation

Funder Name

International Collaborative Cancer Group (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration