

# UKCCCR trial to study the effect on breast cancer mortality of annual mammographic screening starting at age 40

<b>Submission date</b> 06/04/2000	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 06/04/2000	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 19/08/2020	<b>Condition category</b> Cancer	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

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### Contact details

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## Additional identifiers

### Protocol serial number

G9000793

## Study information

**Scientific Title**

UKCCCR trial to study the effect on breast cancer mortality of annual mammographic screening starting at age 40

**Acronym**

The 'Age' trial

**Study objectives**

To determine the effectiveness of mammographic screening starting at age 40, compared with starting at age 50, in reducing mortality from breast cancer.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Ethics approval from Central London REC 98/2/40.

**Study design**

Randomised controlled trial

**Primary study design**

Interventional

**Study type(s)**

Screening

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

Breast cancer

**Interventions**

Women in the intervention group are offered annual screening by mammography until the year of their 48th birthday. Screening is by 2-view mammography at 1st screen and single view subsequently. All women in both intervention and control groups will be invited for screening in the NHSBSP between the age of 50-52.

**Intervention Type**

Other

**Phase**

Not Applicable

**Primary outcome(s)**

Deaths from breast cancer in women free of the disease at trial entry in the two groups. Information on prognostic factors of all breast cancers is also collected.

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

31/12/2010

# Eligibility

## Key inclusion criteria

Women aged 40-41 identified from Health Authorities registers

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Adult

## Sex

Female

## Key exclusion criteria

Women under care for breast cancer may be removed from the prior notification list by the GPs prior to randomisation

## Date of first enrolment

30/07/1990

## Date of final enrolment

31/12/2010

# Locations

## Countries of recruitment

United Kingdom

England

## Study participating centre

Queen Mary University of London

London

United Kingdom

EC1M 6BQ

# Sponsor information

## Organisation

Medical Research Council (MRC) (UK)

# Funder(s)

## Funder type

Research council

## Funder Name

Medical Research Council (MRC) (UK)

## Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

## Funding Body Type

Government organisation

## Funding Body Subtype

National government

## Location

United Kingdom

## Funder Name

Long-term follow up funded by National Institute for Health Research (NIHR) / Health Technology Assessment (HTA)

# 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?
<a href="#">Results article</a>	results	09/12/2006		Yes	No
<a href="#">Results article</a>	results	01/09/2015		Yes	No
<a href="#">Results article</a>	results	01/04/2016		Yes	No
<a href="#">Results article</a>	results	01/09/2020	19/08/2020	Yes	No
	protocol	01/07			

<a href="#">Protocol article</a>		/1999		Yes	No
<a href="#">Other publications</a>	Implications of pathologist concordance	01/03 /2002		Yes	No
<a href="#">Other publications</a>	assessment of contamination in the control group:	01/04 /2010		Yes	No
<a href="#">Participant information sheet</a>	Participant information sheet	11/11 /2025	11/11 /2025	No	Yes