

A trial to study the place of surgery in elderly women with breast cancer treated by tamoxifen

Submission date 01/07/2001	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 01/07/2001	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 13/06/2014	Condition category 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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Additional identifiers

Protocol serial number
CRCBCTG6

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

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Breast

Interventions

1. Group A: Optimal surgery plus tamoxifen, 20 mg twice daily
2. Group B: Tamoxifen, 20 mg twice daily

All patients receive tamoxifen until there is evidence of progressive disease.

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. Aged >70 years
2. Histological, cytological or unequivocal mammographic confirmation of carcinoma of the breast
3. Operable breast cancer
4. No metastases
5. No previous malignancy, except skin cancer or adequately treated in situ carcinoma of the cervix
6. Fit and willing to receive either treatment

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Senior

Sex

Female

Key exclusion criteria

1. Patients with bilateral breast cancer are excluded
2. Patients with Paget's disease without a palpable lump, in situ carcinoma or an impalpable lesion shown on mammograph alone are excluded

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**

UK Co-ordinating Committee for Cancer Research (UKCCCR)

ROR

<https://ror.org/054225q67>

Funder(s)

Funder type
Not defined

Funder Name
Not provided at time of registration

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?
Results article	results	01/08/1997		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes