

Randomised trial of primary systemic therapy versus conventional therapy in operable breast cancer

Submission date 19/08/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 19/08/2002	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 24/10/2019	Condition category Cancer	<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

UKCCCR Register Co-ordinator
MRC Clinical Trials Unit
222 Euston Road
London
United Kingdom
NW1 2DA

Additional identifiers

Protocol serial number

SCTO37

Study information

Scientific Title

Randomised trial of primary systemic therapy versus conventional therapy in operable breast cancer

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: Mastectomy, wide local excision and axillary clearance plus adjuvant tamoxifen or CMF chemotherapy (depending on menstrual and nodal status)
2. Group B: Primary systemic therapy either endocrine or CAP chemotherapy (depending on oestrogen receptor status) followed by mastectomy, wide local excision and axillary clearance, and adjuvant endocrine or chemotherapy depending on prior response of tumour

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

31/10/1995

Eligibility**Key inclusion criteria**

1. Operable carcinoma of the breast
2. Age <70 years
3. No medical contraindications to protocol treatments

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/1990

Date of final enrolment

31/10/1995

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

UKCCCR Register Co-ordinator

London

United Kingdom

NW1 2DA

Sponsor information**Organisation**

Scottish Cancer Therapy Network (UK)

Funder(s)**Funder type**

Research organisation

Funder Name

Scottish Therapy Network (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration