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

Submission date 19/08/2002	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 19/08/2002	**Overall study status** Completed	
Last Edited 24/10/2019	**Condition category** Cancer	

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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Not Specified

Participant information sheet

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 measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/1990

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

Age group
Adult

Sex
Female

Target number of participants
Not provided at time of registration

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
England

United Kingdom

Study participating centre

UKCCCR Register Co-ordinator
London
United Kingdom
NW1 2DA

Sponsor information

Organisation
Scottish Cancer Therapy Network (UK)

Sponsor details
Trinity Park House
South Trinity Road
Edinburgh
United Kingdom
EH5 3SQ

Sponsor type
Research organisation

Funder(s)

Funder type
Research organisation

Funder Name
Scottish Therapy Network (UK)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration