

A Randomised Trial of Adriamycin and Taxotere vs. Adriamycin and Cyclophosphamide in Breast Cancer

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
19/08/2002	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
19/08/2002	Completed	<input checked="" type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
30/05/2012	Cancer	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr --

Contact details

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

Additional identifiers

Protocol serial number

BR 9809

Study information

Scientific Title

Study objectives

Added 12/02/10:

To compare the clinical and pathologic response rates of doxorubicin and cyclophosphamide (AC) with doxorubicin and docetaxel (AD) as primary chemotherapy in women with primary or locally advanced breast cancer.

Please note that as of 12/02/10 this record has been updated. All changes can be found in the relevant field with the above update date.

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)

Treatment

Health condition(s) or problem(s) studied

Breast cancer

Interventions

1. Adriamycin 50 mg/m² intravenous (iv) 3 weekly + taxotere 75 mg/m² iv 3 weekly (NB Taxotere must be administered prior to Adriamycin)
2. Adriamycin (60 mg/m² iv) 3 weekly + cyclophosphamide (600 mg/m² iv) weekly. Maximum of 6 cycles of combination chemotherapy.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Doxorubicin (Adriamycin®), docetaxel (Taxotere®), cyclophosphamide

Primary outcome(s)

Added 12/02/10:

Overall (complete and partial) clinical response rates

Key secondary outcome(s)

Added 12/02/10:

Number of relapses

Completion date

01/07/2002

Eligibility

Key inclusion criteria

1. Histologically proven breast cancer (BrCa)
2. Potentially operable disease >3cm diameter, locally advanced disease (including T4 lesions) or inflammatory BrCa
3. Patients older than 18 years

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Female

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/07/1999

Date of final enrolment

01/07/2002

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

UKCCR 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 Cancer Therapy Network (UK)

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	01/05/2005		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes