ISRCTN44725655 https://doi.org/10.1186/ISRCTN44725655

Comparison of second-line hormonal agents medroxyprogesterone acetate and aminoglutethimide in advanced 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 27/01/2015	**Condition category** Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
- -

Contact details
UKCCCR Register Co-ordinator
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Additional identifiers

Protocol serial number
BR3004

Study information

Scientific Title
Comparison of second-line hormonal agents medroxyprogesterone acetate and aminoglutethimide in advanced 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)
Treatment

Health condition(s) or problem(s) studied
Breast cancer

Interventions
Patients are randomised to receive either:
1. Group A: Medroxyprogesterone acetate 250 mg four times daily until disease progression or failure of treatment.
2. Group B: Aminoglutethimide 250 mg twice daily plus hydrocortisone 20 mg twice daily until disease progression or failure of treatment. There is a crossover option on disease progression or failure of treatment.

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Medroxyprogesterone acetate, aminoglutethimide, hydrocortisone

Primary outcome(s)
Not provided at time of registration.

Key secondary outcome(s))
Not provided at time of registration.

Completion date
01/08/1996

Eligibility

Key inclusion criteria

1. Advanced breast cancer
2. Relapsed or failed following tamoxifen treatment
3. Postmenopausal

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/08/1991

Date of final enrolment
01/08/1996

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
UKCCCR Register Co-ordinator
London
United Kingdom
NW1 2DA

Sponsor information

Organisation
Ciba-Geigy Pharmaceuticals (Switzerland)

ROR
https://ror.org/02f9zrr09

Funder(s)

Funder type
Industry

Funder Name
Ciba-Geigy Pharmaceuticals, Farmitalia Carlo Erba

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes