

A Double Blind, Randomised, Placebo Controlled Study of Farlutal in the Treatment of Weight Loss in Patients Undergoing Radiotherapy for Head and Neck Cancer

Submission date 19/08/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 19/08/2002	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 03/03/2008	Condition category Cancer	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

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Additional identifiers

Protocol serial number

MPA8801

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

Head and Neck Cancer

Interventions

1. Group A: Oral medroxyprogesterone acetate 500 mg twice daily for a maximum of 12 weeks
2. Group B: Placebo tablets twice daily for a maximum of 12 weeks

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

19/08/2005

Eligibility**Key inclusion criteria**

1. Aged >18 years
2. Squamous cell carcinoma of the head and neck
3. Not known to have or suspected of having early breast cancer
4. No known hypersensitivity to medroxyprogesterone acetate
5. No medical contraindications to treatment

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Lower age limit

18 years

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

19/08/2000

Date of final enrolment

19/08/2005

Locations**Countries of recruitment**

United Kingdom

England

Italy

Study participating centre

UKCCCR Register Co-ordinator

London

United Kingdom

NW1 2DA

Sponsor information**Organisation**

Farmitalia Carlo Erba (Italy)

ROR

<https://ror.org/03htt2d69>

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