

A study of adjuvant chemotherapy for malignant glioma

Submission date 28/02/2001	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 28/02/2001	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 12/04/2012	Condition category Cancer	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

http://www.ctu.mrc.ac.uk/research_areas/study_details.aspx?s=101

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number

BR05

Study information

Scientific Title

Study objectives

To assess whether the addition of chemotherapy to a standard radiotherapy regime influences the survival of adult patients with malignant glioma (astrocytomas grades 3 & 4) and to assess the ability of in vitro chemosensitivity testing to identify patients with a better prognosis.

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

Cancer

Interventions

1. One group receives chemotherapy added to a standard radiotherapy regimen
2. The other group receives none

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Chemotherapy

Primary outcome(s)

Survival time

Key secondary outcome(s))

Not provided at time of registration

Completion date

30/06/1996

Eligibility**Key inclusion criteria**

1. Age 18-69
2. No previous treatment except surgery, corticosteroids, anticonvulsants or diuretics

3. Entry within 6 weeks of surgery
4. Other malignancies
5. Fitness for treatment

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

69 years

Sex

All

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/09/1988

Date of final enrolment

30/06/1996

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

MRC Clinical Trials Unit

London

United Kingdom

NW1 2DA

Sponsor information

Organisation

Medical Research Council (MRC) (UK)

Funder(s)**Funder type**

Research council

Funder Name

Medical Research Council (MRC) (UK)

Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

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	15/01/2001		Yes	No