

Medical Research Council Trial of Immediate versus Deferred Treatment for Intracerebral Brain Tumours Presenting With Epilepsy

Submission date 19/08/2002	Recruitment status Stopped	☐ Prospectively registered
		☐ Protocol
Registration date 19/08/2002	Overall study status Stopped	☐ Statistical analysis plan
		☐ Results
Last Edited 15/11/2019	Condition category Nervous System Diseases	☐ Individual participant data
		☐ 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

BR08

Study information

Scientific Title

Medical Research Council Trial of Immediate versus Deferred Treatment for Intracerebral Brain Tumours Presenting With Epilepsy

Study objectives

Not provided at time of registration

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Brain and Nervous System

Interventions

1. Group A: Immediate treatment with clinicians choice of one or more of surgery, radiotherapy or chemotherapy, plus best medical treatment
2. Group B: Deferred treatment, with best medical treatment only until intervention becomes clinically necessary

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

10/12/1993

Reason abandoned (if study stopped)

Participant recruitment issue

Eligibility**Key inclusion criteria**

1. Patients with a clinically definite history of epilepsy who have Computed Tomography (CT) and /or Magnetic Resonance (MR) imaging supporting a probable diagnosis of intracerebral tumour

2. Clinical picture, CT or MR imaging is unequivocal and allows exclusion of cerebral abscess or extra-axial tumour (meningioma)
3. No other medical condition likely to effect the feasibility of follow-up
4. No previous neurosurgery, radiotherapy or chemotherapy
5. No other previous or concurrent malignant disease, except non melanomatous carcinoma of the skin
6. Age >16

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

01/12/1988

Date of final enrolment

10/12/1993

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

UKCCCR Register Co-ordinator

London

United Kingdom

NW1 2DA

Sponsor information

Organisation

Medical Research Council (MRC) (UK)

Funder(s)

Funder type

Government

Funder Name

Medical Research Council (UK)

Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, Medical Research Committee and Advisory 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