

A randomised double-blind controlled trial of BCNU as second-line therapy in metastatic oesophago-gastric, colorectal and pancreatic cancer

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| Submission date 19/08/2002 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered |
| | | <input type="checkbox"/> Protocol |
| Registration date 19/08/2002 | Overall study status Completed | <input type="checkbox"/> Statistical analysis plan |
| | | <input type="checkbox"/> Results |
| Last Edited 15/12/2015 | Condition category Cancer | <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

RMH E/C 1105

Study information

Scientific Title

A randomised double-blind controlled trial of BCNU as second-line therapy in metastatic oesophago-gastric, colorectal and pancreatic 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

Colon, oesophagus, pancreas, rectum, stomach cancers

Interventions

1. Arm 1: BCNU 200 mg in 500 ml 5% dextrose
2. Arm 2: 5% dextrose alone

Intervention Type

Drug

Phase

Not Applicable

Drug/device/biological/vaccine name(s)

BCNU

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

31/03/2001

Eligibility

Key inclusion criteria

Any patient who has progressed on or has a short disease free interval with folinic acid modulated 5-Fluorouracil (5FU) protracted infusional 5FU regimens

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

01/01/1995

Date of final enrolment

31/03/2001

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

MRC Clinical Trials Unit

London

United Kingdom

NW1 2DA

Sponsor information**Organisation**

The Royal Marsden NHS Foundation Trust (UK)

ROR

<https://ror.org/0008wzh48>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Royal Marsden Hospital (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration