

UK Hepatic Artery Pump Trial Two: Continuous Hepatic-Artery Infusion for Palliation in Colorectal Liver Metastases

Submission date 19/08/2002	Recruitment status No longer recruiting	☐ Prospectively registered ☐ Protocol
Registration date 19/08/2002	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 29/10/2019	Condition category Cancer	☐ 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
UKCCCR Register Co-ordinator
MRC Clinical Trials Unit
222 Euston Road
London
United Kingdom
NW1 2DA

Additional identifiers

Protocol serial number
UKHAP2

Study information

Scientific Title
UK Hepatic Artery Pump Trial Two: Continuous Hepatic-Artery Infusion for Palliation in Colorectal Liver Metastases

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

Cancer of Colon, Liver, Rectum

Interventions

1. Regimen A: Intra-hepatic continuous infusion of floxuridine for 5 days followed by systemic 5-fluorouracil and folinic acid for 5 days delivered via an arterial port-a-cath.
2. Regimen B: Systemic 5-fluorouracil and folinic acid for 5 days delivered via an arterial port-a-cath.

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

5-fluorouracil and folinic acid

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/09/1996

Eligibility**Key inclusion criteria**

1. Aged <75 years
2. Colorectal liver metastases, less than 45% liver involvement as assessed by Computed

Tomography (CT) scan
3. Karnofsky score more than 80%
4. Normal Bilirubin level
5. No prior chemotherapy

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/1990

Date of final enrolment

01/09/1996

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

UKCCCR Register Co-ordinator

London

United Kingdom

NW1 2DA

Sponsor information

Organisation

Colon Cancer Concern (UK)

ROR

<https://ror.org/03ngjs524>

Funder(s)

Funder type

Charity

Funder Name

Colon Cancer Concern (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration