

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	<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 29/10/2019	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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

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 measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/1990

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

Age group

Adult

Sex

Not Specified

Target number of participants

Not provided at time of registration

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

England

United Kingdom

Study participating centre
UKCCCR Register Co-ordinator
London
United Kingdom
NW1 2DA

Sponsor information

Organisation

Colon Cancer Concern (UK)

Sponsor details

c/o Bowel Cancer UK
7 Rickett Street
London
United Kingdom
SW6 1RU
+44(0)020 7381 9711
admin@bowelcanceruk.org.uk

Sponsor type

Charity

Website

<http://www.bowelcanceruk.org.uk/home>

ROR

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

Funder(s)

Funder type

Charity

Funder Name

Colon Cancer Concern (UK)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration