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

	<ul><li>Prospectively registered</li></ul>
19/08/2002 No longer recruiting	<pre>Protocol</pre>
Overall study status	Statistical analysis plan
Completed	Results
Condition category	Individual participant data
Cancer	<ul><li>Record updated in last year</li></ul>
	Completed  Condition category

# 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-acath.

# 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** 

# 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