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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

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