

# The effect of rofecoxib on colorectal liver metastases

<b>Submission date</b> 01/07/2001	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 01/07/2001	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 04/01/2012	<b>Condition category</b> Cancer	<input type="checkbox"/> Individual participant data

**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**  
CRC COX2

## Study information

**Scientific Title**

**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**

Colorectal cancer, liver metastases

**Interventions**

1. Rofecoxib 25 mg/day
2. Placebo

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

Not provided at time of registration

**Key secondary outcome(s))**

Not provided at time of registration

**Completion date**

05/01/2002

## Eligibility

**Key inclusion criteria**

1. Age greater than or equal to 18 years
2. Either sex
3. Liver resection deemed clinically appropriate for management of metastatic Colorectal Cancer (CRC) liver disease
4. Duration between decision to perform liver resection and surgery greater than 2 weeks
5. Ability to give written informed consent
6. Telephone at home

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

18 years

**Sex**

All

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/01/1998

**Date of final enrolment**

05/01/2002

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre**

UKCCCR Register Co-ordinator

London

United Kingdom

NW1 2DA

**Sponsor information****Organisation**

Cancer Research UK (CRUK) (UK)

**ROR**

<https://ror.org/054225q67>

# Funder(s)

## Funder type

Industry

## Funder Name

Merck and Co Inc. (USA)

# Results and Publications

## Individual participant data (IPD) sharing plan

## IPD sharing plan summary

Not provided at time of registration

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/09/2003		Yes	No