

Please complete this form and photocopy. Send the original immediately to your trials unit:

MRC: RADICALS Data Manager, MRC Clinical Trials Unit, Institute of Clinical Trials & Methodology, 90 High Holborn 2nd Floor, London, WC1V 6LJ

CCTG: Clinical Trials Assistant, CCTG, Queen's University, 10 Stuart Street, Kingston, Ontario, K7L 3N6, Canada

Patient's initials **Date of birth** **MRC Patient ID No.**

[illegible]

Responsible investigator:

Institution:

Please provide information on co-enrolment.

Initial box

For interventional (treatment) cancer trials, please initial the box to confirm that the trial team is aware of the patient's participation in this study. (For further information on co-enrolment, see the RADICALS CRF Completion Guidelines.)

Please complete **either** section A **or** section B.

A

B

1. Trial registration source:
e.g. ISRCTN registry (isrctn.com) or ClinicalTrials.gov (clinicaltrials.gov)

2. Trial registration number:

Trial registration number (ISRCTN or NCT number) should be available on the trial protocol, website or online registers such as isrctn.com; clinicaltrials.gov; or WHO ICTRP (apps.who.int/trialsearch/)

i. Trial name/acronym:

ii. Sponsor:

iii. Number of randomisation arms in the study:

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Date of randomisation into other study:

The diagram shows a 16-bit register divided into four fields: **d** (2 bits), **m** (4 bits), **v** (4 bits), and **CoEnr1_d** (6 bits). The **CoEnr1_d** field is highlighted with a green box.

Please confirm, by initialling the boxes, that you have read and understood the statements below:

Initial box

- I confirm that the above patient is co-enrolled into another clinical trial whilst participating in RADICALS and that RADICALS follow-up will not be impeded as a result of this co-enrolment.
- I also confirm that the information provided above is correct to the best of my knowledge, and that I have taken a copy for the patients site file.

Signed by
(Only MRC authorised person or CCTG investigator)

Date

d	m	y
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