

CANcer Diagnosis Decision rules

Submission date 12/12/2012	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 12/03/2013	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 20/03/2025	Condition category Cancer	<input type="checkbox"/> Individual participant data <input checked="" type="checkbox"/> Record updated in last year

Plain English Summary

<http://www.cancerresearchuk.org/cancer-help/trials/a-study-help-gps-decide-sooner-who-needs-tested-lung-or-bowel-cancer-candid>

Study website

<https://www.southampton.ac.uk/candid/index.page>

Contact information

Type(s)

Scientific

Contact name

Ms Natalie Thompson

Contact details

PhD Student, Research Programme Manager - The STREAM Study, Archivist (PCPS) & Mental Health First Aider

School of Primary Care, Population Sciences and Medical Education

University of Southampton

Aldermoor Health Centre

Aldermoor Close

Southampton

United Kingdom

SO16 5ST

-

n.thompson@soton.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

13492

Study information

Scientific Title

CANcer Diagnosis Decision rules: an observational study

Acronym

CANDID

Study hypothesis

This study seeks to work out which of the symptoms and examination findings are the most effective in the early prediction of lung or colon cancer.

The aim is to recruit up to 20,000 patients who consult their GP half with lung symptoms and the other half with colorectal symptoms. Clinical information will be collected using standardised internet based forms.

The clinical prediction 'rules' or decision aids developed from these studies will then be tested, with 2000 patients for each condition, for validity.

Ethics approval required

Old ethics approval format

Ethics approval(s)

South Central Oxford A, 31/07/2012, ref: 12/SC/0328

Study design

Non-randomised observational cohort study

Primary study design

Observational

Secondary study design

Cohort study

Study setting(s)

Hospital

Study type(s)

Diagnostic

Participant information sheet

Not available in web format, please use the contact details to request a patient information sheet

Condition

Cancer / primary care study

Interventions

Willing patients will complete lifestyle questionnaires and provide blood or saliva samples (including for genetic analysis). A notes review will also be undertaken. The National Cancer and Mortality Registries will then be monitored to see which patients develop cancer, and statistical analysis will determine the most important clinical variables that predict cancer.

Follow Up Length: 60 month(s)

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Participant appearing in cancer or mortality registries; Timepoint(s): up to 5 years post recruitment

Secondary outcome measures

No secondary outcome measures

Overall study start date

31/01/2013

Overall study end date

31/12/2022

Eligibility

Participant inclusion criteria

Male and female over 35 with lung or colorectal symptoms

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

UK Sample Size: 20000

Participant exclusion criteria

1. Already has a diagnosis of cancer
2. Unable to give informed consent

Recruitment start date

31/01/2013

Recruitment end date

30/09/2017

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Aldermoor Health Centre

Southampton

United Kingdom

SO16 5ST

Sponsor information

Organisation

University of Southampton (UK)

Sponsor details

Research Governance Office

George Thomas Building 37

Room 4055

Highfield

Southampton

England

United Kingdom

SO17 1BJ

Sponsor type

University/education

Website

<http://www.southampton.ac.uk>

ROR

<https://ror.org/01ryk1543>

Funder(s)

Funder type

Government

Funder Name

National School for Primary Care Research (UK)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

30/06/2023

Individual participant data (IPD) sharing plan

The data sharing plans for the current study are unknown and will be made available at a later date.

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol file	version v4	03/07/2015	31/07/2020	No	No