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Evaluation of open access nurse practitioner based diagnostic service in patients with rectal bleeding

Submission date 23/01/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 23/01/2004	**Overall study status** Completed	
Last Edited 03/12/2019	**Condition category** Cancer	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
SPGS782/SW

Study information

Scientific Title
Evaluation of open access nurse practitioner based diagnostic service in patients with rectal bleeding

Study objectives
To determine if the outlook from colorectal cancer can be improved by the provision of a nurse-based flexible sigmoidoscopy service for all patients with rectal bleeding.

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

Health condition(s) or problem(s) studied
Cancer (neoplasms): Colon; Cancer (neoplasms): Rectum

Interventions
1. Provision of a nurse-based flexible sigmoidoscopy service for all patients with rectal bleeding
2. Standard care

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure

Three-year incidence of advanced (Dukes' stage CID) colorectal cancer in intervention and control groups; delay between first GP contact and pathological diagnosis in above groups; cost and logistics of providing service to intervention group (including mobile service to subgroup of practices).

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/12/1995

Completion date
30/09/1999

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

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
01/12/1995

Date of final enrolment
30/09/1999

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

University of Southampton
Southampton
United Kingdom
SO16 6YD

Sponsor information

Organisation
NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
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+44 (0)20 7307 2622
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Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Executive South West (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