

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

Submission date 23/01/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 23/01/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 03/12/2019	Condition category Cancer	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
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

Study type(s)

Not Specified

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(s)

Three-year incidence of advanced (Dukes' stage C/D) 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).

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/09/1999

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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**

United Kingdom

England

Study participating centre

University of Southampton

Southampton

United Kingdom

SO16 6YD

Sponsor information**Organisation**

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

Funder(s)

Funder type

Government

Funder Name

NHS Executive South West (UK)

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes