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Prospective randomised controlled trial to assess the effect of colonoscopy patient information leaflets on patient knowledge and anxiety

Submission date
12/09/2003

Recruitment status
No longer recruiting

Registration date
12/09/2003

Overall study status
Completed

Last Edited
07/06/2017

Condition category
Surgery

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[ ] Results
[ ] Individual participant data
[ ] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0263107686

Study information

Scientific Title
Prospective randomised controlled trial to assess the effect of colonoscopy patient information leaflets on patient knowledge and anxiety

Study objectives
Do patient information leaflets for colonoscopy improve patient knowledge and anxiety scores?

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)
Hospital

Study type(s)
Other

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
Surgery: Colonoscopy

Interventions
1. Standard pre-colonoscopy consultation
2. Standard consultation plus an information leaflet

Intervention Type
Procedure/Surgery

Phase
Not Specified

Primary outcome measure
1. Patient knowledge scores
2. Patient anxiety scores (STAI)
3. Patient satisfaction scores

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/06/2001

Completion date
01/06/2007

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
80 patients from General Surgery

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/06/2001

Date of final enrolment
01/06/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre

Department of Surgery
London
United Kingdom
W1P 7LD

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

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

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
University College London Hospitals NHS Trust (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