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Effect of Metamucil on Bowel Preparation for Colonoscopy

Submission date 09/12/2003	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [X] Results [_] Individual participant data
Registration date 10/12/2003	**Overall study status** Completed	
Last Edited 29/08/2007	**Condition category** Surgery	

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
N/A

Study information

Scientific Title

Study objectives
Not provided at time of registration

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)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Bowel Preparation for Colonoscopy

Interventions
Patients randomised to 4 days of Metamucil versus Placebo prior to consuming a standard lavage, followed by colonoscopy

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Metamucil

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/01/2002

Completion date
01/01/2003

Eligibility

Key inclusion criteria
Patients scheduled for colonoscopy using a standard lavage preparation

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
64

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/01/2002

Date of final enrolment
01/01/2003

Locations

Countries of recruitment
United States of America

Study participating centre
4646 John R
Detroit
United States of America
48201

Sponsor information

Organisation
Procter and Gamble Company (USA)

Sponsor details
P.O. Box 8006
Mason,Ohio
United States of America
45040

Sponsor type
Industry

ROR
https://ror.org/04dkns738

Funder(s)

Funder type
Industry

Funder Name
Procter and Gamble Company (USA)

Funder Name
Wayne State University (USA)

Alternative Name(s)
Wayne State, WSU

Funding Body Type
Government organisation

Funding Body Subtype
Universities (academic only)

Location
United States of America

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	02/02/2004		Yes	No