

Comparison of small bowel barium (EZ-Paque) and pre-thickened barium (EZ-Varibar) for videofluoroscopic assessment of dysphagia

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 18/10/2016	Condition category Signs and Symptoms	<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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0226118475

Study information

Scientific Title

Comparison of small bowel barium (EZ-Paque) and pre-thickened barium (EZ-Varibar) for videofluoroscopic assessment of dysphagia

Study objectives

To assess whether the use of pre-thickened barium preparations is superior to the current practice of diluting and thickening powered barium to meet the patients' requirements.

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)

Diagnostic

Participant information sheet

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

Health condition(s) or problem(s) studied

Dysphagia

Interventions

Randomised comparative trial:

1. Small bowel barium (EZ-Paque)
2. Pre-thickened barium (EZ-Varibar)

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Completion of videofluoroscopy assessment.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/12/2002

Completion date

01/06/2004

Eligibility

Key inclusion criteria

50 consecutive patients referred for videofluoroscopy

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

50

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/12/2002

Date of final enrolment

01/06/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Wythenshawe Hospital
Manchester
United Kingdom
M23 9LT

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

Industry

Funder Name

Cook UK Ltd

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