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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	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 12/09/2003	**Overall study status** Completed	
Last Edited 18/10/2016	**Condition category** Signs and Symptoms	

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 videofluroscopy 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 videofluroscopy

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