

# The use of methylene blue dye in the diagnosis of dysplasia in Barrett's Oesophagus Surveillance

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 12/02/2010	<b>Condition category</b> Digestive System	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

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### Contact details

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## Additional identifiers

### Protocol serial number

N0436121390

## Study information

### Scientific Title

**Study objectives**

The purpose of this study is to compare the use of MB directed biopsies (MBDB) to random biopsies (RB) in the diagnosis of dysplasia.

**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**

Digestive System: Dysplasia

**Interventions**

1. Methylene blue directed biopsies
2. Random biopsies

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

The frequency of dysplasia between methylene blue directed biopsies and random biopsies.

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

01/01/2006

**Eligibility****Key inclusion criteria**

Barrett's oesophagus patients with recent diagnosis of dysplasia

**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**

01/01/2003

**Date of final enrolment**

01/01/2006

## **Locations**

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

Room 190A, Gastroenterology Department

Leeds

United Kingdom

LS1 3EX

## **Sponsor information**

**Organisation**

Department of Health (UK)

## **Funder(s)**

**Funder type**

Government

**Funder Name**

Leeds Teaching Hospitals NHS Trust (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?
<a href="#">Results article</a>	results	01/08/2006		Yes	No