

# Nasogastric and intravenous erythromycin as prokinetic in intensive care patients: a randomized controlled study

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| <b>Submission date</b><br>12/09/2003   | <b>Recruitment status</b><br>No longer recruiting | <input type="checkbox"/> Prospectively registered<br><input type="checkbox"/> Protocol                       |
| <b>Registration date</b><br>12/09/2003 | <b>Overall study status</b><br>Completed          | <input type="checkbox"/> Statistical analysis plan<br><input type="checkbox"/> Results                       |
| <b>Last Edited</b><br>16/09/2014       | <b>Condition category</b><br>Signs and Symptoms   | <input type="checkbox"/> Individual participant data<br><input type="checkbox"/> Record updated in last year |

## 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0259108904

# Study information

## Scientific Title

### Study objectives

Erythromycin's effect on gastric emptying in intensive care.

### 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)

Treatment

## Participant information sheet

### Health condition(s) or problem(s) studied

Signs and Symptoms:

### Interventions

Nasogastric erythromycin compared to intravenous erythromycin

### Intervention Type

Other

### Phase

Not Applicable

### 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**

01/07/2004

## Eligibility

**Key inclusion criteria**

Patients who exhibit a failure to absorb nasogastric feed

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Not Specified

**Target number of participants**

20

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/01/2002

**Date of final enrolment**

01/07/2004

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

The Walton Centre for Neurology & Neurosurgery

Liverpool

United Kingdom

L9 7LJ

## 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**

Government

**Funder Name**

The Walton Centre for Neurology and Neurosurgery 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