

# The effect of intraoperative fluid status optimisation guided by oesophageal doppler monitoring on outcome of patients following urgent or emergency laparotomy

<b>Submission date</b> 18/05/2001	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 18/05/2001	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 16/11/2011	<b>Condition category</b> Signs and Symptoms	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<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

EudraCT/CTIS number

IRAS number

**ClinicalTrials.gov number**

**Secondary identifying numbers**

G0100043

## **Study information**

**Scientific Title**

**Acronym**

FLOES

**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)**

Hospital

**Study type(s)**

Not Specified

**Participant information sheet**

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

Fluid status in emergency laparotomy

**Interventions**

In the intervention group pulse pressure variation measurements were used to guide fluid boluses of 6% Hydroxyethylstarch 130/0.4

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome measure**

Serum urea, creatinine and cystatin C levels were measured prior to and at the end of surgery and postoperatively on day 1, day 3 and day 5.

**Secondary outcome measures**

Not provided at time of registration.

**Overall study start date**

01/05/1999

**Completion date**

01/05/2002

## Eligibility

**Key inclusion criteria**

Not provided at time of registration.

**Participant type(s)**

Patient

**Age group**

Adult

**Sex**

Not Specified

**Target number of participants**

30

**Key exclusion criteria**

Not provided at time of registration.

**Date of first enrolment**

01/05/1999

**Date of final enrolment**

01/05/2002

## Locations

**Countries of recruitment**

Scotland

United Kingdom

**Study participating centre**  
**Department of Anaesthesia**  
Glasgow  
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## Sponsor information

### Organisation

Medical Research Council (MRC) (UK)

### Sponsor details

20 Park Crescent  
London  
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clinical.trial@headoffice.mrc.ac.uk

### Sponsor type

Research council

### Website

<http://www.mrc.ac.uk>

## Funder(s)

### Funder type

Government

### Funder Name

Medical Research Council (MRC) (UK)

### Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, MRC

### Funding Body Type

Government organisation

### Funding Body Subtype

National government

### Location

United Kingdom

# 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?
<a href="#">Results article</a>	results	01/06/2008		Yes	No