

# 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
		<input type="checkbox"/> Protocol
<b>Registration date</b> 18/05/2001	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
<b>Last Edited</b> 16/11/2011	<b>Condition category</b> Signs and Symptoms	<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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## Additional identifiers

### Protocol serial number

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

## Study type(s)

Not Specified

## 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(s)

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.

## Key secondary outcome(s))

Not provided at time of registration.

## Completion date

01/05/2002

# Eligibility

## Key inclusion criteria

Not provided at time of registration.

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Adult

## Sex

Not Specified

## 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

United Kingdom

Scotland

## Study participating centre

Department of Anaesthesia

Glasgow

United Kingdom

G4 0SF

# Sponsor information

## Organisation

Medical Research Council (MRC) (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

## 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