

# Multi centre randomised prospective trial of duodenal stenting versus surgical gastrojejunostomy for palliative relief of gastric outlet obstruction secondary to inoperable peri-ampullary cancers

<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> 23/05/2012	<b>Condition category</b> Surgery	<input type="checkbox"/> Individual participant data

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

**Protocol serial number**  
N0256107988

# Study information

## Scientific Title

## Study objectives

Which method of gastric bypass offers the best palliation with lowest cost and least mortality and morbidity?

## 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)

Treatment

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

Surgery: Gastric bypass

## Interventions

Patients randomised to receive duodenal stenting or surgical bypass. Followed up until death with analysis of morbidity, mortality, survival, cost, readmission rates, palliation of symptoms and quality of life.

## Intervention Type

Procedure/Surgery

## Phase

Not Specified

## Primary outcome(s)

30 Day mortality, morbidity, cost, quality of life, survival after treatment, recurrent hospital admissions and reasons, successful placement of stents, return to nutrition and level of return to nutrition.

## Key secondary outcome(s)

Not provided at time of registration

## Completion date

03/01/2004

# Eligibility

## Key inclusion criteria

35 patients in each group.

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Not Specified

## Sex

Not Specified

## Key exclusion criteria

Does not match inclusion criteria

## Date of first enrolment

03/01/2002

## Date of final enrolment

03/01/2004

# Locations

## Countries of recruitment

United Kingdom

England

## Study participating centre

University Department of Surgery

London

United Kingdom

NW3 2QG

# Sponsor information

## Organisation

Department of Health (UK)

# Funder(s)

## Funder type

Hospital/treatment centre

## Funder Name

The Royal Free Hampstead 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	01/06/2007		Yes	No
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes