

# Prevention of radiographic contrast media induced renal injury by administration of intravenous N-acetylcysteine in vascular patients undergoing angiography/angioplasty

<b>Submission date</b> 12/09/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 12/09/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 03/08/2009	<b>Condition category</b> Urological and Genital Diseases	<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

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## Additional identifiers

**Protocol serial number**  
N0256108007

## Study information

## Scientific Title

### Study objectives

To assess if administration of N-acetylcysteine prevents or reduces the risk of acute renal damage following administration of radiographic contrast media in patients with vascular disease undergoing angiography/angioplasty.

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Not provided at time of registration

### Study design

Randomised double blind placebo controlled trial

### Primary study design

Interventional

### Study type(s)

Prevention

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

Urological and Genital Diseases: Renal injury

### Interventions

Intravenous N-acetylcysteine vs placebo

### Intervention Type

Drug

### Phase

Not Specified

### Drug/device/biological/vaccine name(s)

N-acetylcysteine

### Primary outcome(s)

Service outcomes development.

### Key secondary outcome(s)

Not provided at time of registration

### Completion date

01/12/2003

## Eligibility

### Key inclusion criteria

100 patients

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

All

**Key exclusion criteria**

Does not meet inclusion criteria.

**Date of first enrolment**

01/03/2002

**Date of final enrolment**

01/12/2003

**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	01/12/2004		Yes	No