

# Multi-centre randomised controlled trial of nurse practitioners and pre-registration house officers (PRHO) in pre-operative workup

<b>Submission date</b> 25/04/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 25/04/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 11/03/2009	<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

### Contact name

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### Contact details

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## Additional identifiers

### Protocol serial number

HTA 94/40/38

## Study information

## Scientific Title

### Study objectives

To determine whether pre-operative assessment carried out by an appropriately trained nurse (ATN) is equivalent in quality to that carried out by a pre-registration house officer (PRHO).

To assess whether pre-assessments carried out by ATNs and PRHOs are equivalent in terms of cost.

To determine whether assessments carried out by ATNs are acceptable to patients.

To investigate the quality of communication between senior medical staff and ATNs.

More details can be found at: <http://www.hta.ac.uk/993>

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Not provided at time of registration.

### Study design

Prospective randomised equivalence trial

### Primary study design

Interventional

### Study type(s)

Not Specified

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

Pre-operative assessment

### Interventions

The study design was principally a prospective randomised equivalence trial but was accompanied by additional qualitative assessment of patient and staff perceptions, and an economic evaluation.

The intervention consisted of a pre-operative assessment carried out by either an appropriately trained nurse (ATN) or a PRHO. Of the patients who completed the study with a full evaluation, 926 patients were randomised to the PRHO arm of the trial and 948 to the ATN arm. Three ATNs took part in the study, one from each centre, together with a total of 87 PRHOs.

### Intervention Type

Procedure/Surgery

### Phase

Not Specified

### Primary outcome(s)

Immediately following the initial assessment of a patient by a PRHO or an ATN, one of a number of clinical research fellows, all specialist registrars in anaesthetics, repeated the assessment and recorded it on a study form, together with a list of investigations required. The clinical research fellow then evaluated the competency of the initial assessor by comparing the quality of their

assessment with their own. Any deficiencies in ordering of investigations and referral to other specialities were met in order to maximise patient care. Three areas of ATN and PRHO performance were judged separately, history taking, examination and ordering of tests, and each was graded into one of four categories, the most important of which was under-assessment, which would possibly have affected peri-operative management. In the case of ordering of tests, it was possible to have both over- and under-assessed a patient on different tests.

**Key secondary outcome(s)**

Not provided at time of registration.

**Completion date**

31/10/1999

## Eligibility

**Key inclusion criteria**

All patients attending at one site for assessment prior to general anaesthetic for elective general, vascular, urological or breast surgery were potentially included in the study. Of 1,907 patients who were randomised, 1874 completed the study with a full evaluation.

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

All

**Key exclusion criteria**

Not provided at time of registration.

**Date of first enrolment**

01/10/1997

**Date of final enrolment**

31/10/1999

## Locations

**Countries of recruitment**

United Kingdom

England

**Study participating centre**  
**University Surgical Unit**  
Southampton  
United Kingdom  
SO16 6YD

## Sponsor information

**Organisation**  
Department of Health (UK)

**ROR**  
<https://ror.org/03sbpja79>

## Funder(s)

**Funder type**  
Government

**Funder Name**  
NIHR Health Technology Assessment Programme - HTA (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	07/12/2002		Yes	No