

# Co-managed care in a peripheral ophthalmic clinic

<b>Submission date</b> 23/01/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 23/01/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 17/09/2010	<b>Condition category</b> Eye Diseases	<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**  
PSI14-20

## Study information

**Scientific Title**

**Study objectives**

The aim of the study was to test the key hypothesis that: Co-management of care in a peripheral ophthalmic clinic will improve the efficiency and cost-effectiveness of, and patient satisfaction with, day-care management of cataract surgery. Three objectives were identified:

1. To assess the value of introducing a trained ophthalmic nurse into the care of patients with cataract attending the peripheral clinic
2. To ascertain the best site for pre-operative assessment (peripheral or main hospital)
3. To evaluate the post-operative management of patients including possible role changes in the multi-disciplinary team

**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**

Cataract

**Interventions**

Patients were randomly allocated to receive either the experimental (introduction of a trained ophthalmic nurse) or control (standard) model of care. N = 31 - intervention, N = 32 - control

**Intervention Type**

Other

**Phase**

Not Applicable

**Primary outcome(s)**

For both groups, visual outcomes (acuity and function), anxiety and depression, costs to patients and the NHS, and patient satisfaction were the outcome measures selected. Observational data relating to development of staff roles were also collected. All data were collected by an independent researcher.

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

10/07/1997

# Eligibility

## Key inclusion criteria

Patients with cataracts

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Adult

## Sex

All

## Key exclusion criteria

Does not match inclusion criteria

## Date of first enrolment

01/07/1995

## Date of final enrolment

10/07/1997

# Locations

## Countries of recruitment

United Kingdom

England

## Study participating centre

School of Nursing, Midwifery and Health Visiting

Manchester

United Kingdom

M13 9PL

# Sponsor information

## Organisation

Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

# Funder(s)

## Funder type

Government

## Funder Name

NHS Primary and Secondary Care Interface National Research and Development Programme (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	01/06/1999		Yes	No
<a href="#">Protocol article</a>	protocol	01/06/1999		Yes	No