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Co-managed care in a peripheral ophthalmic clinic

Submission date	Recruitment status	[_] Prospectively registered
23/01/2004	No longer recruiting	[X] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
23/01/2004	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
17/09/2010	Eye Diseases	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Heather Waterman

Contact details
School of Nursing, Midwifery and Health Visiting
University of Manchester
Oxford Road
Manchester
United Kingdom
M13 9PL
+44 (0)161 275 5334
Heather.waterman@man.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

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 measure
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.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/07/1995

Completion date
10/07/1997

Eligibility

Key inclusion criteria
Patients with cataracts

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
63

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
England

United Kingdom

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)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Primary and Secondary Care Interface National Research and Development Programme (UK)

Results and Publications

Publication and dissemination plan
Not provided at time of registration

Intention to publish date

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?
Protocol article	protocol	01/06/1999		Yes	No
Results article	results	01/06/1999		Yes	No