

A randomised controlled trial to assess outcomes associated with three modalities of telephone follow-up, by qualified nurses, of recently discharged day surgery patients.

Submission date 23/01/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 23/01/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 20/07/2011	Condition category Other	<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

N/A

Study information

Scientific Title

Study objectives

The aim of the study is to determine the effect on outcomes following day surgery of a range of levels of telephone interventions by qualified nurses for patient guidance support.

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)

Not Specified

Health condition(s) or problem(s) studied

Not applicable: service delivery

Interventions

Three different modes of telephone intervention follow-up

1. Once at day 1
2. Twice, at days 1 and 3
3. Once at 7 days after discharge

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

1. Suitability and timing of post-discharge telephone support, measured by a tool adapted from Orem's self-care deficit theory
2. Compliance with discharge instructions measured by a tool adapted for telephone interview from the Revised Ways of Coping Checklist (WCCL)
3. Frequency of use of Primary Health Care Services
4. GPs assessment of the appropriateness of referrals and contacts

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/04/1997

Eligibility

Key inclusion criteria

Patients aged 17 or above, attending the day surgery centre at Taunton & Somerset Hospital for therapeutic procedures not normally requiring GP, District Nurse or hospital out-patient follow-up. Other criteria include patient consent and patient access to a telephone.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/08/1995

Date of final enrolment

30/04/1997

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Taunton and Somerset NHS Trust

Taunton

United Kingdom

TA1 5DA

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

NHS Executive South West (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?
Results article	results	01/12/1998		Yes	No