

To evaluate the impact of a Carer Support Needs Assessment Tool (CSNAT) intervention in hospice home care

Submission date 22/06/2012	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 22/06/2012	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 06/09/2016	Condition category Other	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Most people prefer to be cared for and to die in the familiar surroundings of their own home. Being able to make this choice very much depends on the efforts of family carers. While taking on a caring role is normally done willingly, it can result in considerable burden and negative effects on carers health. Therefore it is important to ensure that carers are well supported to enable them to look after patients at home while also maintaining their own health and wellbeing. So far health professionals have had no clear guidelines to help them ensure that they assess and address carer support needs regularly and in detail. So we developed a simple assessment form, the Carer Support Needs Assessment Tool (CSNAT), based on what carers told us their main support needs were in end of life home care. The procedures for CSNAT assessment, prioritisation and follow-up together form the CSNAT intervention. The aim of the study is to find out whether the using the CSNAT intervention with carers during care giving makes a difference to carer outcomes in bereavement.

Who can participate?

All adult carers of patients looked after by the hospice home care (HHC) services can take part in the study.

What does the study involve?

We will ask six HHC services to use the CSNAT intervention with carers of patients under their care. We will then compare bereavement outcomes for carers who had usual care with those who had the intervention. The HHC services will start using the CSNAT at three month intervals so we can make a comparison between services who have started the intervention with those who have not, and also make a comparison before and after the start of the intervention within each service. We will measure outcomes using a postal survey of carers 4-5 months after bereavement. To reduce any potential risks, HHC will exclude carers that are believed to suffer particularly complicated grief in bereavement, and contact details to HHC bereavement support will be provided. The study will also ask the views of the HHC nurses and some of the carers who

had the intervention on what made the intervention successful or not so successful. Finally, we will look at HHC CSNAT records to see whether some factors, such as how often assessments were done, seem to make a difference.

What are the possible benefits and risks of participating?

Participants in the CSNAT intervention group may receive better support during care giving and experience better health, well being and lower levels of grief in bereavement.

Where is the study run from?

University of Manchester (UK).

When is the study starting and how long is it expected to run for?

The study started in May 2012 and will run for 2 years.

Who is funding the study?

National Institute for Health Research (NIHR) (UK).

Who is the main contact?

Gunn Grande

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Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

12422

Study information

Scientific Title

A randomised trial to evaluate the impact of a Carer Support Needs Assessment Tool (CSNAT) intervention in hospice home care

Acronym

CSNAT

Study objectives

To test whether a formalised, comprehensive procedure for carer support needs assessment, prioritisation and follow up (CSNAT intervention) improves quality of care and carer outcomes in end of life home care, compared to usual care.

Ethics approval required

Old ethics approval format

Ethics approval(s)

NRES Committee North West - Greater Manchester East, 23/03/2012, ref: 12/NW/0206

Study design

Stepped wedge cluster randomised trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Quality of life

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Topic: Primary Care Research Network for England, Generic Health Relevance and Cross Cutting Themes; Subtopic: Not Assigned, Generic Health Relevance (all Subtopics); Disease: All Diseases, Health Services Research

Interventions

CSNAT, Formalised, comprehensive procedure for carer support needs assessment, prioritisation and follow up using the Carer Support Needs Assessment Tool (CSNAT)

Control, informal carer assessment as usual

Follow Up Length: 5 month(s)

Intervention Type

Other

Phase

Phase II/III

Primary outcome measure

SF-12 4-5 months post bereavement

Secondary outcome measures

1. Quality of care; Timepoint(s): 4-5 months post bereavement
2. Patient's place of death; Timepoint(s): Date of death
3. Texas Revised Inventory of Grief; Timepoint(s): 4-5 months post bereavement

Overall study start date

21/09/2012

Completion date

31/12/2013

Eligibility**Key inclusion criteria**

1. The main bereaved carer of patients who have been supported by participating hospice home care services (HHCs)
2. Target Gender: Male & Female ; Lower Age Limit 18 years

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Planned Sample Size: 1003; UK Sample Size: 1003

Key exclusion criteria

1. Age <18 years
2. Identified by HHCs as experiencing particularly complicated grief in bereavement

Date of first enrolment

21/09/2012

Date of final enrolment

31/12/2013

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

School of Nursing, Midwifery and Social Work

Manchester

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Sponsor information

Organisation

University of Manchester (UK)

Sponsor details

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Sponsor type

University/education

ROR

<https://ror.org/027m9bs27>

Funder(s)

Funder type

Government

Funder Name

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