A controlled evaluation of the health benefits of a participative community music programme for older people (Silver Song Clubs)

Submission date	Recruitment status	[X] Prospectively registered	
25/08/2009	No longer recruiting	[X] Protocol	
Registration date	Overall study status	Statistical analysis plan	
28/09/2009 Last Edited	Completed Condition category	Results	
		[] Individual participant data	
04/07/2011	Other	Record updated in last year	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Stephen Clift

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

The health benefits of a participative community music programme for older people (Silver Song Clubs): a controlled evaluation single centre randomised unblinded trial

Study objectives

Primary hypothesis:

Singing groups for older people improve both physical and mental aspects of quality of life when compared with usual activities.

Secondary hypotheses:

- 1. Singing groups for older people are more cost effective than usual activities
- 2. Singing groups for older people reduce anxiety and depression when compared with usual activities

Ethics approval required

Old ethics approval format

Ethics approval(s)

Added 16/03/2010:

Surrey Research Ethics Committee approved on the 20th January 2010 (ref: 10/H1109/5)

Study design

Single centre randomised unblinded controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

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

General health status

Interventions

Intervention: 90 - 120 minutes group singing and related activities, weekly for 12 weeks, with follow up measures for a further 12 weeks.

Control: Usual activities

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Health related quality of life measured with 12-item short form health survey (SF-12) at baseline, 12 weeks (post-intervention) and 24 weeks.

Secondary outcome measures

- 1. Health utility, measured with EQ-5D at baseline, 12 weeks (post-intervention) and 24 weeks
- 2. Anxiety and depression measured with Hospital Anxiety and Depression Scale (HADS) at baseline, 12 weeks (post-intervention) and 24 weeks
- 3. Service use questionnaire at baseline, 12 and 24 weeks

Overall study start date

01/01/2010

Completion date

31/12/2011

Eligibility

Key inclusion criteria

- 1. Community dwelling volunteers
- 2. Aged 60+ years, either sex
- 3. Able to give informed consent and complete questionnaires

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

240

Key exclusion criteria

Individuals unable to give informed consent or complete questionnaires.

Date of first enrolment

Date of final enrolment 31/12/2011

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Sidney De Haan Research Centre for Arts & Health
Folkestone
United Kingdom
CT20 1JG

Sponsor information

Organisation

Canterbury Christ Church University (UK)

Sponsor details

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Sponsor type

University/education

Website

http://www.canterbury.ac.uk/

ROR

https://ror.org/0489ggv38

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research (NIHR) (UK) - Research for Patient Benefit (RfPB) (ref: PB-PG-0408-16038)

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	protcol	28/02/2011		Yes	No