A pilot study to assess whether delivery of a health and well-being programme set within a community based football club is feasible and acceptable to hard to reach men

Submission date	Recruitment status	[X] Prospectively registered
05/08/2019	Stopped	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
22/08/2019	Stopped	Results
Last Edited	Condition category	Individual participant data
28/05/2021	Other	Record updated in last year

Plain English summary of protocol

Background and study aims

This study is designed to assess whether men who wouldn't usually access health programmes through the NHS or other health services find a programme that targets physical health and mental health and wellbeing acceptable. The 12-week health and wellbeing programme (called the 12th Man) is delivered through a community-based football club, Newcastle United Foundation. Previously published research and recently collected data from our research group suggest that community-based football clubs are a successful means of recruiting hard to reach groups of men. Hard to reach groups of men are those who do not usually engage with traditional health programmes provided by the NHS or other healthcare providers. In addition, these groups of men generally do not consider themselves unhealthy despite living unhealthy lives. Using the pride and passion for football in regions like the North East of England can help to improve the health of men from this population. That is what has inspired Newcastle United Foundations '12th Man' programme. The 12th Man programme has been developed taking in to account views and feedback from study participants who received this programme during a previous small scale study. Findings from our previous research have enabled researchers to develop a 12-week programme that targets physical, mental and social health elements designed specifically for men aged between 30 and 65 years. During the 12-week programme, participants will take part in behaviour change workshops that cover common health issues that men experience and lifestyle behaviours. These issues include: exercise, diet, stress, sleep, anger, happiness, social support and will teach self-help strategies. Participants will also take part in a variety of physical activity and exercise sessions that will provide an introduction to activities that they can access beyond the end of the programme. The activities are located around the region. The aim of this study is to assess whether participants like and are willing to complete the 12th Man programme.

Who can participate? Men aged 30 – 65 from the North East of England What does the study involve?

Participants are randomly allocated to one of two groups. The first group is the intervention (the 12th Man programme delivered face to face by a coach), and the second is a comparator group (the 12th man programme delivered online). Measurements are taken at four timepoints: at the start of the study, immediately after, 6 months after, and 12 months after the end of the programme. The researchers monitor their ability to recruit the number of participants required and whether participants remain in the study until the end. They also seek the views of participants to find out their thoughts about the programme including reasons why they did or did not find it useful and beneficial. They also assess changes to physical, mental and social health outcomes at the start and end of the programme. These changes include body mass index, blood pressure, hip to waist ratio and markers of health outcomes within blood samples. Mental health measures include self-efficacy, self-esteem and quality of life.

What are the possible benefits and risks of participating?

Benefits of taking part in this research for the participants may include improvements to physical and mental health and improvements in social connectedness. The results of this study will inform a larger trial investigating the long term implications of participating in the 12th Man intervention on hard to reach groups of men. This may in turn demonstrate a need for this intervention to be extended throughout other football clubs around the country which may ultimately impact the health of a large number of individuals. Risks may include: discomfort during blood sampling; musculoskeletal injury; cardiorespiratory complications; trips, slips and falls; and the potential for psychological damage. These have all been addressed in the attached risk assessment for this study and has been reviewed by the ethics panel.

Where is the study run from? Newcastle United Football Club (UK)

When is the study starting and how long is it expected to run for? January 2018 to July 2021

Who is funding the study?

- 1. Newcastle University (UK)
- 2. Newcastle United Foundation (UK)

Who is the main contact? Mr Oliver Bell oliver.bell@nufc.co.uk

Contact information

Type(s)

Scientific

Contact name

Mr Oliver Bell

ORCID ID

http://orcid.org/0000-0002-5789-9219

Contact details

St James Park
Newcastle Upon Tyne
United Kingdom
NE1 4ST
+44 (0)7746971855
oliver.bell@nufc.co.uk

Additional identifiers

EudraCT/CTIS numberNil known

IRAS number

ClinicalTrials.gov number
Nil known

Secondary identifying numbers 12thManpRCT

Study information

Scientific Title

A pilot randomised controlled trial to assess feasibility and acceptability of a behavioural intervention targeting physical and mental health of hard to reach men via a football club - the 12th Man

Acronym

12th Man Pilot RCT

Study objectives

A pilot randomised controlled trial to assess the feasibility and acceptability of the 12th Man intervention with hard to reach men. The intervention has been developed and delivered via a football club to reach and engage with men who would not ordinarily contact NHS services or health providers to address physical and mental health support needs.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 06/08/2019, Faculty of Medical Science Research Ethics Committee, part of Newcastle University's Research Ethics Committee (Medical Sciences Faculty Office, Faculty of Medical Sciences, Newcastle University, Newcastle Upon Tyne, NE2 4HH, UK; Tel: +44 (0)191 208 5301; Email: marjorie.holbrough@ncl.ac.uk), ref: 1713/13707

Study design

Single-centre two-armed randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Other

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet.

Health condition(s) or problem(s) studied

Physical and mental health

Interventions

The proposed study is not disease-specific. Men aged between 30 and 65 years who are free from any condition that would prevent or restrict them from participating in physical activity are the focus of this intervention. The purpose of the intervention is to improve physical and mental health.

Participants will be randomly allocated 1:1 to the intervention or comparator group using statistical software with random block sizes of 2, 4 and 6. Randomisation will be completed by an individual from outside of the research team.

Intervention arm: The 12th Man intervention is a 12-week behavioural programme that targets physical and mental health and wellbeing in hard to reach men. The 12 weekly sessions will each last 2 hours in duration and will be delivered face-to-face in a group setting. They will each involve a one hour workshop and a one-hour physical activity session utilising local facilities and clubs. In addition, there will be a 6-month reunion meeting held post completion of the programme.

During week one, each participant will be given a copy of the intervention workbook that provides a brief overview of each weekly session. The workbook also includes interactive tasks to complete during and/or outside of sessions. Participants are advised that the aim is to complete the workbook by the end of the intervention.

Participants are also invited to join the Facebook Group for their individual 12th Man cohort and the group for all 12th Man participants.

Comparator arm: The 12th Man intervention delivered online

The comparator group will receive the 12th Man intervention delivered online without face-to-face weekly contact with the coach or other participants. Participants will be invited to the same study visits as the intervention group to allow the collection of outcome data and these data collections will take place in various NUFC buildings including the football stadium and the training ground. These locations are intended to increase adherence to outcome measurement sessions.

The comparator group will receive 12 weekly videos emailed to them (one per week). The sessions will be recorded prior to the beginning of the 12th Man intervention and will be generic – i.e. they will not be tailored in the same way as the 12th Man intervention delivered face-to-face, however, participants will be able to use the content in a tailored fashion supported by their workbooks that will be given during week one of the intervention. The videos will cover all topics delivered to the participants in the intervention group.

In addition to the 12 weekly videos, comparator group participants will also be added to WhatsApp and Facebook groups (subject to consent). Here they will be updated with all the physical activity opportunities available to them in their local community, similar to the intervention group. They will be encouraged to access these but will not be guided by a weekly coach.

Fasted (> 6 hours) venous blood samples will be taken by a trained phlebotomist using a 2 x 10ml vacutainer tubes treated with di-potassium ethylene diamine tetra-acetic acid (EDTA) and serum using standard operating procedure. Samples will be stored at 4 C (either refrigerator or cool bag with ice) until processing at a laboratory within 24 hours. All blood samples will be centrifuged at 3000 rpm for 20 minutes at 4 C with the corresponding plasma and serum supernatant transferred to aliquots and stored at -80°C until later analysis.

12-month post-intervention follow-up (i.e. a 15-month study period).

Intervention Type

Behavioural

Primary outcome measure

- 1. Feasibility and acceptability:
- 1.1. Feasibility assessed by recording recruitment, retention and adherence rates of study participants including willingness to be randomised
- 1.2. Acceptability assessed qualitatively via focus group discussions upon completion of the study

Outcomes measured at baseline, immediately post intervention, and 6 and 12 months post intervention

Secondary outcome measures

Measured at baseline, immediately post intervention, and 6 and 12 months post intervention:

- 1. Fasting measures of plasma, serum or whole blood insulin, glucose, non-esterfied fatty acids, interleukin-6, tumour necrosis factor alpha, C-reactive protein, haemoglobin A1c, triglycerides and cholesterol
- 2. Body height (cm) and body weight (kg) collected to analyse body mass index (BMI)
- 3. Diastolic and systolic blood pressure (mmHg) measured with an automated blood pressure monitor. A body tape measure will be used to measure hip to waist ratio
- 4. Self-reported physical activity measured using IPAQ-short form (Ingledew and Markland., 2008)
- 5. Planning and coping planning of behaviour change measured using self-reported questionnaire adapted from Sniehotta, Scholz and Schwarzer., 2003
- 6. Behaviour regulations in exercise measured using BREQ-2 (Markland and Tobin 2004)
- 7. Quality of life measured using EQ-5D-5L (Herdman et al., 2011)
- 8. General self-efficacy measured using self-reported questionnaire from Jerusalem and Schwarzer 1979
- 9. Self-esteem measured using self-reported questionnaire from Rosenberg, 1986

10. Perceived and received social support assessed through questionnaires adapted from previous research (Freeman and Rees., 2008)

Overall study start date

01/01/2018

Completion date

01/07/2021

Reason abandoned (if study stopped)

Participant recruitment issue

Eligibility

Key inclusion criteria

- 1. Men
- 2. Aged 30 65 years
- 3. Consent to randomisation

Participant type(s)

Other

Age group

Adult

Sex

Male

Target number of participants

92

Key exclusion criteria

- 1. Failure to return pre-screening questionnaire
- 2. Condition or disability which would prevent participation in physical activity or exercise

Date of first enrolment

01/03/2021

Date of final enrolment

01/06/2021

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Newcastle United Football Club

St James Park Barrack Road Newcastle Upon Tyne United Kingdom NE1 4ST

Sponsor information

Organisation

Newcastle University

Sponsor details

Institute of Cellular Medicine
William Leech Building
Medical School
Newcastle upon Tyne
England
United Kingdom
NE2 4HH
+44 (0)191 208 7865
emma.stevenson@newcastle.ac.uk

Sponsor type

University/education

Website

http://www.ncl.ac.uk/

ROR

https://ror.org/01kj2bm70

Organisation

Newcastle United Foundation

Sponsor details

St James Park
Barrack Road
Newcastle Upon Tyne
United Kingdom
NE1 4ST
+44 (0)191 201 8451
steve.beharall@nufc.co.uk

Sponsor type

Charity

Website

https://nufoundation.org.uk/

Funder(s)

Funder type

University/education

Funder Name

Newcastle University

Alternative Name(s)

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

United Kingdom

Funder Name

Newcastle United Foundation

Results and Publications

Publication and dissemination plan

Findings from the Pilot RCT will be communicated to local, regional and national groups and via social media.

Throughout the trial period findings will be disseminated at academic and professional conferences.

Intention to publish date

01/11/2021

Individual participant data (IPD) sharing plan

The data sharing plans for the current study are unknown and will be made available at a later date.

IPD sharing plan summary
Data sharing statement to be made available at a later date