# Developing a way of improving the prescribing of many drugs for older people who live in their own home and are cared for by general practitioners

Submission date	Recruitment status No longer recruiting	<ul><li>Prospectively registered</li></ul>			
18/01/2016		Protocol			
Registration date	Overall study status	Statistical analysis plan			
22/01/2016	Completed	[X] Results			
<b>Last Edited</b> 16/02/2023	<b>Condition category</b> Other	[] Individual participant data			

#### Plain English summary of protocol

Background and study aims

In the last century, advances in medicine have led to people living much longer as previously fatal diseases are now treatable. Many people, particularly those over the age of 60, are living with two or more long-term medical conditions (multimorbidity). These patients often need to take a range of different medications to control their conditions (polypharmacy). In some cases, the prescribing of medications may not be appropriate and patients are given too much or too little of what they actually need. Many studies have tried to improve this and ensure that patients are prescribed all the medicines they actually need, however many have been unsuccessful. This may be because studies have not considered the views and opinions of patients or the way that health professionals make decisions about which medicines are needed. The aim of this study is to try to design a programme in order to improve polypharmacy prescribing practices of GP's for older people.

### Who can participate?

GP practices in Northern Ireland willing to take part in the study, the GPs who work there and adult patients over the age of 65 who take at least 4 different medications.

#### What does the study involve?

GPs who work at participating practices that consent to take part in the study are given access to an online video which shows how to best prescribe multiple medications to older patients in a normal consultation. The video also contains feedback about why this approach works well. Appropriate patients are then invited to come in to see their GP for an appointment to review their medications. After each consultation, both the GPs and the patients complete anonymous feedback questionnaires to provide their opinions about the consultations. The results of these questionnaires are then reviewed to find out how acceptable the approach is. There are no specific follow-up appointments, but for the patients taking part, their medical records are reviewed 4 weeks later in order to see if there has been any change to their medication following the consultation.

What are the possible benefits and risks of participating? There are no direct benefits or risks to participants taking part in the study.

Where is the study run from?

The study is run from Queen's University Belfast and takes place in two GP practices in Northern Ireland (UK)

When is the study starting and how long is it expected to run for? April 2015 to September 2015

Who is funding the study? Dunhill Medical Trust (UK)

Who is the main contact? Prof. Carmel Hughes

# Contact information

#### Type(s)

Scientific

#### Contact name

**Prof Carmel Hughes** 

#### Contact details

Queen's University Belfast School of Pharmacy 97 Lisburn Road Belfast United Kingdom BT9 7BL

# Additional identifiers

Protocol serial number

2

# Study information

#### Scientific Title

Development of an intervention to improve appropriate polypharmacy for older people in primary care

#### **Study objectives**

The overall aim of this project is to develop and feasibility test an intervention to improve the prescribing of appropriate polypharmacy for older people in primary care, focusing on healthcare professionals' (HCPs') encounters with patients in daily practice.

#### Ethics approval required

#### Old ethics approval format

#### Ethics approval(s)

Health and Social Care Research Ethics Committee A, 29/05/2015, ref: 15/NI/0104

#### Study design

Single-centre feasibility study

#### Primary study design

Interventional

#### Study type(s)

Other

#### Health condition(s) or problem(s) studied

Polypharmacy in older people

#### **Interventions**

A feasibility study is conducted in order to assess the usability and acceptability of an intervention to improve the prescribing of appropriate polypharmacy for older patients in primary care. The intervention consists of a video demonstration of how general practitioners (GPs) can prescribe appropriate polypharmacy during a typical consultation with older patients and will be delivered as an online video. The video also includes feedback emphasising the positive outcomes of this type of approach.

During the recruitment phase of the study, a meeting is held between the researcher and GPs within each practice. During this meeting GPs are provided with an overview of the feasibility study protocol and intervention. Following the provision of written informed consent, GPs are provided with access to the online video, before information letters are issued by the practice to patients meeting inclusion criteria, inviting them to attend a scheduled consultation with their GP about their medicines.

Baseline assessments are conducted on the date that each practice chooses to invite patients for scheduled consultations with recruited GPs. These assessments involve the collection of data from three sources: GPs, patients and recruited patients' medical records. GPs complete a feedback questionnaire after they perform medication reviews during scheduled consultations with recruited patients. Recruited patients are also asked to complete a feedback questionnaire after the scheduled consultations with their GPs. There is no direct follow-up with GP or patient participants after the scheduled consultations are completed. Data is also extracted from recruited patients' medical records at baseline and follow-up (one month after the scheduled consultation). The extracted data includes details of patient demographics (i.e. age, gender), clinical conditions and prescribed medications (both acute list items and repeat list items).

#### Intervention Type

Behavioural

## Primary outcome(s)

- 1. The usability and acceptability of the intervention for GPs is determined using a feedback questionnaire which recruited GPs complete at baseline (after medication reviews during scheduled consultations)
- 2. The usability and acceptability of the intervention for patients is determined using a feedback

questionnaire which recruited patients complete at baseline (after medication reviews during scheduled consultations)

## Key secondary outcome(s))

- 1. Feasibility of recruitment procedures is assessed by determining whether recruitment targets have been met at the end of the study
- 2. Feasibility of the data collection procedures is determined by applying validated assessments of prescribing appropriateness (i.e. STOPP/START criteria, Medication Appropriateness Index) and prescribing regimen complexity (i.e. Medication Regimen Complexity Index) to clinical data extracted from recruited patients' medical records at baseline and follow-up (four weeks after date of scheduled consultations)

#### Completion date

30/09/2015

# **Eligibility**

#### Key inclusion criteria

General practice inclusion criteria:

- 1. Located in Northern Ireland
- 2. The practice was willing to implement the study
- 3. Individual general practitioners (GPs) willing to provide written informed consent.

#### Patient inclusion criteria:

- 1. Over the age of 65
- 2. Receiving four or more regular medicines
- 3. Not cognitively impaired
- 4. Resident in the community

#### Participant type(s)

Patient

## Healthy volunteers allowed

No

#### Age group

Senior

#### Sex

All

#### Key exclusion criteria

- 1. Not meeting the inclusion criteria
- 2. Not proving written informed consent

#### Date of first enrolment

01/06/2015

#### Date of final enrolment

31/08/2015

# Locations

#### Countries of recruitment

**United Kingdom** 

Northern Ireland

## Study participating centre Queen's University Belfast

School of Pharmacy 97 Lisburn Road Belfast United Kingdom BT9 7BL

# Sponsor information

#### Organisation

Queen's University Belfast

#### **ROR**

https://ror.org/00hswnk62

# Funder(s)

#### Funder type

Hospital/treatment centre

#### **Funder Name**

**Dunhill Medical Trust** 

#### Alternative Name(s)

The Dunhill Medical Trust, Dunhill Medical Trust, DunhillMedical, DMT

#### Funding Body Type

Private sector organisation

## **Funding Body Subtype**

Trusts, charities, foundations (both public and private)

#### Location

**United Kingdom** 

# **Results and Publications**

# Individual participant data (IPD) sharing plan

Not provided at time of registration

# IPD sharing plan summary

Not expected to be made available

# **Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	20/07/2017		Yes	No
HRA research summary			28/06/2023	No	No
Other publications	Development of intervention	16/11/2016	16/02/2023	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes