The safety in primary care study: a randomised, controlled feasibility study

Submission date	Recruitment status	Prospectively registered		
08/06/2018	No longer recruiting Overall study status	[X] Protocol		
Registration date		Statistical analysis plan		
12/06/2018	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
01/02/2019	Other			

Plain English summary of protocol

Background and study aims

Research on patient safety has focused largely on hospital settings, and there is limited knowledge about patient safety in primary care. This study aims to assess the effectiveness of an intervention to improve patient safety in primary care. The purpose of this study is to evaluate whether a future larger study of the patient safety intervention should be carried out.

Who can participate?

Staff working in general practices in the Republic of Ireland and in Northern Ireland

What does the study involve?

Participating practices are randomly allocated to one of two groups to either receive the intervention over a 9-month period or to continue care as usual. The interventions include repeated completion of a safety questionnaire and feedback on these findings, and the review of patient records to identify any patients who may have been harmed as part of their care.

What are the possible benefits and risks of participating?

The findings from the study may increase knowledge of how often patients are harmed in primary care, contribute to improved patient safety practices in primary care, and inform future research on patient safety improvement. The benefits of participating are that there will be an increased awareness of patient safety in the practice that may lead to improved patient care. The risks are that participants may become distressed if they realise they contributed to a patient safety incident.

Where is the study run from?

- 1. NUI Galway (Ireland)
- 2. Queen's University Belfast (UK)

When is the study starting and how long is it expected to run for? September 2015 to June 2017

Who is funding the study? Health Research Board (Ireland)

Who is the main contact? Dr Paul O'Connor paul.oconnor@nuigalway.ie

Contact information

Type(s)

Scientific

Contact name

Dr Paul O'Connor

Contact details

Discipline of General Practice NUI Galway 1 Distillery Road Galway Ireland H91 TK33 +353 (0)91492897 paul.oconnor@nuigalway.ie

Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

The safety in primary care study: a randomised, controlled feasibility study

Acronym

SAP-C

Study objectives

The purpose of this feasibility study is to evaluate the:

- 1. Willingness of practices to participate in the study
- 2. Retention of control and intervention practices
- 3. Response rates to questionnaires
- 4. Feedback from the intervention group on the feasibility, usefulness, and sustainability of the intervention
- 5. Effects of the intervention on safety climate

Ethics approval required

Old ethics approval format

Ethics approval(s)

- 1. Irish College of General Practitioners' Research Ethics Committee, 14/01/2016
- 2. Office for Research Ethics Committees of Northern Ireland, 23/02/2016, ref: 16/NI/0008

Study design

Cluster randomised controlled trial

Primary study design

Interventional

Study type(s)

Other

Health condition(s) or problem(s) studied

General practice

Interventions

Simple randomisation was employed, whereby participating practices were assigned to either the intervention or control group.

Five practices received the intervention over a 9-month period. The intervention consisted of: 1) repeated safety climate (SC) measurement (using GP-SafeQuest questionnaire) and feedback, and 2) patient record reviews using a specialised trigger tool to identify instances of undetected patient harm.

The five practices in the control group continued care as usual.

Intervention Type

Behavioural

Primary outcome(s)

The evaluation of the study's implementation process was the primary outcome. Outcomes of interest were:

- 1. Willingness of practices to participate in the study, measured during the recruitment phase prior to the intervention
- 2. Response rates to safety climate questionnaire, measure at baseline and month 9
- 3. Feasibility questionnaire at month 9
- 4. Retention of control and intervention practices, measured at month 9
- 5. interviews on the feasibility, usefulness, and sustainability of the intervention at month 9

Key secondary outcome(s))

None

Completion date

30/06/2017

Eligibility

Key inclusion criteria

Staff working in general practices in the Republic of Ireland or Northern Ireland that have expressed a willingness to be involved in the study

Participant type(s)

Health professional

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

People who do not work in a general practice that agreed to participate in the study

Date of first enrolment

01/02/2016

Date of final enrolment

21/03/2016

Locations

Countries of recruitment

United Kingdom

Northern Ireland

Ireland

Study participating centre Discipline of General Practice

NUI Galway Galway Ireland H91 TK33

Study participating centre Department of General Practice and Primary Care

Queen's University Belfast Belfast United Kingdom BT9 7HR

Sponsor information

Organisation

HRB Primary Care Clinical Trials Network Ireland

ROR

https://ror.org/003hb2249

Funder(s)

Funder type

Research organisation

Funder Name

Health Research Board

Alternative Name(s)

Health Research Board, Ireland, An Bord Taighde Sláinte, HRB

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

Ireland

Results and Publications

Individual participant data (IPD) sharing plan

Requests for access to data should be made to Dr Paul O'Connor (paul.oconnor@nuigalway.ie). The data will be available from December 2018. Data that could be made available is anonymous safety climate data, and the feasibility questionnaire data. Sharing data was not explicitly mentioned in the consent form. Therefore, requests for data sharing will have to be considered on a case-by-case basis by the members of the trial steering committee.

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	30/01/2019	01/02/2019	Yes	No
Protocol article	protocol	16/09/2016		Yes	No
HRA research summary			28/06/2023		No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes