ISRCTN61739603 https://doi.org/10.1186/ISRCTN61739603

Seniors Health Investigation Network - Blood Pressure Monitoring Program

Submission date	Recruitment status	[_] Prospectively registered
27/05/2005	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
01/06/2005	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
15/04/2008	Circulatory System	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Acronym
SHINE - Dundas Pilot

Study objectives
To determine the best method (mail or telephone) for
family physicians offices to invite patients to attend a session, and to identify factors that predicted attendance. In preparation for a larger scale randomized controlled trial, we also performed a cost analysis of invitation method and success rate, investigated the operational and methodological aspects of the proposed intervention, and queried patients willingness to continue to attend community pharmacy blood pressure sessions and their preferences for the time and frequency of the sessions.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Single-centre

Study setting(s)
GP practice

Study type(s)
Diagnostic

Participant information sheet

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

Interventions
Regular patients 65 years or older (n = 235) were
randomly allocated to invitation by mail or telephone to attend pharmacy cardiovascular health awareness sessions led by volunteer peer health educators.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
The primary end-point
was the overall attendance among patients invited by mail compared to those invited by telephone.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/03/2001

Completion date
27/04/2001

Eligibility

Key inclusion criteria
Patients were eligible to participate if they were 65
years of age or older, considered by the physician to be regular patients, community dwelling, and able to attend a cardiovascular health promotion session in a local pharmacy.

Participant type(s)
Patient

Age group
Senior

Sex
Both

Target number of participants
235

Key exclusion criteria
Patients were excluded if
they suffered from dementia or a serious, non-cardiovascular disease or condition, or were non-English
speaking and could not attend with an English-speaking companion.

Date of first enrolment
01/03/2001

Date of final enrolment
27/04/2001

Locations

Countries of recruitment
Canada

Study participating centre
1200 Main Street West HSC-2V12
Hamilton
Canada
L8N 3Z5

Sponsor information

Organisation
Biovail Pharmaceuticals Canada, formerly Crystaal Corporation

Sponsor details
7150 Mississauga Road
Mississauga, Ontario
Canada
L5N 8M5

Sponsor type
Industry

ROR
https://ror.org/01zgpn844

Funder(s)

Funder type
Industry

Funder Name
Biovail Corporation Canada, formerly Crystaal Corporation

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?
Results article	Results:	19/08/2005		Yes	No