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Can the use of a Link-Worker improve Attendance for Diabetic Retinal Screening in the Asian population of Coventry and Warwickshire?

Submission date
22/11/2007

Recruitment status
No longer recruiting

Registration date
22/02/2008

Overall study status
Completed

Last Edited
22/02/2012

Condition category
Nutritional, Metabolic, Endocrine

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
1

Study information

Scientific Title

Acronym
LWADRS

Study objectives
The use of a link-worker will improve attendance for Diabetic Retinal Screening in the Asian population of Coventry and Warwickshire.

Ethics approval required
Old ethics approval format

Ethics approval(s)
No formal approval is required as this trial is an operational research.

Study design
Group asssigned randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Screening

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
Diabetic retinopathy

Interventions
The participating GP surgeries are randomised to the intervention and control groups.

Intervention group: A link worker is assigned to each participating surgery. All patients who do not attend their first screening visit have a second visit arranged. In the 48 hours prior to this the link worker contacts the patient to ensure they are aware of the appointment and remind them to attend. The link worker takes on an educational and facilitator role.

Control group: Usual care only

Duration of intervention will depend on recruitment at each surgery and the care required by participants.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Attendance at retinal screening

Secondary outcome measures
No secondary outcome measures

Overall study start date
01/07/2007

Completion date
01/07/2008

Eligibility

Key inclusion criteria
All patients attending retinal screening service in Coventry and Warwickshire

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
4 GP surgeries - target number 400 patients

Key exclusion criteria
None

Date of first enrolment
01/07/2007

Date of final enrolment
01/07/2008

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Hospital of St Cross
Rugby
United Kingdom
CV22 5PX

Sponsor information

Organisation
University of Warwick, Medical School (UK)

Sponsor details
-
Coventry
England
United Kingdom
CV4 7AL

Sponsor type
University/education

Website
http://www2.warwick.ac.uk

ROR
https://ror.org/01a77tt86

Funder(s)

Funder type
University/education

Funder Name
University of Warwick, Medical School (UK)

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	01/12/2004		Yes	No