

A randomised controlled trial of health visitor safety advice plus low cost safety equipment for families living in deprived areas

Submission date
23/01/2004

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
23/01/2004

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
06/02/2019

Condition category
Injury, Occupational Diseases, Poisoning

☐ 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

RBK99XX4

Study information

Scientific Title

A randomised controlled trial of health visitor safety advice plus low cost safety equipment for families living in deprived areas

Study objectives

To assess the effectiveness of health visitor safety advice coupled with access to low cost safety equipment for low income families with children under 5, living in deprived areas.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval as of 06/02/2019:

Queen's Medical Centre, Nottingham University Hospital NHS Trust Ethics Committee, 13/07/1999, ref. EX039903.

Previous ethics approval:

Queen's Medical Centre Research Ethics Committee, Nottingham

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Prevention

Participant information sheet

Health condition(s) or problem(s) studied

Injury

Interventions

The intervention group will receive a 20 min safety consultation with the health visitor based on the principles used in adult learning, plus opportunistic reinforcement of safety education at future consultations. Low cost equipment available will include stair gates, fireguards, smoke alarms, cupboard locks and window catches. Control group received standard care.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Primary outcome measure: the proportion of families in which at least one child experiences a medically attended unintentional injury (primary and secondary care).

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/11/1999

Completion date

30/04/2002

Eligibility**Key inclusion criteria**

All families with children under 5 in practices with a Townsend score of >0.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/1999

Date of final enrolment

30/04/2002

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Division of General Practice

Nottingham

United Kingdom

NG7 2UH

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details

The Department of Health

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

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dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

NHS Executive Trent (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	01/06/2003		Yes	No
Results article	results	01/12/2004		Yes	No
Results article	results	22/01/2005		Yes	No
Results article	results	01/11/2005		Yes	No
Results article	results	01/12/2005		Yes	No
Results article	results	01/05/2006		Yes	No