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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

Registration date
23/01/2004

Overall study status
Completed

Last Edited
06/02/2019

Condition category
Injury, Occupational Diseases, Poisoning

[ ] 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

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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
+44 (0)20 7307 2622
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