A randomised controlled trial of an injury prevention programme in primary schools

Submission date	Recruitment status No longer recruiting	Prospectively registered	
23/01/2004		☐ Protocol	
Registration date 23/01/2004	Overall study status Completed	Statistical analysis plan	
		[X] Results	
Last Edited	Condition category	Individual participant data	
08/08/2011	Injury, Occupational Diseases, Poisoning		

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers MCH 10-08

Study information

Scientific Title

Study objectives

The aim of the study is to reduce school and sports and leisure injuries, reduce hazards and increase injury prevention awareness.

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

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Injury, occupational diseases, poisoning: Musculoskeletal injury; Injury, occupational diseases, poisoning: Burns

Interventions

An injury prevention advocate will work with schools and the local injury prevention alliance, with emphasis on locally appropriate environmental interventions, advocacy and training. An evaluation officer will conduct evaluations of process, impact and outcome, using risk questionnaires to measure impact of the study and the level of self-reported injuries and a local surveillance system to measure health outcomes.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

To implement a range of outcome, impact and process measures for the evaluation of the programme and to develop a package of materials that highlight areas of good practice within schools that can be replicated elsewhere.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/10/1996

Completion date

01/10/2001

Eligibility

Key inclusion criteria

The two year intervention will target 7 and 9 year old school pupils.

Participant type(s)

Patient

Age group

Child

Lower age limit

7 Years

Upper age limit

9 Years

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

01/10/1996

Date of final enrolment

01/10/2001

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Department of Child Health
Newcastle Upon Tyne
United Kingdom
NE2 4HH

Sponsor information

Organisation

Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

Sponsor details

The Department of Health Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

http://www.doh.gov.uk

Funder(s)

Funder type

Government

Funder Name

NHS Mother and Child Health National Research and Development Programme (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	01/03/1997		Yes	No