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

Submission date	Recruitment status	Prospecti
23/01/2004	No longer recruiting	[] Protocol
Registration date	Overall study status	[] Statistica
23/01/2004	Completed	[X] Results
Last Edited 08/08/2011	Condition category Injury, Occupational Diseases, Poisoning	[_] Individua

] Prospectively registered

] Statistical analysis plan

] Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s) Scientific

Contact name Dr Elizabeth Towner

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 **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