# A randomised controlled trial of the effectiveness of an education package in reducing baby walker use

Submission date	Recruitment status	Prospectively registered		
23/01/2004	No longer recruiting	☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
23/01/2004	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
06/02/2019	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

RBK99XX7

# Study information

#### Scientific Title

A randomised controlled trial of the effectiveness of an education package in reducing baby walker use

#### **Study objectives**

Is an education package delivered by midwives and health visitors aimed at reducing baby walker use effective?

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Ethical approval as of 06/02/2019:

Queen's Medical Centre, University Hospital NHS Trust Ethics Committee, 13/07/1999, ref. EX029902.

North Nottinghamshire Local Research Ethics Committee, 12/06/2000, ref. NNHA/494.

#### Previous ethical approval:

Queen's Medical Centre, Nottingham, ref. EX029902.

# Study design

Randomised controlled trial

# Primary study design

Interventional

# Secondary study design

Randomised controlled trial

# Study setting(s)

GP practice

## Study type(s)

Prevention

## Participant information sheet

# Health condition(s) or problem(s) studied

Pediatric injury

#### **Interventions**

i. Evidence based educational package developed to reduce baby walker use. The package will be presented at one ante-natal consultation from 28 weeks (by midwives) and in 2 post-natal

consultations, at the birth review and 3-4 month hip check (by health visitors). ii. No exposure to package.

#### **Intervention Type**

Other

#### **Phase**

**Not Specified** 

#### Primary outcome measure

The primary outcome measure is prevalence of baby walker possession and use.

#### Secondary outcome measures

Not provided at time of registration

#### Overall study start date

01/10/1999

#### Completion date

30/04/2002

# **Eligibility**

#### Key inclusion criteria

The study population will comprise all pregnant women (of at least 28 weeks gestation) registered with 34 participating practices.

## Participant type(s)

**Patient** 

#### Age group

Adult

#### Sex

Female

## Target number of participants

Not provided at time of registration

#### Key exclusion criteria

Not provided at time of registration

#### Date of first enrolment

01/10/1999

#### 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	01/03/2003		Yes	No
Results article	results	01/03/2003		Yes	No
Results article	results	01/09/2003		Yes	No
Results article	results	01/12/2003		Yes	No
Results article	results	01/08/2005		Yes	No
Results article	results	01/09/2006		Yes	No
Results article	results	01/06/2018		Yes	No