

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

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

Protocol serial number

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.

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Prevention

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(s)

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

Key secondary outcome(s)

Not provided at time of registration

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

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

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

United Kingdom

England

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)

Funder(s)

Funder type

Government

Funder Name

NHS Executive Trent (UK)

Results and Publications

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