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The effects of parenting intervention on quality of life and parental confidence in management of young asthmatic children

Submission date 21/10/2010	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 21/10/2010	**Overall study status** Completed	
Last Edited 18/04/2017	**Condition category** Respiratory	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
8624

Study information

Scientific Title

The effects of parenting intervention on quality of life and parental confidence in management of young asthmatic children

Study objectives
The study will be a randomised controlled trial comparing care as usual versus care as usual supplemented by group-delivered Triple P parenting seminars and telephone support in parents /carers of 2 - 7 year old asthmatic children.

Ethics approval required
Old ethics approval format

Ethics approval(s)
MREC approved, ref: 09/H1017/115

Study design
Single-centre randomised interventional treatment pilot/feasibility study

Primary study design
Interventional

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Topic: Respiratory; Subtopic: Respiratory (all Subtopics); Disease: Respiratory

Interventions
Triple P parenting seminars and telephone support.

Follow-up length: 9 months
Study entry: registration only

Intervention Type
Other

Phase
Not Applicable

Primary outcome(s)
Paediatric Asthma Caregiver's Quality of Life Questionnaire (PACQLQ), measured at 3 and 9 months

Key secondary outcome(s))
Not provided at time of registration

Completion date
09/01/2012

Eligibility

Key inclusion criteria
1. Parents of children with frequent episodic or chronic persistent asthma who are taking regular preventive medication
2. Parents must have an asthmatic child aged 2 - 7 years, either sex
3. Diagnosis of asthma in children, based on:
3.1. An observation of recurrent episodes of wheeze (preferably witnessed by a medical observer)
3.2. A clinical response to a bronchodilator medication (eg. Salbutamol)
3.3. An absence of other rarer lung conditions (e.g., cystic fibrosis, chronic neonatal lung disease) which may account for these symptoms

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Child

Lower age limit
2 years

Upper age limit
7 years

Sex
All

Key exclusion criteria
1. The child has a significant disability
2. The parents are currently seeing a professional for the child's behaviour difficulties, are currently receiving psychological help or counselling, or have significant learning difficulties. For the purposes of the study, in order to participate in the seminars, parents will be excluded if they don't understand English.

Date of first enrolment
01/06/2010

Date of final enrolment
09/01/2012

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Manchester Royal Infirmary
Manchester
United Kingdom
M13 9WL

Sponsor information

Organisation
Central Manchester and Manchester Children's University Hospital (CMMCUH) NHS Trust (UK)

ROR
https://ror.org/00he80998

Funder(s)

Funder type
Government

Funder Name
National Institute for Health Research (NIHR) (UK) Research for Patient Benefit (RfPB) programme

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration