

Development and validation of a caregiver quality of life questionnaire for wheezy preschool children

Submission date 24/06/2010	Recruitment status No longer recruiting	☐ Prospectively registered
		☐ Protocol
Registration date 24/06/2010	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 30/09/2016	Condition category Respiratory	☐ Individual participant data
		☐ Record updated in last year

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
7819

Study information

Scientific Title

Development and validation of a caregiver quality of life questionnaire for wheezy preschool children

Acronym

QOL

Study objectives

To establish a valid and reliable measure of health-related quality of life for caregivers of wheezy preschool children.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Kent Research Ethics Committee, July 2006, ref: 06/Q1905/8

Primary study design

Interventional

Study design

Single centre non-randomised interventional treatment trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Topic: Respiratory; Subtopic: Respiratory (all Subtopics); Disease: Respiratory

Interventions

Item reduction stage:

Asking carers of wheezy preschool children to complete a quality of life (QOL) questionnaire plus two data sheets. It takes about half an hour to complete the questionnaire.

Validation stage:

The final questionnaire will be given to parents to complete on three occasions (over approximately 8 weeks):

1. At first clinic visit
2. 1 - 2 weeks later before any change in treatment has been made
3. 6 weeks after a change in asthma treatment intended to bring the asthma symptoms under control

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

To have a validated quality of life questionnaire for caregivers of wheezy preschool children

Key secondary outcome(s)

No secondary outcome measures

Completion date

31/12/2010

Eligibility

Key inclusion criteria

1. Boys and girls
2. History of 2 or more wheezing episodes
3. Age 1 year to 5 years 11 months

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

1 Years

Upper age limit

5 Years

Sex

All

Key exclusion criteria

Other major medical problem

Date of first enrolment

01/01/2010

Date of final enrolment

31/12/2010

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Royal Sussex County Hospital
Brighton
United Kingdom
BN2 5BE

Sponsor information

Organisation

Brighton and Sussex University Hospitals NHS Trust (UK)

Funder(s)

Funder type

Charity

Funder Name

Asthma UK (UK)

Alternative Name(s)

asthmalunguk, Asthma + Lung UK, Asthma UK

Funding Body Type

Private sector organisation

Funding Body Subtype

Research institutes and centers

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary