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Parental satisfaction with interventions for childhood ADHD/hyperkinetic disorder

Submission date	Recruitment status	[ ] Prospectively registered
12/09/2003	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
12/09/2003	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
16/06/2014	Mental and Behavioural Disorders	[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr Cheyvonne Ogunde

Contact details
Calderdale and Huddersfield NHS Trust
Huddersfield Royal Infirmary
Lindley
Huddersfield
United Kingdom
HD3 3EA

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0083122677

Study information

Scientific Title

Study objectives
Satisfaction outcomes will be different for children with hyperkinetic disorders who receive combination treatment compared with medication alone.

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)
Treatment

Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Mental and Behavioural Disorders: Attention-deficit/hyperactivity disorder (ADHD)

Interventions
This is a pilot randomised controlled trial (RCT) to determine whether there is any difference in parental satisfaction (measured during a brief interview and using the 'treatment evaluation inventory') between those having medication alone and those having behavioural modification therapy and medication in combination.

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure
Parental satisfaction

Secondary outcome measures
Not provided at time of registration

Overall study start date
25/11/2002

Completion date
24/11/2003

Eligibility

Key inclusion criteria
60 families

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
60

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
25/11/2002

Date of final enrolment
24/11/2003

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Calderdale and Huddersfield NHS Trust
Huddersfield
United Kingdom
HD3 3EA

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
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
Calderdale and Huddersfield NHS Trust (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