

Snuggledown - Use of sensory blankets for children with autistic spectrum disorder

Submission date 10/08/2011	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 10/08/2011	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 23/06/2015	Condition category Nervous System Diseases	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

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
10090

Study information

Scientific Title
Snuggledown - The use of sensory weighted blankets in children with autistic spectrum disorders and poor sleep: a randomised crossover study

Study objectives

The objective of this trial is to confirm (or refute) that a specially designed weighted blanket is effective in improving total duration of night-time sleep compared to a similar but non-weighted blanket in children with autistic spectrum problems

No randomised clinical trials (RCT) have been published. Previous studies of weighted jackets have not reported any side effects. Care is required to ensure that the child is free to move /remove blanket at any point.(MCF, 2008)

Anecdotal parent reported benefits include a reduced sleep latency time (ie: reduced time to fall asleep) and reduced number of awakenings throughout the night (ie: increased periods of continuous, un-interrupted sleep throughout the night).

Ethics approval required

Old ethics approval format

Ethics approval(s)

South East London REC2, 24/08/2010, ref: 10/H0802/57

Study design

Randomised interventional treatment trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Sleep disorders in paediatrics

Interventions

Sensory blanket (Southpaw Sensory Integration blanket)

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Total sleep time based on actigraphy two weeks after each cross-over period

Key secondary outcome(s))

1. Sleep latency and efficiency (actigraphy)
2. Total sleep time and sleep latency by sleep diary
3. Parental assessment of sleep according to composite sleep disturbance index
4. Changes in behaviour (Aberrant Behaviour Checklist questionnaire)
5. Child's Visual Analogue Scale (VAS) perspective of sleep quality and acceptability of blanket
6. Sensory Behaviour Questionnaire (Green 2009)

Completion date

01/10/2012

Eligibility

Key inclusion criteria

1. Children aged 5 to 15 years and 10 months with a diagnosis of an autistic spectrum disorders (ASD), diagnosed by a community paediatrician, paediatric neurologist or paediatric neurodisability consultant
2. No plans to commence new medication known to influence behaviour or sleep
3. Diagnosis of impaired sleep as defined by a) not falling asleep within one hour of 'lights off' or 'snuggling down to sleep' at age-appropriate times and/or b) less than 7 hours of continuous sleep in 3 out of 5 nights
4. A total score of <4 on questions 18 to 20 on the Children's Sleep Habits Questionnaire indicating the likely absence of sleep apnoea
5. Parental English language sufficient to read and complete questionnaires and sleep diaries
6. Ability to start each night's sleep in own bed
7. Male and female participants

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

5 years

Upper age limit

15 years

Sex

All

Key exclusion criteria

1. Children who are using a weighted blanket
2. Children whose parents are unlikely to be able to use the actigraph or complete sleep diaries or both
3. Children where there may be a problem of major non-concordance with blanket or actigraph
4. Currently participating in a conflicting clinical study
5. Epilepsy (uncontrolled or medication changes)
6. Cerebral palsy

Date of first enrolment

01/04/2011

Date of final enrolment

01/10/2012

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Evelina Children's Hospital

London

United Kingdom

SE1 7EH

Sponsor information

Organisation

Guy's and St.Thomas' NHS Foundation Trust (UK)

ROR

<https://ror.org/00j161312>

Funder(s)

Funder type

Charity

Funder Name

Research Autism (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				

Results article		01/08/2014		Yes	No
Study website	Study website	11/11/2025	11/11/2025	No	Yes