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Evaluation of a patient education package in childhood enuresis.

Submission date	Recruitment status	[ ] Prospectively registered
23/01/2004	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
23/01/2004	Completed	[X] Results
Last Edited	**Condition category**	[ ] Individual participant data
03/12/2008	Mental and Behavioural Disorders	

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
RBF 96X36

Study information

Scientific Title

Study objectives

To evaluate a multimedia education package, developed to educate children about their enuresis, for its effect on: clinical outcome; child and parent satisfaction; and children's levels of self-esteem.

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

Study type(s)
Not Specified

Health condition(s) or problem(s) studied
Mental and behavioural disorders: Other mental disorder; Urological and genital diseases: Other urological and genital disease

Interventions
1. Five clinics will continue to provide standard information to the children and parents
2. Five clinics will utilise the multimedia education package
3. Five clinics will provide the same information as the multi-media package but in printed leaflet form

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
Measures that are routinely monitored in the treatment of enuresis: number of children who become 'dry'; length of treatment time to attain 'dry' status; and relapse rates at the standard 6 month appointment.

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

Completion date
30/06/1999

Eligibility

Key inclusion criteria

The project would follow children through the standardised treatment protocol for enuresis in Leicester Community Trust (28 clinics in all). Five clinics would provide standard information and care, five would use the multimedia package, and five printed information with similar content to that of the package. 300 children, divided into three groups, 100 children for each experimental condition.

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/07/1997

Date of final enrolment
30/06/1999

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Department of Paediatrics
Nottingham
United Kingdom
NG1 1PB

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