

Evaluation of a patient education package in childhood enuresis.

Submission date 23/01/2004	Recruitment status No longer recruiting	☐ Prospectively registered
		☐ Protocol
Registration date 23/01/2004	Overall study status Completed	☐ Statistical analysis plan
		☒ Results
Last Edited 03/12/2008	Condition category Mental and Behavioural Disorders	☐ 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
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

Primary study design

Interventional

Study design

Randomised controlled trial

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