

Evaluation of a patient education package in childhood enuresis.

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| Submission date 23/01/2004 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered |
| | | <input type="checkbox"/> Protocol |
| Registration date 23/01/2004 | Overall study status Completed | <input type="checkbox"/> Statistical analysis plan |
| | | <input checked="" type="checkbox"/> Results |
| Last Edited 03/12/2008 | Condition category Mental and Behavioural Disorders | <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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

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 measure

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.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/07/1997

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

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

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

England

United Kingdom

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

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details

The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

NHS Executive Trent (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

Study outputs

| Output type | Details | Date created | Date added | Peer reviewed? | Patient-facing? |
|---------------------------------|---------|--------------|------------|----------------|-----------------|
| Results article | results | 01/03/2003 | | Yes | No |