The relative efficacy of two brief treatments for sleep problems in young learning disabled (mentally retarded) children: a randomised controlled trial

P Montgomery, G Stores, L Wiggs

Background: Settling and night waking problems are particularly prevalent, persistent, and generally considered difficult to treat in children with a learning disability, although intervention trials are few. Scarce resources, however, limit access to proven behavioural treatments.

Aims: To investigate the efficacy of a media based brief behavioural treatment of sleep problems in such children by comparing (1) face-to-face delivered treatment versus control and (2) booklet delivered treatment versus controls.

Methods: The parents of 66 severely learning disabled children aged 2–8 years with settling and/or night waking problems took part in a randomised controlled trial with a wait-list control group. Behavioural treatments were presented either conventionally face-to-face or by means of a 14 page easy to read illustrated booklet. A composite sleep disturbance score was derived from sleep diaries kept by parents.

Results: Both forms of treatment were almost equally effective compared with controls. Two thirds of children who were taking over 30 minutes to settle five or more times per week and waking at night for over 30 minutes four or more times per week improved on average to having such settling or night waking problems for only a few minutes or only once or twice per week (H = 34.174, df = 2, p < 0.001). These improvements were maintained after six months.

Conclusions: Booklet delivered behavioural treatments for sleep problems were as effective as face-to-face treatment for most children in this population.

METHODS

Participants

The study involved children aged 2–8 years with any form of severe learning disability, confirmed by the general practitioner (GP). Severe sleep problems were defined according to standardised criteria as follows: (1) night waking occurring three or more times a week for more than a few minutes and the child disturbing the parents or going into their room or bed; and/or (2) settling problems occurring three or more times a week with the child taking more than one hour to settle and disturbing the parents during this time. These problems needed to have been present for at least three months and not be explicable in terms of a physical problem such as pain.

Intervention

This consisted of the same instructions to parents in one of two randomly allocated formats:
• Using a brief booklet
• Conventionally, face-to-face.

In both cases the approach was entirely behaviourally and focused on the child's sleep problems and the parents' response to them, covering the following topics:

• **Normal sleep:** setting realistic expectations, indicating the advantages to the whole family when children sleep well.
• **Introduction to behavioural techniques in general:** how behaviours can be triggered by events that precede them, and encouraged (or discouraged) by providing appropriate reinforcement; management strategies such as ignoring, consistency, and reward systems.
• **Monitoring behaviour:** using a sleep diary to record settling, wakeings, daytime naps, sleepiness, etc; and using this information to devise and monitor treatment plans.
• **Good sleep habits:** for example, sleep as a learned behaviour, appropriate surroundings, removal of daytime naps, importance of clear routines, putting children down to sleep while awake but drowsy, removal of bottles, how to deal with possible physical causes of sleep disturbance such as wet nappies or being too hot/cold.
• **Specific techniques for changing undesirable behaviour:**
  - **Settling and nightwaking:** (i) ignoring the child; (ii) checking briefly on the child at increasingly extending intervals with minimal attention; (iii) gradually decreasing the physical contact between child and parent during the episode.
  - **Sleeping in the parents' bed:** (i) raising awareness of the problem; (ii) returning the child to bed as needed using the settling techniques above.
  - **Rewards** for encouraging desirable behaviour.

The intervention was delivered by one of the researchers (PM) either conventionally face-to-face, or briefly via booklet at the second meeting (see Procedure). With the face-to-face approach, time spent with the parent was about 90 minutes and followed precisely the same material as in the booklet. This information was based, with permission, on a booklet written by the Child Psychology Department, Dudley Road Hospital, West Birmingham and expanded by the authors to address the needs of learning disabled children. It consisted of 14 pages of text with cartoons to make the booklet more appealing and illustrate some of the techniques. Written information for parents available in most bookshops and primary care centres contains similar information. There is little variation in the content of these kinds of materials, although none has previously been systematically tested and validated.

The booklet was evaluated using the Flesch Readability Test, a standard method of assessing the level of secondary education required to understand written material. The Flesch index was 74.3 and the Flesch–Kincaid Grade level 7.3, indicating that it is readable by someone educated up to age 13. For comparison, it is simpler than three quarters of UK newspapers.

### Assessments

**Sleep**

A brief sleep problem screening questionnaire, previously shown to be valid and reliable, was used to identify children whose parents reported a severe sleep problem as set out in Table 1, and to establish whether this was likely to respond to a behavioural intervention or whether it might reflect a physical problem such as pain. A sleep history was taken in a semi-structured interview. Parents were given a sleep diary to complete for the two weeks prior to each assessment, primarily to record the times of sleep onset, offset, and night waking, but also to describe how they dealt with the problems. From the diary a Composite Sleep Disturbance Score was calculated by summing the score on each problem.

The minimum entry score was 4, representing, for example, a child with settling problems lasting more than 30 minutes at least five times weekly. The maximum is 8, indicating a child who also wakes in the night for at least 30 minutes more than three nights each week. Since parental report is the main way in which children sleep problems come to light in primary care, the Composite Sleep Disturbance Score was chosen as the primary outcome measure.

**Parental evaluation of response**

Parents were asked what minimum improvement in their child's sleep would be needed for them to consider the intervention worthwhile, following Jones and Verduyn (1983). Fifty five parents (83%) said that if the problem reduced by half they would think the intervention worthwhile, so a reduction in the Composite Sleep Disturbance Score of 50% or more was taken as the secondary outcome measure.

**Evaluation of the booklet**

Participants in the booklet groups were given a brief questionnaire to evaluate it in terms of relevance, ease with which the booklet was understood, and usefulness. Each item could be scored from 0 to 4 (“not at all…” “quite…” “not very…” “not at all …”). The range of scores was, therefore, 0 (worst) to 12 (best).

**Design**

This was a randomised controlled trial with two active treatment groups, conventional and booklet delivery of behavioural advice, as well as a crossover control group. Power calculations based on previous work with conventionally delivered behavioural treatment showing a reduction in the treatment group's Composite Sleep Disturbance Scores from 6.73 to 2.96 (SD 2.24) and in controls from 7.23 to 6.29 (SD 2.70), indicated that group sizes of 20 would give 95% power at the 0.05 level of significance. As the relative difference of effect between the two delivery methods was unknown, a pragmatic decision was taken to recruit as many participants as possible in the time available.

**Procedure**

Screening questionnaires and letters requesting details of their child’s sleep (without the offer of treatment) were sent to the parents of all 268 children attending the special schools or receiving pre-school teacher counsellor services in

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Scoring of the composite sleep disturbance score</th>
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</thead>
<tbody>
<tr>
<td><strong>Problem</strong></td>
<td><strong>Score</strong></td>
</tr>
<tr>
<td>Settling frequency</td>
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</tr>
<tr>
<td></td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>2</td>
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<tr>
<td>Settling duration</td>
<td>0</td>
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<tr>
<td></td>
<td>1</td>
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<td></td>
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<td>Night waking frequency</td>
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<td>1</td>
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<tr>
<td></td>
<td>2</td>
</tr>
<tr>
<td>Night waking duration</td>
<td>0</td>
</tr>
<tr>
<td></td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>2</td>
</tr>
</tbody>
</table>
Oxfordshire, Berkshire, and Buckinghamshire. Parents of eligible children were sent a letter inviting them to participate in the treatment trial and informed consent was obtained at this point from parents and from the child’s GP. Those agreeing were given a baseline assessment of the sleep questionnaire and subjective measurement of their child’s sleep using a sleep diary kept for two weeks before each assessment. A detailed sleep history was taken to assess other clinical features of the children’s sleep.

Participants were then randomly allocated to one of the experimental groups.

- **Brief treatment (booklet):** giving parents a simple booklet, describing appropriate techniques for dealing with various sleep problems.
- **Conventional treatment:** seeing parents in their homes and presenting face-to-face the behavioural strategies used in the booklet to address their child’s sleep problem.
- **Crossover control group:** no intervention for six weeks, at which point they were re-randomised into an active treatment group.

The family received a second visit (on average five days later) and delivery of the appropriate intervention. No contact was then made until six weeks later, when a sleep diary was sent and all measures were repeated in participants’ homes. Total contact time with each family was approximately 180 minutes for face-to-face conventional treatment (of which approximately 90 minutes was spent on the intervention) and 90 minutes with the booklet group for assessments.

All participants received a postal follow up in which parents completed a sleep diary for one week, approximately six months post-intervention. From this the Composite Sleep Disturbance Score was again calculated to assess whether any treatment effects had been maintained. Families failing to return this questionnaire were reminded once by mail and then telephoned for collection of these data.

**Allocation**

Seventy five opaque envelopes were produced for the initial randomisation and lodged with an independent staff member. Each contained a slip of paper with the word “conventional”, “booklet”, or “control” (25 each). The randomisation was performed by this staff member selecting an envelope for each participant immediately after the initial assessment meeting with parents. For the re-randomisation of the control crossover group this process was repeated with a second batch of 26 envelopes, half each with the word “conventional” or “booklet”. The researcher conducting the study was therefore blind to the nature of the treatment allocated until after the post-treatment assessment. Following that point both participant and researcher were aware of the treatment group to which they had been randomised.

**Analyses**

Distributions of the data were not normal, even after transformation, so all analyses presented here are non-parametric. The main comparisons were across three groups (conventional face-to-face treatment, booklet treatment and control). Post hoc multiple comparisons were performed to identify which groups differed.

Care was taken to ensure that treatment content for the conventional group was consistent with the booklet. Taped sessions were compared with the booklet and a selection of these were rated by an experienced clinical psychologist who helped to ensure consistency between the two delivery methods. A selection of Composite Sleep Disturbance Scores was randomly cross checked for consistency by an independent rater who found agreement levels greater than 95%.

**RESULTS**

Of 268 parents approached, 184 (69%) replied to the invitation letter and screening questionnaire. Of these, 102 (55%) were invited to join the study as their children met criteria; 76 responded and actually joined the study, and 66 (42 boys and 24 girls), completed it as shown in fig 1. The groups did not differ significantly on any baseline sociodemographic or medical measures (see table 2). Children were aged between 27 and 101 months. Four children were said to have epilepsy. Thirty of the parents left school before the age of 18, and 36 left aged 19 or older.

**Composite sleep disturbance scores**

**Baseline**

Table 3 shows the Composite Sleep Disturbance Scores by group at baseline. Statistically there was no significant difference between them (Kruskal Wallis test statistic \( H = 0.362, \) df = 2, \( p = 0.834 \)), although sleep problems appeared slightly worse in the group receiving conventional treatment, which may be clinically important. A difference of 1 point on this scale would suggest night waking for up to 30 minutes rather than for a few minutes (see table 1).

**Post-treatment**

Significant group differences were found in Composite Sleep Disturbance Scores over time (\( H = 34.174, \) df = 2, \( p<0.001 \)), as shown in table 3, but the effects of active treatment were very similar, irrespective of delivery method. Post-hoc multiple comparisons using Bonferroni correction for type 1 errors confirmed that the significant differences were between the active treatments and controls. Power calculations indicated that it was not possible to detect a difference between the two active treatments, as a sample of 6325 would be required to achieve 95% power. A replication phase after the controls received treatment was not possible as there were no controls remaining with whom to compare them.

**Six month follow up**

One-week sleep diaries were obtained from 64 of the 66 participating families in order to assess the durability of any treatment effects six months after the children began active treatment. As shown in table 3, the data indicated that treatment effects had been maintained.

**Controls**

No significant improvements in sleep scores were observed during the control phase. However, when treatment was delivered, by whichever method, they made similar improvements to those seen in groups who received treatment immediately.

**Response to treatment**

In addition to the primary outcome measure, the parental evaluation of response as defined by a 50% symptom reduction was also considered. The conventionally treated group included 13 “responders” versus 5 “non-responders”. Comparable figures were 15 versus 7 for the booklet group, and 0 versus 24 for controls. These data support the hypothesis that booklet delivered and therapist delivered treatments were equally effective when compared with controls (\( H = 36.975, \) df = 2, \( p<0.001 \)). In the control-crossover groups similar results were obtained when they received active treatment, with 9 responders versus 3 non-responders in the conventional group and 8 versus 4 in the booklet group.
Evaluation of the booklet
All participants who received the booklet (whether immediately or after a period in the control group) were asked to evaluate it briefly in terms of relevance, ease with which it was understood, and usefulness. Twenty-three replied (68%). The mean score was 10.17 (SD 1.87) (max = 12), indicating that parents found the booklet appropriate and helpful. In 13 cases both parents reported that they used it, in nine it was the mother only, and in one case the father only.

Sociodemographic factors
No social indicators such as family structure or social class were found to relate to outcome with respect to the Composite Sleep Disturbance Score change (for example, number of parents, Fisher exact test statistic ($S$) = 8.51, $p<0.8$; number of siblings, $S = 41.62$, $p<0.4$; social class, $S = 26.80$, $p<0.9$). However, to predict which families are likely to have success with the booklet would require a much larger sample size.19 There were no families for whom literacy was a problem, even though more than a third of the parents left school at age 16 or younger. The importance of ethnic group could not be considered as only eight of the 66 families described themselves as non-Caucasian. Small group sizes prevented statistical analysis of the importance of health, medication, and epilepsy status. However, these possible factors appeared to be similar across groups.

**DISCUSSION**
This study confirms the effectiveness of conventional behavioural treatment for sleep problems in children with learning disabilities (as reported by parents), and shows that brief delivery of this treatment using a booklet did not reduce...
its effect. Treatment benefits were maintained at six months. Previous research in both this kind of population and normal children has shown that the behavioural techniques used here can contribute to the successful management of serious sleep problems. The importance of this study, however, lies in its demonstration that this sort of treatment, which is often hard to access owing to resource constraints, may be presented briefly via a booklet with no loss of effectiveness.

The non-parametric tests used here are, arguably, less powerful than parametric tests, but a convincing treatment effect was still found. Using the parent generated index of satisfaction (sleep problems improving by 50% or more), 75% of the conventional group and 68% of the booklet group were satisfied. The minimal differences between the two delivery methods suggest that, clinically, the booklet has a good therapeutic effect and parents rated it highly. It is short, easy to read, well illustrated, and makes specific reference to the needs of learning disabled children, which is rare in the self-help field.

In addition to the interventions themselves, other factors could affect efficacy. Motivation is often thought to affect motivation in clinical practice.

Table 2 Baseline demographic data of the children and their families

<table>
<thead>
<tr>
<th>Conditions</th>
<th>Total sample</th>
<th>Therapist</th>
<th>Booklet</th>
<th>Controls</th>
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<tr>
<td>frequency</td>
<td>%</td>
<td>frequency</td>
<td>%</td>
<td>frequency</td>
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<td>Autism</td>
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<td>7.6</td>
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<tr>
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</tr>
<tr>
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<td>27.3</td>
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<td>20</td>
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<td>8</td>
<td>40</td>
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<td>Manag/technical</td>
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<td>24.2</td>
<td>4</td>
<td>20</td>
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<tr>
<td>Skilled manual</td>
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<td>13.6</td>
<td>3</td>
<td>15</td>
</tr>
<tr>
<td>Skilled non-manual</td>
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<td>12.1</td>
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<td>5</td>
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<td>Unskilled</td>
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<td>13.6</td>
<td>4</td>
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<td>11</td>
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<td>5</td>
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<tr>
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<tr>
<td>3</td>
<td>1</td>
<td>1.5</td>
<td>0</td>
<td>0</td>
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</table>

*Microcephaly, n = 2; hydrocephalus, n = 2; Angelman syndrome, n = 2; cerebral palsy, n = 2; Williams syndrome, n = 1; triple X syndrome, n = 1.

Table 3 Composite Sleep Disturbance Score (CSDS) over time by treatment group

<table>
<thead>
<tr>
<th>Group</th>
<th>n</th>
<th>Baseline Mean (SD)</th>
<th>Post-treatment Mean (SD)</th>
<th>Follow up (6 mth) Mean (SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conventional treatment</td>
<td>20</td>
<td>6.55 (1.31)</td>
<td>2.4 (1.93)</td>
<td>1.89 (2.02)</td>
</tr>
<tr>
<td>Brief treatment Controls</td>
<td>22</td>
<td>6.18 (1.46)</td>
<td>2.55 (2.76)</td>
<td>2.08 (2.89)</td>
</tr>
<tr>
<td>All participants</td>
<td>66</td>
<td>6.10 (0.43)</td>
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</table>
ACKNOWLEDGEMENTS
The authors would like to thank the children and families who participated in this research, the NHS Research and Development Board for funding it, Christina Crawford for the artwork in the booklet, and Judith Ponsford for her work on the randomisation.

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