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Specialist home-based nursing services for children with acute and chronic illnesses (Review)

Parab CS, Cooper C, Woolfenden S, Piper SM

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[Intervention Review]

Specialist home-based nursing services for children with acute and chronic illnesses

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ABSTRACT

Background

Specialist paediatric home-based nursing services have been proposed as a cost-effective means of reducing distress resulting from hospital admissions, while enhancing primary care and reducing length of hospital stay. This review is an update of our original review, which was published in 2006.

Objectives

To evaluate specialist home-based nursing services for children with acute and chronic illnesses.

Search methods

We searched the following databases in February 2012: the Cochrane Central Register of Controlled Trials (CENTRAL) in *The Cochrane Library* 2012 Issue 2, Ovid MEDLINE, EMBASE, PsycINFO, CINAHL and Sociological Abstracts. We also searched ClinicalTrials.gov and the WHO International Clinical Trials Registry Platform. No language restrictions were applied.

Selection criteria

Randomised controlled trials (RCTs) of children from birth to age 18 years with acute or chronic illnesses allocated to specialist homebased nursing services compared with conventional health care. Outcomes included utilisation of health care, physical and mental health, satisfaction, adverse health outcomes and costs.

Data collection and analysis

Two review authors extracted data from the studies independently and resolved any discrepancies by recourse to a third author. Metaanalysis was not appropriate because of the clinical diversity of the studies and the lack of common outcome measures.

Main results

We screened 4226 titles to yield seven RCTs with a total of 840 participants. Participants, interventions and outcomes were diverse. No significant differences were reported in health outcomes; two studies reported a reduction in the hospital stay with no difference in the hospital readmission rates. Three studies reported a reduction in parental anxiety and improvement in child behaviours was reported in three studies. Overall increased parental satisfaction was reported in three studies. Also, better parental coping and family functioning was reported in one study. By contrast, one study each reported no impact on parental burden of care or on functional status of children. Home



care was reported as more costly for service providers with substantial cost savings for the family in two studies, while one study revealed no significant cost benefits for the family.

Authors' conclusions

Current research does not provide supporting evidence for a reduction in access to hospital services or a reduction in hospital readmission rate for children with acute and chronic illnesses using specialist home-based nursing services; however, the only summary finding across a few studies was that there is a significant decrease in length of hospitalisation. The preliminary results show no adverse impact on physical health outcomes and a number of papers reported improved satisfaction with home-based care. Further trials are required, measuring health, satisfaction, service utilisation and long-term costs.

PLAIN LANGUAGE SUMMARY

Specialist home-based nursing services for children with acute and chronic illnesses

This review aimed to examine whether specialist paediatric home-based nursing services for children with acute and chronic illnesses reduce the number of hospital admissions and length of stay, enhance health care in the community and reduce stress for families at the time of their child's illness. It is an update of our original review published in 2006. We found seven relevant randomised controlled trials (RCTs) of total of 840 children aged from birth to 18 years with acute and/or chronic illnesses receiving either specialist home-based nursing services or conventional health care. The outcomes included use of health care services, physical and mental health, satisfaction, adverse health outcomes and costs. We decided not to combine the results of these RCTs because of the variety in types of services provided, types of participants and the outcome measures used. The results of individual RCTs show improved satisfaction with home-based care with no adverse impact on physical health outcomes for children. There is some evidence that specialist home-based nursing services reduce the length of hospital stay; however, there is no evidence that it leads to a reduction in use of hospital services. Further trials are required, measuring health, satisfaction, service use and long-term costs.



BACKGROUND

Description of the condition

The illness of a child leading to hospitalisation is a stressful event for the family (Kai 1996). It can be distressing for children, as in some emergency departments they may be separated from their families and the security of their own home (Farquahar 1990; Lenihan 1985). While not every child who presents to a hospital is admitted (Cooper 2000), many such children spend several hours in the emergency department (ED) while being observed or assessed (Browne 1996); in addition, children may wait while their carers are provided with education on how to manage the child's illness.

Children presenting at EDs frequently have a non-urgent problem (Bowling 1987; Halfon 1996; Jones 1999; Worsley 1985) and no usual care provider (Beebe 1993; Grover 1994). Many parents take their child to an ED seeking reassurance (Boyle 2000). These presentations, and subsequent admissions, may provide opportunities to give carers the confidence to support and care for their child, as well as to provide education about the disease process and its management. Admission of children is not always determined by the severity of their illness; it can also be influenced by previous medical consultations for the illness as well as parental expectations that the child will not be discharged from the ED (Cooper 2003).

In the current era, the necessity to explore alternative models of service delivery such as home-based nursing services has become more important, based on increasing survival of children with complex conditions, and also as it has become more feasible to provide high-technology interventions at home (Parker 2012). Thus, there is a shift towards parents being more responsible for complex care for children children with chronic illnesses in the home settings with educational as well as clinical management support.

Description of the intervention

The need to reduce hospital admission rates and shorten the length of hospital stay has prompted the development of specialist homebased nursing services for children with acute and chronic illnesses. The home-based nursing care substitutes for acute hospital review and/or admission by providing clinical review, support, education and management of the acutely or chronically unwell child in their own home. If the child requires it, they are provided with streamlined access to hospital services. It has been postulated that these services have the potential to reduce the number of hospital admissions (Meates 1997; Smith 1986), to reduce the length of stay (Meates 1997; Whiting 1997) and to facilitate early discharge by providing a continuum of care from the hospital into the home (Meates 1997). They also provide opportunities to enhance primary care in the community through liaison with general practitioners (GPs) and through links with other community-based health services (Fradd 1994).

Why it is important to do this review

The development of specialist home-based nursing services for children with acute and chronic illnesses has been determined by local circumstance and enthusiastic individuals rather than strategic planning based on evidence (While 2000), resulting in a diverse range of service models. Much of the literature comprises accounts of service development (Box 1993) or descriptive studies of parent or professional views (Madigan 1997; Sartain 2001). Literature related to the safety of models of care is limited (McConnochie 1998; Montalto 1998) and concerns are rapidly increasing about the impact of changes in delivery and financing mechanisms on quality of care (Kelly 1994). Empirical data are needed about the effects of specialist home-based nursing services for children with acute and chronic illnesses, to explore the strengths and weaknesses of different service models, as well as general efficacy.

The literature search for this review was completed in 2005 prior to its first publication (Cooper 2006). Subsequently, a systematic review was published in the UK about international evidence for the effectiveness and costs of paediatric home care. The Parker 2012 review was produced because there had not been a substantial growth in this kind of service despite advocacy by policy makers and professionals over the past few decades. We have conducted this update on a similar basis, to explore the available international evidence regarding specialist home-based nursing services for children and adolescents.

OBJECTIVES

To evaluate the impact of specialist home-based nursing services for children with acute and chronic illnesses.

METHODS

Criteria for considering studies for this review

Types of studies

Trials were eligible for inclusion in the review if:

- The assignment of study participants to the intervention or control group was random or quasi-random, for example, by alternate allocation;
- 2. The study intervention had paediatric nursing outreach services as its major focus;
- 3. There was at least one objective measure, for example, hospital admission data and/or a standardised measure such as a behaviour checklist, and that such measures were used for both intervention and control groups.

Types of participants

Children aged from birth to 18 years with acute and or chronic illnesses. In this review, terms such as 'chronic illness' and 'long-term illness' are used interchangeably.

Types of interventions

Specialist home-based nursing services provided to children with acute illnesses and/or chronic and complex conditions, compared with conventional health care (for example, hospital admission) as the control group. We excluded studies where nurses provided planned illness educational support independent of clinical review and management of children with acute and/or chronic diseases.

Types of outcome measures

- 1. Physical health of participants;
- 2. Mental health of participants;
- 3. Utilisation of emergency departments (EDs);
- 4. Hospital admissions;



- 5. Length of stay in hospital;
- 6. Parental, child and referrer satisfaction;
- 7. Quality of life of children and their carers;
- 8. Cost of treatment for the family and to the service providers.

The study outcomes were measured using the following.

- 1. Hospital data on service utilisation, admissions, length of stay.
- 2. Questionnaires, for example, quality of life questionnaires, health outcome rating scales.
- 3. Satisfaction surveys.

Search methods for identification of studies

Electronic searches

This update is based on database searches run in February 2012 (Appendix 1), which cover the period since the search date of the previous version of this review in August 2005. We used the most recent version of the Cochrane highly sensitive search strategy for identifying randomised trials in Ovid MEDLINE (Lefebvre 2008) and searched two additional sources (ClinicalTrials.gov and International Clinical Trials Registry Platform).

We searched the following databases with no language restrictions.

The Cochrane Central Register of Controlled Trials (CENTRAL), part of the *The Cochrane Library*, 2012 Issue 2, last searched 9 February 2012;

Ovid MEDLINE (1946 to January Week 4 2012), last searched 8 February 2012;

CINAHL (1937 to current), last searched 9 February 2012;

EMBASE (1980 to 2012 Week 05), last searched 9 February 2012

PsycINFO (1806 to Feb Week 1 2012), last searched 9 February 2012; Sociological Abstracts (Sociofile) (1952 to current), last searched 9 February 2012;

ClinicalTrials.gov, all available years searched 10 February 2012; International Clinical Trials Registry Platform, all available years searched 10 February 2012.

Searching other resources

The archives of abstracts from the Pediatric Academic Societies (incorporating the American Ambulatory Pediatric Association and the Society for Pediatric Research) Annual Scientific Meetings from 2000 to 2012 did not yield any unpublished data from randomised trials evaluating specialist home-based nursing services for children. We had no responses to emails to the Ambulatory Care Australia email list and the 'Hospital in the Home' electronic exchange.

Data collection and analysis

Selection of studies

Three review authors working independently (CC, DW, SW) screened titles and abstracts from the searches for the original review and two authors (CP, CC) screened results for this update. We resolved disagreements by consensus. We obtained potentially relevant articles for assessment of the full text and for data extraction.

Data extraction and management

We input data using Review Manager 5.1 (Cochrane Handbook). Data extraction forms were developed a priori and included information about study location, methods, participants, type of ambulatory paediatric intervention, and outcomes. Two review authors (CC, SW) independently performed data extraction for the initial phase of the review and two (CP, CC) for the next phase. We resolved disagreements by negotiation with a third review author (DW for the first phase and SW for the second phase).

Assessment of risk of bias in included studies

We assessed the risk of bias for the five previously included studies and the two newly added studies using the domain-based evaluation tool described in the Cochrane Handbook. Each study was assessed as being at low, high or unclear risk of bias for allocation concealment, blinding, incomplete outcome data and other sources of bias.

Measures of treatment effect

We report mean differences and 95% confidence intervals for continuous variables, where they were available. For binary outcomes, we report the risk ratio and 95% confidence interval. For meta-analysis, where possible in future updates, we will use mean differences for continuous variables, with standardised mean differences if different scales measure the same construct in different studies, and risk ratios for binary outcomes.

Dealing with missing data

We analysed data where possible on an intention-to-treat basis, i.e. with all participants randomised in their original groups. We contacted trial authors to obtain missing or additional information.

Assessment of reporting biases

The tool described in the Cochrane Handbook was used to assess the risk of bias in included studies. Two authors (CP, CC) independently carried out this assessment and there was an agreement among both for conclusions.

Data synthesis

Due to the clinical diversity of the participants' illnesses and the services offered, as well as a lack of common outcome measures, we decided against meta-analysis and statistical assessment of heterogeneity (the I^2 statistic; Higgins 2002), and we therefore provide a narrative summary. In future updates, we intend to include any available data in meta-analyses if it proves feasible.

Subgroup analysis and investigation of heterogeneity

We stratified the results to determine any differences in treatment effects according to age and type of intervention:

- 1. Clinically different interventions, for example, hospital-based versus community-based services;
- 2. Clinically relevant differences between groups of participants:
- children with acute illness, children with chronic illness, children with chronic haematological illnesses;
- children under 10 years, children between 10 and 18 years.



We present results according to the above subgroups. However, statistical subgroup analysis was not possible for the current version of this review.

Sensitivity analysis

Sensitivity analysis (to evaluate the impact of study quality on outcome) was not possible. If a meta-analysis can be undertaken in future updates, we will use sensitivity analyses to determine the impact of risk of bias on our findings.

RESULTS

Description of studies

Results of the search

The original review included five studies. The search was run again from August 2005 until February 2012. A total of 2570 new titles were screened and 152 duplicates were identified. Of the 2418 titles screened, 2347 were not relevant to the review. We read the remaining 71 articles to obtain further information and only two studies (Stevens 2006; Tie 2009) were suitable for inclusion, giving a total of seven studies in this update. See the study flow diagrams Figure 1 and Figure 2.





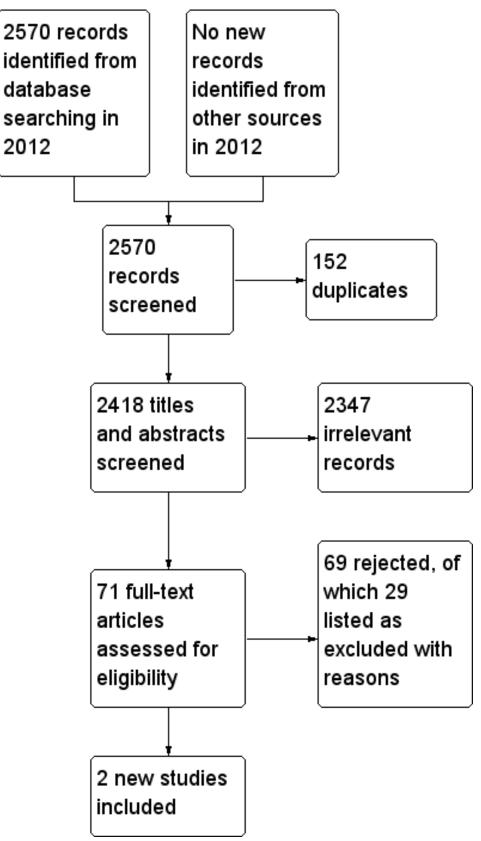
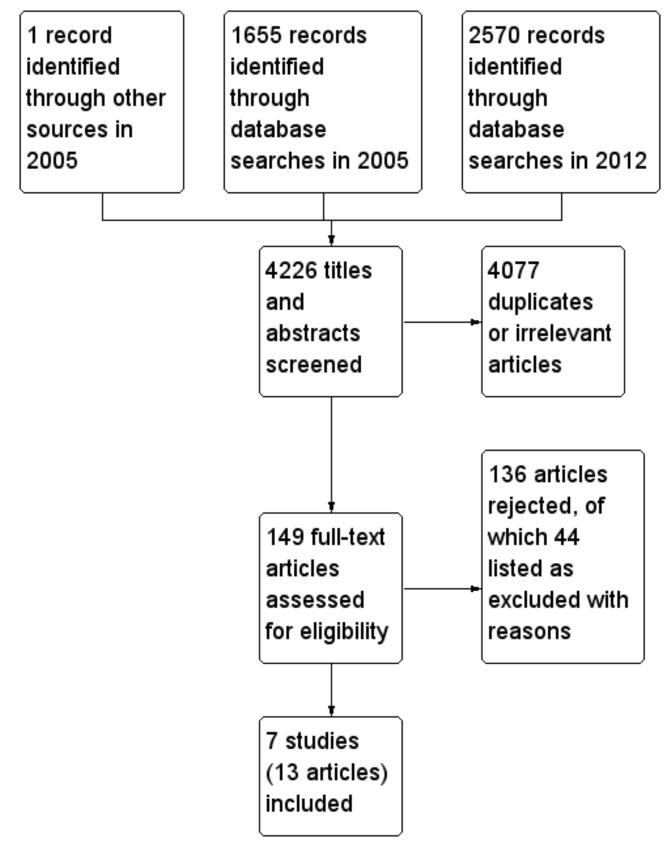




Figure 2. Study flow diagram incorporating all records for lifetime of review





Included studies

Please see Characteristics of included studies table for more detailed information.

The number of participants in the seven included studies ranged from 29 (Stevens 2006) to 399 (Sartain 2001). The total number of participants across all the included studies was 840. Participants in these studies ranged from three months of age through to 17 years of age. Specialist home-based nursing outreach services were provided as an intervention in each study. However, studies differed in respect of the qualifications of the nurses and their availability, including hours of service and number of visits. This is detailed in Table 1, 'Characteristics of Interventions'. No two studies examined the same intervention. In addition, treatments and outcomes varied considerably. Included studies broadly fell under the following categories.

Study design

Five trials used parallel study designs while two had a crossover study design. Participants were allocated to the intervention versus usual posthospitalisation care in two studies (Burke 1997; Stein 1984) and to traditional inpatient care in the other three (Dougherty 1998; Sartain 2001; Tie 2009). Strawczynski 1973 was a cross-over study with participants spending a year in each of the intervention and control (hospital care) groups. Another cross-over study was Stevens 2006, with participants receiving each of hospital (standard care) or home (intervention) chemotherapy alternately over a period of six months in succession.

Interventions for children with acute paediatric illnesses

There were only two studies that evaluated home-based nursing intervention for children with acute illnesses. Sartain 2001 focused on children with acute paediatric illness with breathing difficulty, diarrhoea and vomiting or fever. The intervention, home care, was defined as clinical nursing care in a patient's own home, as an alternative to hospital admission. Treatment in this study was between one and four home visits per day, with the number of days and visits determined by health professionals in conjunction with parents. This study examined readmission rates and length of stay data as outcomes. The second study (Tie 2009) compared home oxygen therapy with traditional inpatient hospital care for children with acute bronchiolitis. The children in the home care group continued with oxygen therapy at home following initial hospitalisation and were supported by the nurse home visiting programme accompanied by parental education. The outcomes assessed were readmission to hospital within seven days of discharge home and the total duration of hospitalisation.

Interventions for children with a range of chronic conditions

Stein 1984 examined home care provided to children with chronic conditions through traditional clinics, inpatient units or a home care office. The minimum package was an initial assessment,

one home visit and one monthly contact thereafter for at least six months. Involvement in home care was reassessed every six months by the treating staff. This study measured outcomes related to psychological adjustment and functional status.

Another study (Burke 1997) examined a community-based stresspoint intervention provided by nursing staff to children with chronic illness. The intervention began two weeks before a planned hospital admission and continued until two weeks postdischarge. This study collected data throughout the intervention period using various scales to assess outcomes including stress and behavioural responses.

Interventions for children with newly diagnosed insulindependent diabetes mellitus (IDDM)

Dougherty 1998 examined home-based care provided by nursing staff over an extended period to newly diagnosed diabetic children, as an alternative to inpatient care. Nursing staff initially provided twice-daily home visits in the first few days after diagnosis, with transition to diminishing visits or phone calls to the family over two weeks, but remained available to the family for telephone contact for the next 24 months. This study focused on outcomes of diabetic management including metabolic control and diabetesrelated adverse events.

Interventions for children with chronic haematological illnesses

Strawczynski 1973 defined home care as the availability of access to a 24-hour nursing service to treat acute bleeds at home or at school for children with haemophilia, as soon as possible after they occurred. It did not report on the number of days or visits for individual children. This study reported the number of bleeds and reduction in procrastination time for haemophilia as its outcomes.

Another study (Stevens 2006) compared a hospital-based and a home-based programme for children with acute lymphoblastic leukaemia (ALL). Some of the chemotherapy treatment was delivered by the hospital pharmacy to children's homes and was then administered by a trained nurse from a community health service agency. The option of contacting a hospital oncology nurse was available to parents as required. The study outcomes were child's quality of life, the burden on parental caregivers, adverse effects and costs.

Excluded studies

Please see Characteristics of excluded studies table for details of the 41 excluded studies.

Risk of bias in included studies

The risk of bias for seven included studies was assessed using the Cochrane domain-based evaluation tool (Cochrane Handbook: Chapter 8) and is summarised below in Figure 3 and Figure 4.



Figure 3. Risk of bias summary: review authors' judgements about each risk of bias item for each included study

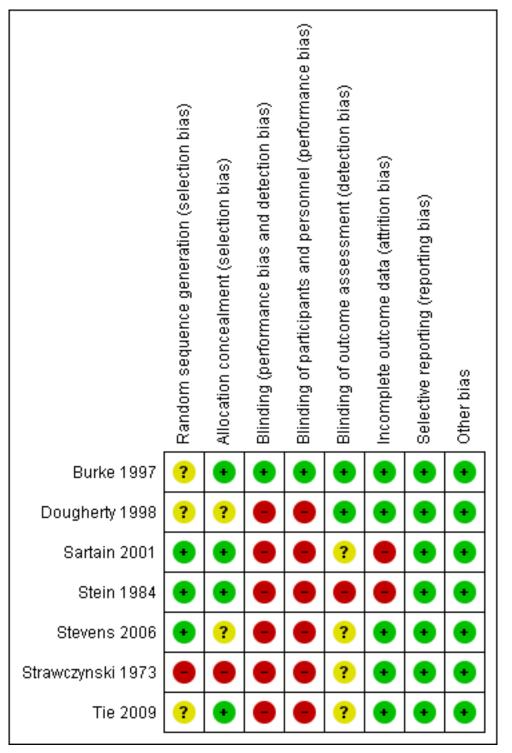
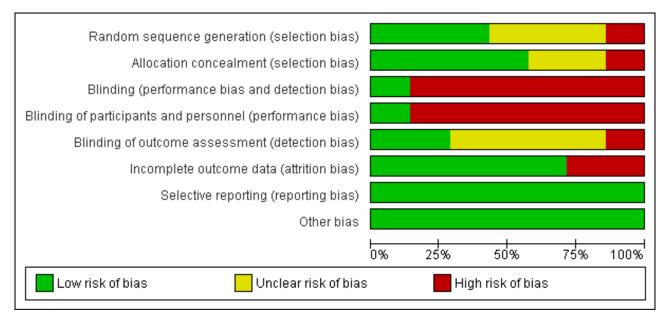


Figure 4. Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies



Allocation

While all seven studies stated that participants were randomised, only three gave information about the methods of randomisation or allocation concealment, and details were not reported for two studies (Dougherty 1998; Tie 2009). Opaque sealed numbered envelopes were used in one study (Sartain 2001) and a dice roll with opaque sealed envelopes in another (Stein 1984). A table of random numbers was used in a third study (Stevens 2006), but with no mention of the allocation concealment. Another study (Burke 1997) used randomisation by a secretary blinded to the study details but the method of allocation concealment was not clarified. Quasirandomisation was used in one study with alternate allocation (Strawczynski 1973).

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Where necessary, we contacted study authors for clarification or missing information, but received no replies.

Blinding

Outcome assessors were reported to be blind to the intervention groups in two of the included studies. Burke 1997 used study nurses to administer the pretest to participants. Research assistants blinded to study group membership administered post-test questionnaires. Study nurses did not have contact with hospital staff during the study. In Dougherty 1998, the principal investigators were not blinded to the intervention; however, results for the intervention and control groups were not disclosed before the end of the trial. Stein 1984 made no attempt to blind interviewers to the intervention status, but there was no direct contact between the clinicians and interviewers. Strawczynski 1973 was not able to blind clinical staff to intervention status or outcome measures. Sartain 2001 had the same researcher conduct all interviews in the participant's home within two weeks of discharge. It was not stated that the interviewer was blind to group allocation; however, the allocation is likely to have been disclosed during the interview. Stevens 2006 and Tie 2009 did not mention details of blinding, but the study methodology implies that blinding of neither participants nor clinicians was possible.

Incomplete outcome data

The number of participants lost to follow-up varied. Three studies (Dougherty 1998; Sartain 2001; Tie 2009) reported no withdrawal of participants. Burke 1997 recruited 50 children and two were lost to extended follow-up (4% loss). Loss to follow-up was 15% for Stein 1984 at six months and 17% at one year. Strawczynski 1973 enrolled 40 patients with four (10%) lost to follow-up. Of these, one died of an intracranial haemorrhage; one developed a circulating anticoagulant, and two moved away from the study city. Stevens 2006 randomised 29 children and two from each group withdrew at the start of the study due to relapse. Throughout the study, at four data collection stages, data were not available for analysis on one or two participants from each group. Only one study (Sartain 2001) conducted an intention-to-treat analysis.

Selective reporting

All studies appear to have reported on all measured outcomes.

Other potential sources of bias

There were no other potential sources of bias in the included studies.

Effects of interventions

Meta-analysis was not appropriate in this review because of the range of outcome measures being reported. In addition, interventions were diverse and complex involving multidisciplinary teams, different hours of service and variable inclusion of educational components (see Table 1). Participants were also diverse: some had chronic illnesses such as diabetes, haemophilia, acute lymphoblastic leukaemia (ALL), and others had an acute presentation. Results are therefore presented as a written summary of individual studies only, without data synthesis.

Physical health of participants (including harmful effects)

Stein 1984 reported presentations for concurrent illness during the period of the study and found no significant difference in general health between the intervention and control groups of children with a range of chronic conditions.

Burke 1997 identified improved parental ability to cope with illness in a chronically ill child three months postintervention (P < 0.001).

Dougherty 1998 reported on metabolic control and diabetesrelated adverse events in children with newly diagnosed insulindependent diabetes mellitus (IDDM). At 24 months, the home care group had lower mean HbA1c levels (biological measure of diabetic control), 6.1% versus 6.8% (P < 0.02). There was no difference between the two groups for adverse diabetes-related clinical events.

Strawczynski 1973 reported on the number of bleeds and procrastination time in children with haemophilia presenting with a bleed. Overall, significantly more bleeding episodes were reported in the home care programme, with the majority of these being mild to moderate bleeds. Severe bleeds were significantly more likely in the hospital programme (P value not reported). Procrastination time was defined as the time elapsed between the bleed occurring and reporting for treatment. Reported procrastination time decreased in the intervention group (mean 17 hours, compared to 29 hours in the control group; P value and significance not reported).

Stevens 2006 used Pediatric Oncology Quality of Life Scale (POQOLS) to assess quality of life, and reported no significant long-term difference between the hospital and home treatment groups in values for sensitivity to restrictions in physical functioning and the ability to maintain a normal physical routine.

Mental health of participants

Stein 1984 reported on psychological adjustment for children with a range of chronic conditions. For children five years and older, there was a significant improvement in the child's psychological adjustment at six months postintervention (P < 0.05), with some diminution by 12 months postintervention (P < 0.10). Mothers in the home care group showed improvement in mental health symptoms compared to those in the control group (P = 0.07).

Burke 1997 reported on the behaviour and anxiety of children with a range of chronic conditions. There was no significant difference between the two groups for child behaviour. However, at three months postintervention, intervention group parents were less anxious, with this difference being statistically significant. The intervention group had significantly less discrepancy between desired and actual family functioning at the end of three months of intervention.

Stevens 2006 used Child Behavior Checklist scores to identify behavioural problems and social competence among the study participants and reported no significant difference between intervention and control groups at any of the follow-up periods.

The utilisation of emergency departments (EDs)

No studies offered data for this outcome.

Hospital admissions

Sartain 2001 reported that there were no significant differences between home care and hospital care readmissions over 90 days, at 21 (10.0%) and 15 (7.9%) respectively (P = 0.49). Of those in home care, 14 were readmitted once (7%), two were readmitted twice (1%) and one child was readmitted three times (1%). For hospital care, 13 (7%) were readmitted once and one (< 1%) twice. No child was readmitted more than twice. No other studies reported on this outcome.

Tie 2009 reported that hospital readmission rates were the same for children with acute bronchiolitis within seven days of discharge for traditional hospital care versus home care. One child from each group of 22 participants required readmission.

Length of stay in hospital

Four studies reported on length of service provision. This was reported in several ways, including bed days, inpatient stay and use of hospital-based services. Sartain 2001 reported that children with acute illnesses who were in the control group, i.e. usual hospital treatment, were more likely to have a longer stay than those in the home care group, with mean bed stay of 2.37 versus 1.37 days (P < 0.0001).

Dougherty 1998 reported that there was no difference between both groups in the mean days spent in the intensive care unit at diagnosis. However, mean hospital inpatient stay was lower with home-based care (2.2 +/- 1.6 nights, 70 total) compared to hospital-based care (4.7 +/- 1.6 nights, 147 total). The use of services while hospitalised was lower for the home-based group, logistically because of shorter initial hospital stay. However, they did report that children receiving home care used more diabetes nursing hours during the 24-month period, 58.9 hours per child compared with 17.3 hours for standard care (P value and significance not stated), which implies that the intervention required more intensive nursing support.

Strawczynski 1973 reported the total number of treatment days for children with haemophilia in the home care group was 2030 compared with 1644 for the hospital-based group. However, only 241 from the home care group needed hospitalisation during the treatment period (P value and significance not reported).

Tie 2009 reported that children in the 'hospital in the home' treatment group spent significantly less time in a hospital bed (55.2 hours) than those in the hospital group (96.9 hours) (P = 0.001).

Parental, child and referrer satisfaction

Sartain 2001 collected parent-reported data for children with acute paediatric illnesses from a subset of the intervention group, which found that 36 of 40 (90%) parents and 7 of 11 (66%) children would prefer to access a 'Hospital at Home' service. There was a perception that children receiving the 'Hospital at Home' service recovered more quickly in their own environment and that there was less social disruption and financial burden for the family.

Stein 1984 identified overall greater satisfaction from respondents, with medical care provided in the home care group (P < 0.05) for children with a range of chronic conditions.

Quality of life of children and their carers

At three months post-intervention, Burke 1997 reported that for children with chronic illnesses, the home care group had greater satisfaction in family functioning (P < 0.001); greater parental ability to cope (P < 0.001); greater family ability to cope (P < 0.001); a greater ability in personal and social care coping (P < 0.01); and a greater perception of helpfulness from healthcare providers and institutional sources (P < 0.001).

Stein 1984 did not observe a significant difference for the impact of chronic illness on families between those receiving home care and those in the control group (P = 0.14).

Strawczynski 1973 reported significantly better school attendance (P value not stated) in the home care group, with an average of 2.5 school days missed compared to six days in the hospital programme for children with haemophilia.

Stevens 2006 reported no evidence of an effect of the location (i.e. home or hospital) where chemotherapy was administered in relation to parental burden of care using Caregiving Burden Scale (CBS) at different stages during the study period. Another important finding was that the age of the child was a significant predictor, with inverse effect being observed on quality of life, caregiver burden and adverse events.

Costs

Sartain 2001 assessed costs for children with acute paediatric illnesses in a separate paper (Bagust 2002). A number of parents thought that hospital care was more costly for them, with 20% of parents in the home care programme commenting on savings resulting from remaining at home and 30% of parents in the hospital care group commenting on the financial cost of the child staying in hospital.

Dougherty 1998 found that for children with newly diagnosed IDDM, there were social cost savings. This means that there were savings across hospital, physicians, tax payers and families. There were also parental cost savings (difference between out-of-pocket expenses and the value of their time diverted to care for their child from other activities) of CAD 188 for each child on the home care programme (P < 0.001) and an increased cost to the hospital of CAD 87 per child on the home care programme (P < 0.001).

Strawczynski 1973 reported total costs for providing both the home care and the hospital programme but did not give separate information on costs for the intervention and control groups.

Stevens 2006 did not find any significant difference between costs for the family (societal perspective) associated with home chemotherapy and standard care (P = 0.79).

DISCUSSION

Summary of main results

Two studies of children with acute paediatric illnesses (Sartain 2001; Tie 2009) reported a statistically significant reduction in the duration of hospital stay, with no significant difference in readmission rates

A variety of assessment tools were used in the studies included in this review to measure the functional impact or quality of life.

Burke 1997 found that home-care parents postintervention had better coping skills, family functioning and a reduction in anxiety than those in the traditional care group. No differences in child behaviour were reported in either group. A similar finding of no significant differences in child behaviour was reported in two other studies (Dougherty 1998; Stevens 2006) which used Child Behaviour Checklist (CBCL) for the assessment. A reduction in parental anxiety was noted in two other studies, with Burke 1997 and Stein 1984 reporting an improvement in mental health symptoms for mothers in the study group. Another study (Strawczynski 1973) reported better school attendance in the home-care children. Stein 1984 did not detect any significant impact of the intervention on the functional status of the participating children; Stevens 2006 reported that neither home or hospital location for treatment affected the parental burden of care. Overall, parental satisfaction with the home-based nursing care was reported in three studies (Sartain 2001; Stein 1984; Strawczynski 1973) which included semistructured parental interviews.

Cost effectiveness was not comprehensively addressed by any of the included studies. Three studies presented limited cost data. Dougherty 1998 suggested that the financial cost to the hospital of providing the home-based care programme may cost more than the hospital-based care, but suggested that there are substantial cost savings for the family from the home-care programme. This view was supported by Sartain 2001, who presented qualitative data on cost savings for families in the home-care programme versus increased expenditure for families in the hospital-care group. Interestingly, Stevens 2006 did not find any difference in family costs from a societal perspective in either home- or hospital-based treatment groups.

Overall completeness and applicability of evidence

There was wide clinical diversity between studies due to differences in participants, disease or condition requiring care, interventions and outcomes measured. There was also variation in the statistical methods used by studies. For these reasons, we did not conduct meta-analyses, which should be taken into consideration when drawing conclusions from this review. Few benefits of homebased nursing care were evident in the included studies. However, current studies (Dougherty 1998; Sartain 2001; Stevens 2006) do not take into account the original set-up costs of traditional inpatient services, or the long-term cost savings for children and families achieved by a less disruptive model of care. Further research is required that examines the long-term costs to both service providers and families before an economic rationalisation of home care can be attempted.

Quality of the evidence

This systematic review included seven randomised controlled trials with 840 participants ranging from three months to 17 years. Only one study (Burke 1997) blinded participants and personnel while only two blinded the process of outcome assessment (Burke 1997; Dougherty 1998). The risks of performance and detection bias should be considered when interpreting results of the relevant studies. Also, some studies have a limited sample size which impacts on the statistical power of the study.



Potential biases in the review process

The authors followed the process of systematic review as detailed in the Cochrane Handbook and have tried to minimise the likelihood of potential biases in the review process.

Agreements and disagreements with other studies or reviews

The findings of reduction of hospital stay and hospital readmission rates with home-based nursing care are consistent with some findings from the previously published literature (Meates 1997; Smith 1986; Whiting 1997). A recent systematic review (Parker 2012) on the paediatric home care services reported some evidence of reduced burden and costs for families, particularly in relation to acute admission avoidance. This is consistent with some of the cost-effectiveness findings in the studies included in this review. A mixed-methods study based in the UK (Spiers 2012) enumerates the diversity and complexity as well as variable clinical coverage of the service delivery models which are categorised as 'care closer to home'. This is consistent with the diverse clinical settings of the included studies in this review.

AUTHORS' CONCLUSIONS

Implications for practice

This systematic review has demonstrated a limited evidence base for home care programmes, with few randomised controlled trials undertaken, clinical and methodological heterogeneity between trials that have been completed. At present, there is insufficient evidence to support the effectiveness of specialist paediatric nurse home visiting for acute and chronic illnesses in reducing hospital admissions and Emergency Department utilisation. However, there is suggestive evidence that home care programmes may lead to greater parent satisfaction, improved quality of life and a reduction in the length of hospital stay. The cost effectiveness of these programmes is still to be determined.

Implications for research

Additional randomised controlled trials are required to examine home care programmes for children with acute and chronic

illnesses across the full age range of birth to 18 years. Adequate sample sizes must be used to generate the statistical power to detect differences between the intervention and control groups. Clinically relevant outcomes need to be defined for utilisation of Emergency Departments and length of hospital stay; physical health determinants; mental health determinants; and satisfaction measures. It would be useful if there were a standardised and consistent method of measuring these clinically relevant outcomes across studies, in particular, a definition of readmission to hospital and a standardised questionnaire for measuring parental satisfaction. While data for disease-specific diagnostic groupings would be the ideal, in reality ambulatory care programmes are usually set up in local populations to address a range of acute and chronic conditions. These include, but are not limited to, ambulatory sensitive conditions such as asthma and gastroenteritis.

Currently there are no data to support comprehensive costings for home care programmes from the perspective of a health organisation and the family. Complete evaluation of home care programmes requires estimation of both direct costs to healthcare providers and indirect costs to the child and their family, as well as to society. While healthcare provider costs may be readily assessable, the indirect costs of time off work, loss of earnings and childcare expenses must also be considered. The recently published systematic review (Parker 2012) on this topic refers to the need for additional high-quality research and collection of goodquality service level data with details about effects on the health economy. It also mentions that the evidence for paediatric homebased services continues to grow slowly.

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* Indicates the major publication for the study



CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Burke 1997

| Methods | Randomised controlled trial Sample size calculation |
|---------------|--|
| Participants | 50 children with chronic illness/physical disability for at least 8 months Age 1 to 17 (mean 6.2 years, SD 4.7) 26 boys (52%), 24 girls (48%) Follow up 96% at 3 months |
| Interventions | Stress point intervention vs usual care |
| Outcomes | Scales of Independent Behaviour Vernon Post-hospital Behaviour Questionnaire State-trait Anxiety Inventory Feetham Family Functioning Survey Coping Health Inventory for Parents |
| | 2 weeks and 3 months post intervention |

Notes

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence genera- tion (selection bias) | Unclear risk | Comment: Method of random sequence generation not mentioned. |
| Allocation concealment (selection bias) | Low risk | Quote; "A secretary blind to study details randomly assigned children to exper- imental or control groups within age and diagnosis strata." |
| Blinding (performance bias and detection bias) All outcomes | Low risk | Quote: " Participants, clinic staff, hospital staff and research assistants were blind to group membership". |
| Blinding of participants and personnel (perfor- mance bias) All outcomes | Low risk | As above |
| Blinding of outcome as- sessment (detection bias) All outcomes | Low risk | As above |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Comment: outcome data were reported for all participants. |
| Selective reporting (re- porting bias) | Low risk | Comment: outcome data were presented for all reported outcomes as mea- sured. |
| Other bias | Low risk | Comment: none were identified. |

Dougherty 1998

| Methods | Randomised controlled trial Intention-to-treat analysis not conducted | |
|---------------|---|--|
| Participants | 63 children with newly diagnosed insulin-dependent diabetes mellitus (IDDM) Intervention group: mean age 10.7 yrs, 13 boys (41%), 19 girls (59%) Control group: Usual care, mean 9.8 yrs, 15 boys (48%), 16 girls (52%). 100% follow-up | |
| Interventions | Home and outpatient care for education and treatment of diabetes vs inpatient care | |
| Outcomes | Mean hours per child of service provision Metabolic control Knowledge, adherence and psychosocial effects Parental time and costs Social cost effects Diabetes-related adverse effects Follow-up over 24 months | |

Notes

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence genera- tion (selection bias) | Unclear risk | Comment: Method of random sequence generation not mentioned. |
| Allocation concealment (selection bias) | Unclear risk | Comment: Allocation concealment details not mentioned. |
| Blinding (performance bias and detection bias) All outcomes | High risk | Quote: " Although blinding of the patients, parents and treating team to treat- ment group was impossible, they were blind to the aggregate group data about patient characteristics, insulin treatment and clinical or psychosocial re- sults during the study period." |
| Blinding of participants and personnel (perfor- mance bias) All outcomes | High risk | As above |
| Blinding of outcome as- sessment (detection bias) All outcomes | Low risk | Quote: " In addition, an independent evaluation team was responsible for all data collection and analysis." |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Comment: outcome data were reported for all participants. |
| Selective reporting (re- porting bias) | Low risk | Comment: outcome data were presented for all outcomes reported as mea- sured. |
| Other bias | Low risk | Comment: none was identified. |



Sartain 2001

| Methods | Randomised controlled trial Sample size calculation Intention-to-treat analysis conducted | |
|---------------|---|--|
| Participants | 399 children with acute paediatric illness (breathing difficulty, diarrhoea and vomiting or fever) Median age 13 to 15 months (range 6 weeks to 12 years) | |
| | 240 boys (60%), 159 girls (40%) 100% follow-up | |
| Interventions | 'Hospital at Home' vs usual inpatient paediatric care | |
| Outcomes | Readmissions within 90 days of discharge Length of stay Qualitative assessment of parental and patient satisfaction Cost effectiveness of both types of care | |
| | Data collected over 17 months | |

Notes

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|---|
| Random sequence genera- tion (selection bias) | Low risk | Quote: "Patients were randomised using a sealed numbered and opaque enve- lope technique." |
| Allocation concealment (selection bias) | Low risk | As above |
| Blinding (performance bias and detection bias) All outcomes | High risk | Comment: It is unlikely that participants and personnel were blinded to inter- vention measures of "hospital at home care" vs "conventional hospital care" in this study. |
| Blinding of participants and personnel (perfor- mance bias) All outcomes | High risk | As above |
| Blinding of outcome as- sessment (detection bias) All outcomes | Unclear risk | Comment: The independent clinician verified the diagnoses for all participants readmitted but it is unclear whether all study data were assessed by independent researchers. |
| Incomplete outcome data (attrition bias) All outcomes | High risk | Comment: The qualitative data collection included a purposive sample of 40 families which is 1/10th of the total study population. |
| Selective reporting (re- porting bias) | Low risk | Comment: outcome data were presented for all outcomes reported as mea- sured. |
| Other bias | Low risk | Comment: none was identified. |

Stein 1984 Methods

Randomised controlled trial



Stein 1984 (Continued)

Trusted evidence. Informed decisions. Better health.

| | Set in USA Intention-to-treat analy | ysis not conducted |
|---|---|---|
| Participants | 219 children (age below 11 years) with diverse range of chronic conditions (defined as presence of a physical condition for more than 3 months; children with moderate to severe mental retardation excluded). 85% follow-up at 6 months, 83% follow-up at 1 year. | |
| Interventions | Paediatric home care v | rs standard care |
| Outcomes | Satisfaction with care Child's psychological a Mother's psychiatric sy Impact on family Functional status meas Data collected at enrol | mptoms |
| Notes | | |
| Risk of bias | | |
| Bias | Authors' judgement | Support for judgement |
| Random sequence genera- tion (selection bias) | Low risk | Quote: "This was done using opaque sealed envelopes prepared in sets by a roll of dice to determine the order of the first assignment." |
| Allocation concealment (selection bias) | Low risk | Quote: "All scoring and randomisation procedures were carried out by a mem- ber of the research staff independently of the clinicians, and the responsible clinician was notified of the group assignment." |
| Blinding (performance bias and detection bias) All outcomes | High risk | Quote: "Because of the nature of the intervention and likelihood of discovering group assignment during data collection, no attempt was made for patients, physicians or interviewers to be blind to group assignment." |
| Blinding of participants and personnel (perfor- mance bias) All outcomes | High risk | As above |
| Blinding of outcome as- sessment (detection bias) All outcomes | High risk | As above |
| Incomplete outcome data (attrition bias) All outcomes | High risk | Quote: " A complete data set of all three interviews exists on 174 subjects or 80% of all subjects." |
| Selective reporting (re- porting bias) | Low risk | Comment: outcome data were presented for all outcomes reported as mea- sured. |
| Other bias | Low risk | Comment: none was identified. |

Stevens 2006

Methods

_

Randomised, cross-over study, 6 months subsequent phases of either home of hospital chemotherapy

| itevens 2006 (Continued) | | | |
|---|---|---|--|
| Participants | 15 allocated to home a Total 23 children receiv | lymphoblastic leukaemia (ALL) in the year prior to enrolment in study nd then hospital chemotherapy and 14 vice versa ved allocated treatment d intervention or relapsed | |
| Interventions | | pased with hospital-based chemotherapy by administered by a trained nurse | |
| Outcomes | Child's quality of life (QOL) Effects on parental caregivers Adverse effects Costs | | |
| | Data assessed at baseline, 3 months (postphase 1), 6 months (postphase 1), 3 months (postphase 2) and 6 months (postphase 2) | | |
| Notes | Cross-over study with r | no time gap between cross-over of two study arms | |
| Risk of bias | | | |
| Bias | Authors' judgement | Support for judgement | |
| Random sequence genera- tion (selection bias) | Low risk | Quote: " Using a table of random numbers, eligible children were randomly assigned by the study site manager to either the hospital (standard care) or home (treatment) chemotherapy group for phase 1 (6 months)." | |
| Allocation concealment (selection bias) | Unclear risk | Comment: the allocation concealment is not mentioned. | |
| Blinding (performance bias and detection bias) All outcomes | High risk | Comment: The study method implies that blinding of neither participants nor clinicians was possible. | |
| Blinding of participants and personnel (perfor- mance bias) All outcomes | High risk | As above | |
| Blinding of outcome as- sessment (detection bias) All outcomes | Unclear risk | Comment: Not mentioned whether data were analysed by independent clini- cians blinded to the outcome assessment. | |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Comment: outcome data were reported for all participants. | |
| Selective reporting (re- porting bias) | Low risk | Comment: outcome data were presented for all outcomes reported as mea- sured. | |
| | | | |

Strawczynski 1973

Methods

Quasi-randomised controlled trial



Strawczynski 1973 (Continued)

| Participants | 40 children with haemophilia 2 to 15 years of age 90% follow-up |
|---------------|--|
| Interventions | Home care for bleeds vs hospital |
| Outcomes | Number of bleeds Total number of treatment days Procrastination time School attendance Days in hospital Costs Duration 2 years |
| Notes | |

Risk of bias

| Bias | Authors' judgement | Support for judgement |
|---|--------------------|--|
| Random sequence genera- tion (selection bias) | High risk | Quote: "The patients were assigned to alternate groups when they presented with their first bleeding episode." |
| Allocation concealment (selection bias) | High risk | Comment: alternate group allocation would allow prediction of the allocation sequence. |
| Blinding (performance bias and detection bias) All outcomes | High risk | Comment: The study method implies that blinding of neither participants or clinicians was possible. |
| Blinding of participants and personnel (perfor- mance bias) All outcomes | High risk | As above |
| Blinding of outcome as- sessment (detection bias) All outcomes | Unclear risk | Comment: Not mentioned whether data was analysed by independent clini- cians blinded to the outcome assessment. |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Comment: outcome data were reported for all participants. |
| Selective reporting (re- porting bias) | Low risk | Comment: outcome data were presented for all outcomes reported as mea- sured. |
| Other bias | Low risk | Comment: none was identified. |

Tie 2009

| Methods | Prospective randomised controlled trial, pilot study |
|--------------|---|
| Participants | 44 out of 58 eligible children with acute bronchiolitis enrolled Age group 3 to 24 months Oxygen saturation on arrival > 92% in both groups |



| ie 2009 (Continued) | Observed and clinically stable for > 24 hrs in hospital 100% follow-up | | | |
|---|--|--|--|--|
| Interventions | Home oxygen therapy vs hospital care | | | |
| Outcomes | Readmission to hospital within 7 days of discharge home Total duration of hospitalisation | | | |
| Notes | | | | |
| Risk of bias | | | | |
| Bias | Authors' judgement | Support for judgement | | |
| Random sequence genera- tion (selection bias) | Unclear risk | Comment: The method of random sequence generation not mentioned. | | |
| Allocation concealment (selection bias) | Low risk | Quote: "Researchers were blinded and only following informed consent was the management allocation revealed." | | |
| Blinding (performance bias and detection bias) All outcomes | High risk | Comment: The nature of study intervention, 'hospital in the home' vs tradi- tional hospitalisation indicates that blinding of participants or clinicians was not possible. | | |
| Blinding of participants and personnel (perfor- mance bias) All outcomes | High risk | As above | | |
| Blinding of outcome as- sessment (detection bias) All outcomes | Unclear risk | Coment: Not mentioned whether data were analysed by independent clini- cians blinded to the outcome assessment. | | |
| Incomplete outcome data (attrition bias) All outcomes | Low risk | Comment: outcome data were reported for all participants. | | |
| Selective reporting (re- porting bias) | Low risk | Comment: outcome data were presented for all outcomes reported as mea- sured. | | |
| Other bias | Low risk | Comment: none was identified. | | |

Characteristics of excluded studies [ordered by study ID]

| Study | Reason for exclusion | |
|----------------|--|--|
| Affleck 1989 | Home support rather than specialist home-based nursing | |
| Alexander 1988 | Clinic-based intervention | |
| Badr 2006 | Parental education by nurses (not home-based nursing) | |
| Bang 1999 | Community-based health visitor trial | |
| Beckwith 1988 | Home support rather than specialist home-based nursing | |



| Study | Reason for exclusion | |
|----------------------|---|--|
| Black 1995 | Home visitors for non-organic failure to thrive | |
| Blakeslee 1997 | Non-randomised comparison study | |
| Brooten 1986 | Low birth weight is not an acute or chronic illness | |
| Brown 2002 | Asthma education (not home-based nursing) | |
| Broyles 2000 | Clinic-based intervention | |
| Butz 1994 | Asthma education (not home-based nursing) | |
| Butz 2006 | Asthma education (not home-based nursing) | |
| Caliskan Yilmaz 2010 | Quasi-experimental study design (no randomisation) | |
| Casiro 1993 | Low birth weight is not an acute or chronic illness | |
| Catov 2005 | Asthma education (not home-based nursing) Quasi-experimental design (non-randomised) | |
| Crowin 2005 | Age group 16 years and older with no details available of study participants aged between 16 and 18 age. Authors contacted but no response received | |
| Deaves 1993 | Asthma education (not home-based nursing) | |
| Dolinar 2000 | Asthma education (not home-based nursing) | |
| Donati 1987 | Non-randomised comparison study | |
| Greineder 1999 | Asthma education (not home-based nursing) | |
| Hatziagorou 2010 | Not a randomised controlled trial | |
| Heikens 1994 | Community health aides for protein calorie malnutrition | |
| Hughes 1991 | Asthma education (not home-based nursing) | |
| Keefe 2006 | Parent education programme (not home-based nursing) | |
| Krieger 2009 | Asthma education (not home-based nursing) | |
| Lindberg 2002 | Non-randomised comparison study | |
| Madge 1997 | Asthma education (not home-based nursing) | |
| McConnochie 1999 | Asthma education (not home-based nursing) | |
| Mitchell 1986 | Asthma education (not home-based nursing) | |
| Park 2002 | Non-randomised comparison study | |
| Raynor 1999 | Specialist health visitor for non-organic failure to thrive | |
| Rieger 1995 | Non-randomised comparison study | |



| Study | Reason for exclusion | |
|------------------|--|--|
| Schuttelaar 2010 | Nurse practitioner care in the hospital outpatient clinic, no home-based treatment | |
| Shin 2009 | Parental education for early intervention (not home-based nursing) | |
| Svahn 2002 | Non randomised comparison study, inappropriate age group | |
| Svahn 2008 | Non-randomised comparison study | |
| Talabere 1993 | Asthma education (not home-based nursing) | |
| Tiberg 2011 | Home care in hospital grounds, not home-based care | |
| Wang 1995 | Presurgery assessments, not home nursing | |
| Wolter 1997 | Participants with cystic fibrosis give their own therapy - no nurses | |
| Wright 1998 | Home visitors for non-organic failure to thrive | |

ADDITIONAL TABLES

Table 1. Characteristics of interventions showing diversity of services

| Study | Study set- tings | Type of illness - intervention | Hours Available | No. visits possi- ble | Staff Qualifica- tions | Information provided |
|-------------------|--|---|---|---|--|---|
| Burke 1997 | Queen's Uni- versity, Uni- versity of Toronto, On- tario, Canada | Children with chronic condi- tions - stress- point interven- tion for parents | Not report- ed. Direct phone con- tact with nurse related to 'stres- sors' rather than clini- cal events. | Not reported. | "master's pre- pared nurse". | Mailed summaries and reminders of families own stress points and coping strategies. |
| Dougherty 1998 | The Montre- al Children's Hospital, Mon- treal, Quebec, Canada | Newly diag- nosed children with IDDM - Home based treatment | 24 hour telephone consulta- tion. | Home visits once/ twice daily for first 2 - 3 days, teach- ing visit at clin- ic after 2 weeks, follow-up outpa- tient visits at 2 - 3 months. | Diabetes treat- ment nurse, as part of a team of dia- betes-specialist physician, psy- chologist and social worker. | Teaching session, in- struction and supervision in practical and theoreti- cal aspects of treatment. |
| Sartain 2001 | Arrow Park Hospital, Up- ton, Wirral, UK | Acute illness (breathing diffi- culty, diarrhoea and vomiting or fever) - Hos- pital at home | 24 hours, 7 days. Ser- vice un- til 2300, on-call overnight. | 1 - 4 daily | Not specified. Participants under care of hospital consul- tants. | Patient information booklets detailing pos- sible course of illness, signs and symptoms of potential deterioration, appropriate treatment, contact information. |



Table 1. Characteristics of interventions showing diversity of services (Continued)

scheme treatment

| | | ment | | | | |
|------------------------|---|---|--|---|--|--|
| Stein 1984 | Paediatric Ambulatory Care Division, Albert Ein- stein College of Medicine, Bronx, New York | Heteroge- nous group of children with chronic illness- es - Paediatric home care | Not speci- fied. | Minimum pack- age: initial home visit, one con- tact (by person or by telephone) each month for 6 months. Most par- ticipants received more. | Team of gen- eralist paedi- atricians, pae- diatric nurse practition- ers and social worker. | "Teach child and fami- ly about condition and train them in self-care skills". |
| Stevens 2006 | University of Toronto and Sick Kid's Hos- pital, Toronto, Canada | Children with acute lym- phoblastic leukemia - Home chemotherapy | Not speci- fied. | Home visits for administration of chemotherapy as per the protocol. | Trained nurse from a commu- nity health ser- vices agency administered chemotherapy, routine contact by community nurse to the on- cologist if con- cerns. | Parents encouraged to contact primary nurse at the oncology clinic at the hospital as required |
| Strawczyn- ski 1973 | The Montre- al Children's Hospital, Mon- treal, Quebec, Canada | Management of bleeding in he- mophilic chil- dren | 24-hour service when in- cident (bleed- ing in chil- dren with haemophil- ia) oc- curred. | Not specified. | Specially trained nurse. | No educational interven- tion reported. |
| Tie 2009 | Department of General Paediatrics, Princess Mar- garet Hospital for children, Perth, West- ern Australia | Acute bronchi- olitis - home oxygen therapy | Not speci- fied | "Hospital in the home (HiTH)" nurse home vis- it within 12 hours of hospital dis- charge, minimum of 2 home visits, one phone con- tact with parents in every 24-hour period. | Not specified. | Parents were educated on home oxygen use and instructed on how to ob- serve their children for signs of clinical deterio- ration. |

APPENDICES

Appendix 1. Search strategies February 2012

Cochrane Central Register of Controlled Trials (CENTRAL), part of the Cochrane Library, 2012 (Issue 1), searched 9 February 2012

#1MeSH descriptor: [Home Care Services] 1 tree(s) exploded#2MeSH descriptor: [Nurse Clinicians] this term only#3MeSH descriptor: [Community Health Nursing] this term only



#4MeSH descriptor: [Nurse Practitioners] this term only #5MeSH descriptor: [Home Nursing] 3 tree(s) exploded #6((home or homebased or domicil*) near/3 treat*):ti,ab,kw #7((home or homebased or domicil*) near/3 (care or caring)):ti,ab,kw #8((home or homebased or domicil*) near/3 healthcar*):ti,ab,kw #9(nursing near/3 outreach):ti,ab,kw #10((paediatric or pediatric) near/3 ambulat*):ti,ab,kw #11((home or home based or domicil*) near/3 nurs*) #12(hospital near/3 home):ti,ab,kw #13#1 or #2 or #3 or #4 or #5 or #6 or #7 or #8 or #9 or #10 or #11 or #12 #14(baby or babies or infant* or toddler* or child* or preschool* or pre next school* or teen* or adolescen*):ti,ab,kw #15MeSH descriptor: [Child] 1 tree(s) exploded #16MeSH descriptor: [Infant] 1 tree(s) exploded #17MeSH descriptor: [Adolescent] this term only #18#14 or #15 or #16 or #17 #19#13 and #18 Ovid MEDLINE 1946 to Jan Week 4 2012, searched 8 February 2012 1 exp Home Care Services/ (36908) 2 Home Care Services, Hospital-Based/ (1474) 3 Community Health Nursing/ (17645) 4 nurse clinicians/ (7117) 5 nurse practitioners/ (14200) 6 exp home nursing/ (8281) 7 ((home or homebased or domicil\$) adj3 treat\$).tw. (3333) 8 ((home or homebased or domicil\$) adj3 (care\$ or caring)).tw. (18937) 9 ((home or homebased or domicil\$) adj3 healthcar\$).tw. (835) 10 (nursing adj3 outreach).tw. (55) 11 (p?ediatric adj3 ambulatory\$).tw. (496) 12 ((home or homebased or domicil\$) adj3 nurs\$).tw. (17291) 13 (hospital adj3 home).tw. (3967) 14 or/1-13 (92524) 15 exp Infant/ (856559) 16 exp child/ (1405066) 17 adolescent/ (1440129) 18 (baby or babies or infant\$ or toddler\$ or child\$ or preschool\$ or pre-school\$ or teen\$ or adolescen\$).tw. (1125366) 19 or/15-18 (2771225) 20 randomized controlled trial.pt. (317638) 21 controlled clinical trial.pt. (83333) 22 randomi#ed.ab. (266278) 23 placebo\$.ab. (128230) 24 drug therapy.fs. (1495532) 25 randomly.ab. (161461)

- 26 trial.ab. (229896)
- 27 groups.ab. (1067026)
- 28 or/20-27 (2777450)
- 29 exp animals/ not humans.sh. (3647344)
- 30 28 not 29 (2358297)
- 31 14 and 19 and 30 (2971)
- 32 limit 31 to yr="2005 -Current" (1009)

EMBASE (Ovid), 1980 to 2012 Week 05, searched 9 February 2012

exp home care/ (46930)
 community health nursing/ (24600)
 exp nurse practitioner/ (15820)
 ((home or homebased or domicil\$) adj3 treat\$).tw. (4377)
 ((home or homebased or domicil\$) adj3 (care\$ or caring)).tw. (21637)
 ((home or homebased or domicil\$) adj3 healthcar\$).tw. (900)
 (nursing adj3 outreach).tw. (70)
 (p?ediatric adj3 ambulatory\$).tw. (841)
 ((home or homebased or domicil\$) adj3 nurs\$).tw. (20285)



10 (hospital adj3 home).tw. (4760)

Trusted evidence. Informed decisions. Better health.

11 or/1-10 (108582) 12 exp child/ (1518691) 13 exp adolescent/ (1132793) 14 (baby or babies or infant\$ or toddler\$ or child\$ or preschool\$ or pre-school\$ or teen\$ or adolescen\$).tw. (1316929) 15 12 or 13 or 14 (2508849) 16 11 and 15 (16654) 17 exp Clinical trial/ (880196) 18 Randomized controlled trial/ (296765) 19 Randomization/ (55652) 20 Single blind procedure/ (14775) 21 Double blind procedure/ (102866) 22 Crossover procedure/ (31796) 23 Placebo/ (192025) 24 Randomi#ed.tw. (372082) 25 RCT.tw. (8444) 26 (random\$ adj3 (allocat\$ or assign\$)).tw. (88417) 27 randomly.ab. (203108) 28 groups.ab. (1321946) 29 trial.ab. (293317) 30 ((singl\$ or doubl\$ or trebl\$ or tripl\$) adj3 (blind\$ or mask\$)).tw. (135186) 31 Placebo\$.tw. (164757) 32 Prospective study/ (181795) 33 (crossover or cross-over).tw. (57030) 34 prospective.tw. (350288) 35 or/17-34 (2628978) 36 16 and 35 (2991) 37 limit 36 to yr="2005 -Current" (1335) PsycINFO (Ovid) 1806 to Feb Week 1 2012, searched 9 February 2012 1 home care/ (3786) 2 ((home or homebased or domicil\$) adj3 treat\$).tw. (1198) 3 ((home or homebased or domicil\$) adj3 (care\$ or caring)).tw. (6838) 4 ((home or homebased or domicil\$) adj3 healthcar\$).tw. (84) 5 (nursing adj3 outreach).tw. (18) 6 ((home or homebased or domicil\$) adj3 nurs\$).tw. (7345) 7 (hospital adj3 home).tw. (912) 8 (p?ediatric adj3 ambulatory\$).tw. (40) 9 (nurse adj3 (specialist\$ or practitioner or clinican)).tw. (1067) 10 or/1-9 (16351) 11 (adolescence 13 17 yrs or childhood birth 12 yrs).ag. (527933) 12 (baby or babies or infant\$ or toddler\$ or child\$ or preschool\$ or pre-school\$ or teen\$ or adolescen\$).tw. (609887) 13 11 or 12 (767428) 14 clinical trials/ (5796) 15 (randomis\$ or randomiz\$).tw. (35991) 16 (random\$ adj3 (allocat\$ or assign\$)).tw. (25127) 17 ((clinic\$ or control\$) adj trial\$).tw. (30613) 18 ((singl\$ or doubl\$ or trebl\$ or tripl\$) adj3 (blind\$ or mask\$)).tw. (17112) 19 (crossover\$ or "cross over\$").tw. (5899) 20 random sampling/ (527) 21 Experiment Controls/ (636) 22 Placebo/ (3125) 23 placebo\$.tw. (27173) 24 exp program evaluation/ (14599) 25 treatment effectiveness evaluation/ (13106) 26 ((effectiveness or evaluat\$) adj3 (stud\$ or research\$)).tw. (46630) 27 or/14-26 (151249) 28 10 and 13 and 27 (293) 29 limit 28 to yr="2005 -Current" (143) CINAHL Plus (EBSCOhost), 1937 to current, searched 9 February 2012



S34 S17 and S32 S33 S17 and S32 S32 S18 or S19 or S20 or S21 or S22 or S23 or S24 or S25 or S26 or S27 or S28 or S29 or S30 or S31 S31 TI (evaluat* study or evaluat* research) or AB (evaluate* study or evaluat* research) or TI (effectiv* study or effectiv* research) or AB (effectiv* study or effectiv* research) OR TI (prospectiv* study or prospectiv* research) or AB(prospectiv* study or prospectiv* research) or TI (follow-up study or follow-up research) or AB(follow-up study or follow-up research) S30 placebo* S29 crossover* or "cross over*" S28 (MH "Crossover Design") S27 (tripl* N3 mask*) or (tripl* N3 blind*) S26 (trebl* N3 mask*) or (trebl* N3 blind*) S25 (doubl* N3 mask*) or (doubl* N3 blind*) S24 (singl* N3 mask*) or (singl* N3 blind*) S23 (clinic* N3 trial*) or (control* N3 trial*) S22 (random* N3 allocat*) or (random* N3 assign*) S21 randomis* or randomiz* S20 (MH "Meta Analysis") S19 (MH "Clinical Trials+") S18 MH random assignment S17 S11 and S16 S16 S13 or S14 S15 S15 (baby or babies or infant* or toddler* or child* or preschool* or pre-school* or teen* or adolescen*) S14 (MH "Adolescence") S13 (MH "Child") OR (MH "Child, Preschool") OR (MH "Infant+") S12 infant S11 S1 or S2 or S3 or S4 or S5 or S6 or S7 or S8 or S9 or S10 S10 (hospital N3 home) S9 ((home or homebased or domicil*) N3 nurs*) S8 (p#ediatric N3 ambulatory) S7 (nursing N3 outreach) S6 ((home or homebased or domicil*) N3 healthcar*) S5 ((home or homebased or domicil*) N3 (care* or caring)) S4 ((home or homebased or domicil*) N3 treat*) S3 (MH "Advanced Practice Nurses+") S2 (MH "Community Health Nursing+") OR (MH "Ambulatory Care Nursing") OR (MH"Specialties, Nursing") S1 (MH "Home Health Care+") Sociological Abstracts (Proquest) 1952 to current, searched 9 February 2012 (SU.EXACT("Home Health Care") OR AB((home OR homebased OR domicil*) NEAR/3 treat*) OR TI((home OR homebased OR domicil*) NEAR/3 treat*) OR AB((home OR homebased OR

domicil*) NEAR/3 (care* OR caring)) OR TI((home OR homebased OR domicil*) NEAR/3

(care* OR caring)) OR AB((home OR homebased OR domicil*) NEAR/3 nurs*) OR TI((home

OR homebased OR domicil*) NEAR/3 nurse*) OR AB((home OR homebased OR domicil*)

NEAR/3 (healthcare*)) OR TI((home OR homebased OR domicil*) NEAR/3 (healthcare*)) OR

AB(nursing NEAR/3 outreach*) OR TI(nursing NEAR/3 outreach) OR AB(p*ediatric NEAR/3

ambulatory) OR TI(p*ediatric NEAR/3 ambulatory) OR AB(hospital NEAR/3 home) OR

TI(hospital NEAR/3 home) OR AB((nurs*) NEAR/3 (specialist* OR clinician* OR practitioner*))

or TI((nurs*) NEAR/3 (specialist* OR clinician* OR practitioner*))) AND (SU.EXACT("Infants"

OR "adolescents" OR "Children") OR AB(baby OR babies OR infant* OR toddler* OR child*



OR preschool* OR preschool* OR teen* OR adolescen*) OR TI(baby OR babies OR infant*

OR toddler* OR child* OR preschool* OR preschool* OR teen* OR adolescen*)))

ClinicalTrials.gov All years, searched 10 February 2012

home nursing NOT "nursing home" | Child

ICTRP All years, searched 10 February 2012

home nursing and child*

WHAT'S NEW

| Date | Event | Description |
|-----------------|--|--|
| 29 January 2013 | New citation required and conclusions have changed | Two new studies included. |
| 30 October 2012 | New search has been performed | Updated search. Two new studies found. 'Risk of bias' assess- ments completed for all included studies. |

HISTORY

Protocol first published: Issue 3, 2003 Review first published: Issue 4, 2006

| Date | Event | Description |
|-----------------|--|---------------------------------|
| 9 November 2008 | Amended | Converted to new review format. |
| 1 August 2006 | New citation required and conclusions have changed | Substantive amendment |

CONTRIBUTIONS OF AUTHORS

All review authors contributed to the writing of the review.

DECLARATIONS OF INTEREST

Drs Woolfenden, Cooper and Piper co-ordinate or have co-ordinated ambulatory paediatric services in New South Wales, Australia. Dr Cooper is employed as a Staff Specialist Paediatrician at the Royal North Shore Hospital, Australia. Dr Parab is employed as a Staff Specialist Paediatrician at Wollongong in Illawarra Shoalhaaven Local Health District, Australia.

INDEX TERMS

Medical Subject Headings (MeSH)

Acute Disease [*nursing]; Chronic Disease [*nursing]; Home Care Services, Hospital-Based [*organization & administration] [*standards]; Home Nursing; Hospitalization [statistics & numerical data]; Length of Stay [statistics & numerical data]; Randomized Controlled Trials as Topic

MeSH check words

Adolescent; Child; Child, Preschool; Humans; Infant; Infant, Newborn