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Medical day hospital care for older people versus alternative forms of care (Review)



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[Intervention Review]

Medical day hospital care for older people versus alternative forms of care

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ABSTRACT

Background

The proportion of the world's population aged over 60 years is increasing. Therefore, there is a need to examine different methods of healthcare provision for this population. Medical day hospitals provide multidisciplinary health services to older people in one location.

Objectives

To examine the effectiveness of medical day hospitals for older people in preventing death, disability, institutionalisation and improving subjective health status.

Search methods

Our search included the Cochrane Effective Practice and Organisation of Care (EPOC) Group Register of Studies, CENTRAL (2013, Issue 7), MEDLINE via Ovid (1950-2013), EMBASE via Ovid (1947-2013) and CINAHL via EbscoHost (1980-2013). We also conducted cited reference searches, searched conference proceedings and trial registries, hand searched select journals, and contacted relevant authors and researchers to inquire about additional data.

Selection criteria

Randomised and quasi-randomised trials comparing medical day hospitals with alternative care for older people (mean/median > 60 years of age).

Data collection and analysis

Two authors independently assessed trial eligibility and risk of bias and extracted data from included trials. We used standard methodological procedures expected by the Cochrane Collaboration. Trials were sub-categorised as comprehensive care, domiciliary care or no comprehensive care.

Main results

Sixteen trials (3689 participants) compared day hospitals with comprehensive care (five trials), domiciliary care (seven trials) or no comprehensive care (four trials). Overall there was low quality evidence from these trials for the following results.

For the outcome of death, there was no strong evidence for or against day hospitals compared to other treatments overall (odds ratio (OR) 1.05; 95% CI 0.85 to 1.28; P = 0.66), or to comprehensive care (OR 1.26; 95% CI 0.87 to 1.82; P = 0.22), domiciliary care (OR 0.97; 95% CI 0.61 to 1.55; P = 0.89), or no comprehensive care (OR 0.88; 95% CI 0.63 to 1.22; P = 0.43).

For the outcome of death or deterioration in activities of daily living (ADL), there was no strong evidence for day hospital attendance compared to other treatments (OR 1.07; 95% CI 0.76 to 1.49; P = 0.70), or to comprehensive care (OR 1.18; 95% CI 0.63 to 2.18; P = 0.61), domiciliary care (OR 1.41; 95% CI 0.82 to 2.42; P = 0.21) or no comprehensive care (OR 0.76; 95% CI 0.56 to 1.05; P = 0.09).

For the outcome of death or poor outcome (institutional care, dependency, deterioration in physical function), there was no strong evidence for day hospitals compared to other treatments (OR 0.92; 95% CI 0.74 to 1.15; P = 0.49), or compared to comprehensive care (OR 1.05; 95% CI 0.79 to 1.40; P = 0.74) or domiciliary care (OR 1.08; 95% CI 0.67 to 1.74; P = 0.75). However, compared with no comprehensive care there was a difference in favour of day hospitals (OR 0.72; 95% CI 0.53 to 0.99; P = 0.04).

For the outcome of death or institutional care, there was no strong evidence for day hospitals compared to other treatments overall (OR 0.85; 95% CI 0.63 to 1.14; P = 0.28), or to comprehensive care (OR 1.00; 95% CI 0.69 to 1.44; P = 0.99), domiciliary care (OR 1.05; 95% CI 0.57 to 1.92; P = 0.88) or no comprehensive care (OR 0.63; 95% CI 0.40 to 1.00; P = 0.05).

For the outcome of deterioration in ADL, there was no strong evidence that day hospital attendance had a different effect than other treatments overall (OR 1.11; 95% CI 0.68 to 1.80; P = 0.67) or compared with comprehensive care (OR 1.21; 0.58 to 2.52; P = 0.61), or domiciliary care (OR 1.59; 95% CI 0.87 to 2.90; P = 0.13). However, day hospital patients showed a reduced odds of deterioration compared with those receiving no comprehensive care (OR 0.61; 95% CI 0.38 to 0.97; P = 0.04) and significant subgroup differences (P = 0.04).

For the outcome of requiring institutional care, there was no strong evidence for day hospitals compared to other treatments (OR 0.84; 95% CI 0.58 to 1.21; P = 0.35), or to comprehensive care (OR 0.91; 95% CI 0.70 to 1.19; P = 0.49), domiciliary care (OR 1.49; 95% CI 0.53 to 4.25; P = 0.45), or no comprehensive care (OR 0.58; 95% CI 0.28 to 1.20; P = 0.14).

Authors' conclusions

There is low quality evidence that medical day hospitals appear effective compared to no comprehensive care for the combined outcome of death or poor outcome, and for deterioration in ADL. There is no clear evidence for other outcomes, or an advantage over other medical care provision.

PLAIN LANGUAGE SUMMARY

Medical day hospital care for the elderly versus alternative forms of care

Day hospitals are one way of delivering healthcare to older people. They are out-patient facilities which older patients attend for a full or near full day and receive multidisciplinary health care 'under one roof.' Sixteen trials involving 3689 participants were included in this review and compared day hospitals with other comprehensive services (including inpatient and outpatient services), home based care and no comprehensive services. Attendance at a day hospital offers benefits compared to providing no treatment which include reducing the risk of needing more help with daily activities such as washing or dressing. Furthermore, patients are less likely to suffer one of the following: dying, being institutionalised or becoming more dependent on others. There is no apparent benefit when day hospitals are compared with other comprehensive services or home care. The economic value of day hospitals when compared with other health care services remains unclear.

SUMMARY OF FINDINGS FOR THE MAIN COMPARISON [Explanation]

Day hospitals compared to alternative or no care for rehabilitation

Patient or population: patients with rehabilitation needs

Intervention: day hospitals
Comparison: alternative care

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence Comments (GRADE)
	Assumed risk	Corresponding risk			
	Alternative or no care	Day hospitals			
Death by the end of fol-	Study population		OR 1.05	3533	⊕⊕○○ L1 2 3
low up Follow-up: median 12 months	p: median 12 127 per 1000 132 per 1000 (0.85 to 10)	(0.85 to 1.28)	(16 studies)	low ^{1,2,3}	
	Moderate				
	66 per 1000	69 per 1000 (57 to 83)			
Death or institutional	Study population		OR 0.85	3030	⊕⊕⊖⊖ I1 2 3
care by the end of fol- low up Follow-up: median 12 months	303 per 1000	270 per 1000 (215 to 331)	(0.63 to 1.14)	(13 studies)	low ^{1,2,3}
	Moderate				
	221 per 1000	194 per 1000 (152 to 244)			

Death or deterioration	, · ·		OR 1.07	1268	0 00
in activities of daily living (ADL) Follow-up: median 12 months	407 per 1000	423 per 1000 (343 to 506)	(0.76 to 1.49)	(7 studies)	low ^{1,2,3}
	Moderate				
	430 per 1000	447 per 1000 (364 to 529)			
Death or poor outcome (institutional care, dis- ability or deterioration) Follow-up: median 12 months			OR 0.92	2831	⊕⊕○○
	365 per 1000	346 per 1000 (299 to 398)	(0.74 to 1.15)	(13 studies)	low ^{1,2,3}
months	Moderate				
	241 per 1000	226 per 1000 (190 to 267)			
Deterioration in ADL in	Study population		OR 1.11	905	⊕⊕○○
survivors Various ADL measures	251 per 1000	271 per 1000 (185 to 376)	(0.68 to 1.8)	(7 studies)	low ^{1,2,3}
	Moderate				
	233 per 1000	252 per 1000 (171 to 354)			

^{*}The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; OR: Odds ratio; ADL: activities of daily living

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Limitations for at least one risk of bias criterion, or some limitations for multiple criteria, sufficient to lower confidence in the estimate of effect

 $^{^{2}}$ Whilst there was evidence of heterogeneity, this was anticipated due to the diversity of the populations and of the interventions

³ Wide Cls

BACKGROUND

The first geriatric day hospital was opened in the UK in 1952 (Farndale 1961). Day hospitals developed rapidly in the United Kingdom in the 1960's as an important component of care provision for older people designed to complement in-patient services (Black 2005). The model has since been widely applied in New Zealand, Australia, Canada, the USA and several European countries.

Geriatric day hospitals provide multi-disciplinary rehabilitation in an outpatient setting and operate in a pivotal position between hospital and home-based services (Ames 1995; Black 2005; Brocklehurst 1973; Petermans 2011). They provide specialist services for older people, which can include examinations and consultations, all concentrated in one location (Bussche 2010).

Although there is considerable descriptive literature on day hospital care (RCP 1994), concern has been expressed that evidence for effectiveness is equivocal (Brocklehurst 1980; Donaldson 1986) and that day hospital care is expensive (NAO 1994).

Concern is often expressed about the most appropriate health and social services required to address the needs of an aging population. In the UK, for example, the largest population increase is seen in the over 85 age group. A range of different services models, of which the day hospital is one, may be appropriate to address these needs. This review sets out to examine the effectiveness and resource implications of geriatric medical day hospital attendance for older people and to compare it with other models of healthcare delivery for an older population. This is an updated Cochrane review first published in Forster 1999a.

Description of the condition

Geriatric day hospitals are not usually specific to one condition. However, many will provide rehabilitation services appropriate to conditions such as stroke that are likely to be seen in an older population.

Description of the intervention

Geriatric day hospitals are out-patient healthcare facilities for older people living in the community. They provide multi-professional treatment on a full or part time basis (Beynon 2009). They serve several functions, including assessment, rehabilitation, and medical, nursing, maintenance, social and respite care (Brocklehurst 1980). Rehabilitation and maintenance comprise the main work of the day hospital: 42% and 23% respectively (RCP 1994), with rehabilitation regarded as the most important function (Brocklehurst 1980). The specific features and services offered by individual geriatric day hospitals are subject to considerable variation. However, they usually include a combination of medical

assessment with support from nurses and allied health professionals, often including physiotherapists and occupational therapists. There is no consensus on what types of healthcare professionals should make up the multi-disciplinary teams (Petermans 2011). Additional services such as chiropody, social work, exercise classes and assistance with bathing and hair washing are offered by some hospitals.

How the intervention might work

Geriatric day hospitals offer a multidisciplinary approach to assessment and rehabilitation, with provision of a variety of services in one location. As a result of assessment and treatment occurring 'under one roof,' the health requirements of older people should be identified and responded to in an appropriate and timely manner. The day hospital can provide out-patient delivery of a Comprehensive Geriatric Assessment (CGA) which has a robust evidence base for inpatient setting use (Ellis 2011). A CGA addresses medical, physical, psychological and social needs, and includes the formation of a plan of care and rehabilitation, with a clear method of implementation. Day hospital staff have specific skills, knowledge and experience related to working with older people. Furthermore, the day hospital environment has the advantage of providing social interaction between patients, a factor which domiciliary services and usual care cannot provide. These factors could result in better outcomes for patients through the provision of effective rehabilitation and other healthcare delivery for an older population.

Why it is important to do this review

Between 1985 and 2010 the proportion of the world's population that is aged over 65 years grew by approximately a quarter from 6.0% (291 million) to 7.6% (524 million), and is expected to increase to 13% by 2035, exceeding a billion people globally (UN 2011). As a result of this increase, providing health care that meets the diverse needs of an older population and is cost effective and efficient will be ever more important. Day hospitals are one way of delivering multidisciplinary rehabilitation to older people in an outpatient setting. This review is necessary to assess the effectiveness of day hospitals across a number of health, cost and resource outcomes.

OBJECTIVES

The primary question was whether older patients attending a geriatric medical day hospital would experience better outcomes (in terms of death, dependency or institutionalisation) than those receiving alternative forms of care.

Secondary questions concerned the impact of day hospital care on patient satisfaction and subjective health outcomes, carer distress and resource use and costs.

METHODS

Criteria for considering studies for this review

Types of studies

We included studies that were of a prospective, controlled design in which there was random assignment of participants to alternative treatment groups (one of which involved day hospital care), not as part of a complex multi-service intervention. Studies which utilised 'quasi' randomisation procedures (for example allocation to groups based on date of birth) were also included.

Types of participants

We included patients receiving medical care (mean/median age of >60 years for individual studies). We are aware that day hospital descriptive studies have indicated that day hospital attendance is determined more by needs than age and that younger patients do attend day hospitals. Our pre-specified participant criterion of age 60 years and over was chosen to pragmatically capture this clinical practice. Studies which were specific to psychiatric patients were excluded.

Types of interventions

We defined a day hospital as an out-patient facility where older patients attend for a full or near full day and receive multidisciplinary rehabilitation in a healthcare setting. This is consistent with previous definitions (Siu 1994) and excluded trials evaluating social day centres, or other types of day hospitals such as psychiatric day hospitals for patients with dementia or psychiatric conditions. We excluded studies on day hospitals that only provided services for single, specific conditions (for example, arthritis). We wanted to assess the effects of providing typical general assessment and rehabilitation services relevant to older people. The inclusion of disease-specific trials would risk incorporating the effects of very specific therapies for specific conditions, which were not the focus of this review.

We anticipated considerable heterogeneity, particularly in the control services, and so pre-specified key subgroup comparisons prior to reviewing the trials.

1) Day hospital care versus comprehensive care - where control patients had access to a range of geriatric medical services (both inpatient and outpatient).

- 2) Day hospital care versus domiciliary care where control patients were provided an approximately equivalent rehabilitation program within their own home or social day centre.
- 3) Day hospital care versus no comprehensive care where control patients did not routinely have access to outpatient rehabilitation services.

Types of outcome measures

We wished to identify outcomes which reflected a previous definition of the purpose of day hospital care: to facilitate and prolong independent living for older people in the community (Donaldson 1987). Effective day hospital care would thus be expected to reduce death, to maintain older people in their own home and to reduce admissions to hospital. The following outcomes were therefore selected, all of which were recorded at the end of scheduled follow up.

Primary outcomes

Primary outcomes were:

- death;
- the need for institutional care;
- dependency;
- global 'poor outcome' comprising death or one of the following (in order of preference): resident in institutional care, severe dependency at end of follow up, or deterioration in physical function during follow up; this outcome was included in anticipation of incomplete data sets.

Secondary outcomes

Secondary outcomes included:

- dependency, measured by activities of daily living (ADL)
- patient satisfaction;
- subjective health status (including mood);
- resource use (in hospital or institutional care) plus overall cost analyses;
 - carer distress.

We considered all studies that met the eligibility criteria for study design, participants and interventions regardless of whether the pre-specified primary or secondary outcomes were reported.

Search methods for identification of studies

For this edition of the review, D Andre, University of Leeds Library, developed search strategies in consultation with the authors. They were peer reviewed by M Fiander, EPOC Trials Search Coordinator. We searched the databases listed below for relevant studies.

Electronic searches

- Effective Practice and Organisation of Care (EPOC) group register of trials (August 2013);
- Cochrane Central Register of Controlled Trials (CENTRAL; *The Cochrane Library*, Issue 7, July 2013; Appendix 1);
 - MEDLINE (1996 to July 2013; Appendix 2);
 - Medline in Process (1996 to August 2013; Appendix 3);
 - EMBASE (1996 to August 2013; Appendix 4);
- Cumulative Index to Nursing and Allied Health Literature (CINAHL; 1996 to August 2013; Appendix 5);
- Allied and Complementary Medicine Database (AMED; 1996 to August 2013; Appendix 6);
- Physiotherapy Evidence Database (PEDro; August 2008; Appendix 7);
- Applied Social Science Index and Abstracts (ASSIA; 1996 to August 2013; Appendix 8);
- International Bibliography of Social Sciences (IBSS; 1996 to August 2013; Appendix 9);
 - PsycINFO (1996 to August week 1, 2013; Appendix 10);
- Health Management Information Consortium Database (HMIC; January 2008 to August 2013; Appendix 11);
- NHS Economic Evaluation Database (NHS EED; searched October 2013; Appendix 12);
- Health Technology Assessment (HTA) Database (searched October 2013; Appendix 12);
 - Web of Knowledge (1996 to August 2013; Appendix 13);
- Web of Science, Conference Proceedings Citation Index Social Science (1990 to 2012; Appendix 14);
 - Google Scholar (searched August 2013; Appendix 15);
 - Index to Theses (1996 to August 2013; Appendix 16);
- Proquest Dissertations and Theses (1996 to August 2013; Appendix 17);
- Current Controlled Trials (searched August 2013; Appendix 18).

Search strategies were comprised of natural language (free text) terms and controlled vocabulary (index) terms. Language limits were not applied. Search strategies for this update have been revised in order to improve sensitivity and precision. Changes were made based on an analysis of indexing terms found on previously included studies and by testing terms from the original strategy for precision. Given these changes, searches have been run retrospectively. The results of this search have been de-duplicated from searches we carried out for the previous update of this review in 2008. The reference list of reviews of potential relevance were also examined (Bours 1998; Mason 2007; Outpatient Service 2004; Prvu Bettger 2007; Petermans 2011).

Searching other resources

• HSRProj (searched August 2013; Appendix 19);

- National Research Register (searched September 2007);
- Australian New Zealand Clinical Trials Registry (May 2008).

Data collection and analysis

Selection of studies

Two review authors independently assessed the titles and abstracts from the electronic searches and excluded obviously irrelevant studies. We obtained full text articles of the remaining studies and at least two review authors independently assessed these against pre-specified inclusion criteria to determine which trials would be eligible for inclusion. Study authors were contacted for further details when necessary. Disagreements were resolved by discussion with other members of the review team.

Data extraction and management

At least two review authors extracted data independently. Disagreements were resolved through group consensus. When possible, we contacted study authors for additional information and data as required.

Assessment of risk of bias in included studies

Two review authors independently assessed risk of bias in the included studies using the tool for assessing risk of bias in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011). We scored each study as being at 'high risk of bias', 'low risk of bias' or 'unclear risk of bias' for each of the following domains, and reported them in the 'Risk of bias' tables.

- Random sequence generation (selection bias).
- Allocation concealment (selection bias).
- Blinding of participants and personnel (performance bias).
- Blinding of outcome assessment (detection bias).
- Incomplete outcome data (attrition bias).
- Selective reporting (reporting bias).
- Other possible bias.

Measures of treatment effect

We calculated odds ratios with 95% confidence intervals for the dichotomous outcomes using standard methods. We used a random effects model as the subjects and interventions would have differed in ways which we anticipated would affect results and we could not assume a common effect size (Borenstein 2009).

We calculated inpatient resource use as the average (mean) use of hospital beds (in days) per patient recruited to each trial group. This figure was calculated for individual trials, and groups of trials, by dividing the total number of bed days by the total number of patients.

Unit of analysis issues

In cross-over trials, we only included data from the first period of the trial in meta-analyses to guard against carry-over effects. Where cluster randomised studies presented an estimate of effect that properly accounted for the cluster design, this was used. Where this was not the case, we assumed that the intra-cluster correlation coefficient (ICC) was the same as for other studies included in the review for that outcome. We calculated an average ICC for the outcome and corrected the values for each unadjusted study by the design effect (Higgins 2011).

Dealing with missing data

Where possible, studies were analysed on an intention-to-treat basis. Patients who were lost to follow up or for whom outcome data were not available were excluded from the initial analysis. However, they were included in 'best case' (all missing data in favour of day hospital care), intermediate and 'worst case' (all missing data in favour of alternative care) sensitivity analyses.

Assessment of heterogeneity

We assessed heterogeneity using I^2 and the Q statistic, with P < 0.1 determining significant heterogeneity (Higgins 2011).

Assessment of reporting biases

We attempted to reduce the risk of reporting bias by undertaking comprehensive searches of multiple databases and trials registers, and contacting authors. Where sufficient studies were included for individual outcomes, we undertook visual inspection of funnel plots to identify any obvious sources of publication bias.

Data synthesis

For patient outcomes, we undertook meta-analyses at the end of follow up for the domains of death, death or institutional care, death or deterioration in ADL, death or poor outcome (institutional care, disability or deterioration) and deterioration in ADL

in survivors. Analyses were based on the published summary data rather than individual patient data. For other patient outcomes - ADL, subjective health status and patient satisfaction - we present a narrative summary and a summary of the data is provided in the Data and analyses section. A summary of carer outcomes is also presented in the Data and analyses section. To investigate resource use, we performed a meta-analysis for the domain of requiring institutional care at the end of follow up. For hospital bed use during follow up and cost we present a narrative summary in the Data and analyses section. We assessed the quality of the evidence using the GRADE approach which results in a quality score of high, moderate, low or very low (GRADEpro 2014).

RESULTS

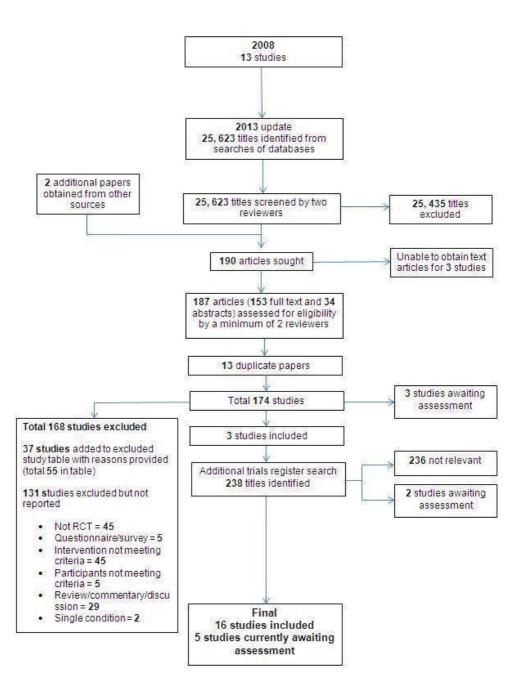
Description of studies

This review update includes 16 randomised controlled trials of medical day hospital versus alternative forms of care for older people. This includes three new studies in addition to the 13 studies from the previous version of this review (Forster 2008).

Results of the search

We screened over 25,000 unique citations and reviewed the full text of 190. 153 full papers and 34 abstracts were obtained and reviewed by a minimum of two reviewers to assess eligibility. Of these, three were included; five are awaiting assessment pending translation or availability of a published report (Studies awaiting classification); the majority of studies were excluded for reasons described in Characteristics of excluded studies; a further 131 studies were excluded for this update but not reported, as they were excluded early in the selection process. A flow diagram of the review update process can be found in Figure 1.

Figure I.



Included studies

Three new studies have been added to this review update (Crotty 2008; Masud 2006; Parker 2009). Five studies are currently awaiting assessment.

Interventions

The current analysis includes 16 trials comprising 37 individual day hospitals. In accordance with the definition of day hospital used, multidisciplinary outpatient rehabilitation was available at all sites. Several of the studies evaluated more than one day hospital; the pilot study undertaken by Vetter 1989 involved two, Masud 2006 and Crotty 2008 each involved three, while a further four trials (Hedrick 1993; Parker 2009; Weissert 1980; Young 1992) each evaluated four day hospitals and Roderick 2001 involved five. The studies were undertaken in various countries including the UK (Burch 1999; Gladman 1993; Masud 2006; Parker 2009; Roderick 2001; Vetter 1989; Woodford 1962; Young 1992), USA (Cummings 1985; Hedrick 1993; Weissert 1980), Australia (Crotty 2008), Canada (Eagle 1991), Hong Kong (Hui 1995), Finland (Pitkala 1991) and New Zealand (Tucker 1984). For further details see Characteristics of included studies.

Comparison groups

Attendance at a day hospital was evaluated against various comparison treatments which were grouped together in the following sub-categories:

- 1) In five studies the comparison treatment was comprehensive care comprising a range of inpatient, outpatient and domiciliary geriatric medical services (Cummings 1985; Eagle 1991; Hedrick 1993; Pitkala 1991; Tucker 1984).
- 2) In seven trials the comparison treatment was domiciliary therapy. This was provided in the patient's home (Crotty 2008; Gladman 1993; Parker 2009; Roderick 2001; Vetter 1989; Young 1992) or day centre (Burch 1999). Three of these trials recruited stroke patients only (Gladman 1993; Roderick 2001; Young 1992) and a fourth was a pilot study (Vetter 1989). In the Nottingham trial patients were randomly allocated to domiciliary rehabilitation or hospital-based rehabilitation in three strata according to discharge ward: health care of older people, general medical unit or stroke unit (Gladman 1993). Hospital-based rehabilitation was provided during day hospital attendance for patients in the older people care stratum and only patients in this stratum have been included in our analysis.
- 3) Four trials compared day hospital attendance against a control group in which patients were eligible for, but not referred to, existing services (Hui 1995; Masud 2006; Weissert 1980; Woodford

1962). In Masud 2006 the control arm received information leaflets on falls prevention and usual care from the primary care service until outcome data was completed, after which time control participants were offered access to the day hospital intervention.

We initially allocated Cummings 1985 and Hui 1995 into their own individual sub-categories according to their comparison group; day hospital versus inpatient care (Cummings 1985) and day hospital versus medical outpatient care (Hui 1995). However, in order to streamline the analysis, these two trials were incorporated into the above categorisation schemes prior to data analysis. The Cummings 1985 trial investigated a day hospital service designed to facilitate early hospital discharge. The service offered to the comparison group was equivalent to comprehensive care and the trial was re-categorised accordingly. The Hong Kong trial (Hui 1995) recruited stroke patients admitted to the same ward and randomised to receive rehabilitation care led by a neurology team or by a geriatrician team. After discharge, patients assigned to a neurologist were followed up at a medical outpatient clinic and the geriatrician patients by day hospital attendance. There were no differences in length of inpatient stay or dependency at discharge and the main treatment difference at final follow up assessment was the type of supporting aftercare: day hospital or medical outpatients. Further discussion with the trialists indicated that this comparison group could best be categorised as 'no comprehensive care'.

Patient characteristics

This review includes studies with a total of 3689 participants. One trial (Hedrick 1993), which was run by the United States Department of Veterans Affairs, recruited largely (96%) male patients. The other trials had a mix of male and female patients. In all but one trial the mean patient age was over 70 years; the New York trial (Cummings 1985) had a mean patient age of 65 years. Four trials (Gladman 1993; Hui 1995; Roderick 2001; Young 1992) recruited only stroke patients. Masud 2006 specifically recruited participants considered at a high risk of falling. The remaining eleven studies recruited patients with a mixture of diagnoses (Burch 1999; Crotty 2008; Cummings 1985; Eagle 1991; Hedrick 1993; Parker 2009; Pitkala 1991; Tucker 1984; Vetter 1989; Weissert 1980; Woodford 1962). The participants usually had a degree of dependency at recruitment as judged by their ADL scores (for further details see Characteristics of included studies).

Excluded studies

The majority of studies were excluded for reasons including a lack of randomisation, intervention that did not meet our criteria for a day hospital, or participants who were not older patients receiving medical care. It should be noted that only those studies which initially appeared to meet the inclusion criteria, but on closer inspection did not, were reported in the Characteristics of excluded studies. For this update, a further 131 studies were excluded but not reported: 45 were not RCTs; in 45 the intervention did not meet our criteria; 29 were review, commentary or discussion papers; five were questionnaires or surveys; and in two the intervention was for a single condition.

Risk of bias in included studies

Ten studies had a low risk of selection bias (method of random se-

quence generation) of which four studies used a computer generated method (Burch 1999; Crotty 2008; Hedrick 1993; Roderick 2001), four used a random number table (Gladman 1993; Hui 1995; Tucker 1984; Woodford 1962) and two used external Internet/web based services (Masud 2006; Parker 2009). Pitkala 1991 had a high risk of bias as randomisation was by date of birth. In five studies, the method of random sequence generation was unreported or unclear (Cummings 1985; Eagle 1991; Vetter 1989; Weissert 1980; Young 1992). For review authors' judgements about each risk of bias item presented as percentages across all included studies see Figure 2, and for review authors' judgements about each risk of bias item for each included study see Figure 3.

Figure 2. Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.

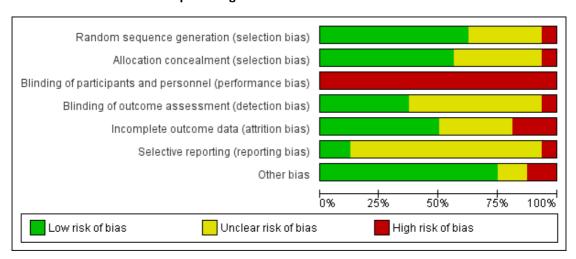


Figure 3. Risk of bias summary: review authors' judgements about each risk of bias item for each included study.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Burch 1999	•	•	•	•	•	?	
Crotty 2008	•	•	•	•	•	•	•
Cummings 1985	?	?	•	?	?	?	?
Eagle 1991	?	?	•	•	•	?	•
Gladman 1993	•	•	•	?	•	?	•
Hedrick 1993	•	•	•	?	•	?	•
Hui 1995	•	•	•	?	?	?	•
Masud 2006	•	•	•	•	•	•	•
Parker 2009	•	•		?	•	•	•
Pitkala 1991				?	?	?	?
Roderick 2001	•	?		•	•	?	•
Tucker 1984	•	?		•	?	?	•
Vetter 1989	?	•	•	?	?	?	•
Weissert 1980	?	?		?		?	•
Woodford 1962	•	?		?		?	
Young 1992	?	•		•	•	?	•

Allocation

Nine studies had an adequate method of allocation concealment (Burch 1999; Crotty 2008; Gladman 1993; Hedrick 1993; Hui 1995; Masud 2006; Parker 2009; Vetter 1989; Young 1992). Methods were unclear in 6 studies (Cummings 1985; Eagle 1991; Roderick 2001; Tucker 1984; Weissert 1980; Woodford 1962). Pitkala 1991 presented with a high risk of bias as their method of randomisation was by date of birth which meant allocation could have been foreseen.

Blinding

Performance bias was a feature of all studies as it was not possible to blind participants due to the nature of the intervention. As a result all studies had a high risk of bias in this domain. Six studies were considered at a low risk for detection bias (blinded outcome assessment; Burch 1999; Crotty 2008; Masud 2006; Roderick 2001; Tucker 1984; Young 1992). Masud 2006 stated that it was not possible to blind researchers to group allocation. However, the review authors considered that the relevant outcome measurements were unlikely to be influenced by a lack of blinding and therefore the risk remained low. The remaining studies were considered to have a high or unclear risk of bias (Cummings 1985; Eagle 1991; Gladman 1993; Hedrick 1993; Hui 1995; Parker 2009; Pitkala 1991; Vetter 1989; Weissert 1980; Woodford 1962).

Incomplete outcome data

Eight studies were considered to be at low risk of bias for attrition (Burch 1999; Crotty 2008; Eagle 1991; Gladman 1993; Hedrick 1993; Masud 2006; Roderick 2001; Young 1992). Three studies were judged to be at high risk of bias. For the Parker 2009 study, losses were similar across the groups but were in excess of 35% by final follow up. For Weissert 1980, 718 participants were excluded for missing data or due to non-adherence. The numbers lost and reasons lost per group were not reported. For Woodford 1962, approximately a third of participants were lost and whilst numbers were balanced across groups, the reasons were not reported. The remaining studies were unclear regarding attrition (Cummings 1985; Hui 1995; Pitkala 1991; Tucker 1984; Vetter 1989).

Selective reporting

Two studies were judged to be at low risk for reporting bias (Masud 2006; Parker 2009). Crotty 2008 was considered at high risk as not all the proposed outcomes reported in the study protocol were included in the available publication. For the remaining studies it was unclear whether selective reporting occurred, or pre-study protocols were unavailable (Burch 1999; Cummings 1985; Eagle 1991; Gladman 1993; Hedrick 1993; Hui 1995; Pitkala 1991; Roderick

2001; Tucker 1984; Vetter 1989; Weissert 1980; Woodford 1962; Young 1992).

Other potential sources of bias

The majority of studies were considered at low risk for other sources of bias (Crotty 2008; Eagle 1991; Gladman 1993; Hedrick 1993; Hui 1995; Masud 2006; Parker 2009; Roderick 2001; Tucker 1984; Vetter 1989; Weissert 1980; Woodford 1962; Young 1992). The Cummings 1985 study was judged as unclear as this was an 'artificial' day hospital established for the purpose of the study and the under-utilisation of the facility may introduced bias. Pitkala 1991 was judged as unclear as 23% of the day hospital group refused the care. Burch 1999 was considered to be at a high risk as 10 of 55 patients transferred from day centre to day hospital.

Further details on how individual studies were scored across the different domains of bias are reported in the risk of bias tables in the Characteristics of included studies.

Effects of interventions

See: Summary of findings for the main comparison Day hospitals compared to alternative care or no care for rehabilitation needs; Summary of findings 2 Day hospitals compared to no comprehensive care for rehabilitation needs; Summary of findings 3 Day hospitals compared to domiciliary care for rehabilitation needs; Summary of findings 4 Day hospitals compared to comprehensive care for elderly persons requiring rehabilitation

The 16 trials included in the review recruited a total of 3689 patients.

Patient outcomes

Nine studies provided final outcome data at 12 months (Burch 1999; Eagle 1991; Gladman 1993; Hedrick 1993; Masud 2006; Parker 2009; Pitkala 1991; Weissert 1980; Woodford 1962), four studies at six months (Crotty 2008; Hui 1995; Roderick 2001; Young 1992), one study at five months (Tucker 1984), one study at three months (Cummings 1985) and one at two months (Vetter 1989).

Death

All 16 trials published data, or provided data on request, for the combined outcome of death at the end of follow up. The pooled OR for all the trials for death at the end of scheduled follow up shows no difference between the day hospital and comparison interventions (odds ratio (OR) 1.05; 95% confidence interval (CI)

0.85 to 1.28; P = 0.66). There was no evidence of a difference when day hospital attendance was compared with comprehensive care (OR 1.26; 95% CI 0.87 to 1.82; P = 0.22), domiciliary care (OR 0.97; 95% CI 0.61 to 1.55; P = 0.89) or no comprehensive care (OR 0.88; 95% CI 0.63 to 1.22; P = 0.43). There was no significant heterogeneity overall (Chi² = 12.04; df = 14; P = 0.60) or for any of the subgroups (P > 0.05) (Analysis 1.1). Outcome data were missing for a total of 102 day hospital patients and 54 controls (representing 3.2% of patients in the comprehensive care subgroup, 0% in the domiciliary subgroup and 7.8% in the no comprehensive care subgroup). Best and worst case sensitivity analyses include the possibility of significant benefit (P < 0.001) or harm (P < 0.01) from day hospital attendance. Visual inspection of funnel plots did not identify any obvious signs of publication bias.

Death or institutional care

Thirteen trials published data, or provided data on request, for death or institutional care by the end of follow up (Burch 1999; Crotty 2008; Eagle 1991; Gladman 1993; Hedrick 1993; Hui 1995; Masud 2006; Pitkala 1991; Tucker 1984; Weissert 1980; Vetter 1989; Woodford 1962; Young 1992). The pooled OR for all the trials for death or institutional care at the end of scheduled follow up shows no difference between the day hospital and comparison interventions. (OR 0.85; 95% CI 0.63 to 1.14; P = 0.28). There was no significant difference between day hospital patients and those receiving comprehensive services (OR 1.00; 95% CI 0.69 to 1.44; P = 0.99), domiciliary care (OR 1.05; 95% CI 0.57 to 1.92; P = 0.88) or no comprehensive services (OR 0.63; 95% CI 0.40 to 1.00; P = 0.05). There were no significant subgroup differences (P = 0.26). There was significant heterogeneity overall for all studies (Chi² = 25.4, df = 11, P = 0.01; I^2 = 57%; Analysis 1.2). Outcome data were missing for a total of 224 day hospital patients and 110 controls (representing 4.2% of patients in the comprehensive care subgroup, 0% in the domiciliary care subgroup and 19.3% in the no comprehensive care subgroup). Best and worst case sensitivity analyses include the possibility of significant benefit (P < 0.0001) or harm (P < 0.0001) from day hospital attendance. Visual inspection of funnel plots did not identify any obvious signs of publication bias.

Death or deterioration in ADL

Seven trials published data on death or deterioration in ADL (Burch 1999; Gladman 1993; Hui 1995; Pitkala 1991; Vetter 1989; Weissert 1980; Young 1992). The pooled OR for all the trials at the end of scheduled follow up shows no difference between the day hospital and comparison interventions (OR 1.07; 95% CI 0.76 to 1.49; P = 0.70). Only Pitkala 1991 provided data for day hospital compared to comprehensive care and the difference was not significant (OR 1.18; 95% CI 0.63 to 2.18, P = 0.61).

There was no difference between day hospital and domiciliary care (OR 1.41; 95% CI 0.82 to 2.42; P = 0.21) or no comprehensive care (OR 0.76; 95% CI 0.56 to 1.05; P = 0.09). There were no significant subgroup differences (P = 0.11) and no significant heterogeneity overall (Chi² = 10.25, df = 6, P = 0.11; $I^2 = 41\%$; Analysis 1.3). Visual inspection of funnel plots did not identify any obvious signs of publication bias.

Death or poor outcome

Thirteen trials published data on death or poor outcome (Burch 1999; Cummings 1985; Eagle 1991; Gladman 1993; Hedrick 1993; Hui 1995; Pitkala 1991; Roderick 2001; Tucker 1984; Vetter 1989; Weissert 1980; Woodford 1962; Young 1992). Roderick 2001 reported data on poor outcome which they defined as "death, recurrent stroke and a six month Barthel score of < 14"; we determined that this was sufficiently similar to our own definition to include in the results. The pooled OR for all the trials at the end of scheduled follow up shows no significant difference between the day hospital and other interventions (OR 0.92; 95% CI 0.74 to 1.15; P = 0.49). There was no significant difference when day hospital was compared with comprehensive care (OR 1.05; 95% CI 0.79 to 1.40; P = 0.74) or domiciliary care (OR 1.08; 95% CI 0.67 to 1.74; P = 0.75). However, there was a significant difference in favour of the day hospital when compared with no comprehensive care (OR 0.72; 95% CI 0.53 to 0.99; P = 0.04), although subgroup results were not significantly different from each other (P = 0.17). There was no significant heterogeneity overall (Chi² = 17.27, df = 12, P = 0.14; I^2 = 31%; Analysis 1.4). Outcome data were missing for 55 day hospital patients and 121 controls (representing 4.6% of patients in the comprehensive care subgroup, 0.5% in the domiciliary care subgroup and 10.3% in the no comprehensive care subgroup). Best and worst case sensitivity analyses included the possibility of significant benefit (P < 0.0001) or harm (P < 0.05) from day hospital attendance. Visual inspection of funnel plots did not identify any obvious signs of publication bias.

Deterioration in ADL among survivors

We wished to examine the influence of day hospital attendance on the functional status of survivors. Although most trials described results in terms of ADL scores, seven different measures were used and reported in different ways. We therefore describe results in terms of recorded deterioration in ADL and the raw ADL results. Seven trials provided data on deterioration in ADL among survivors (Burch 1999; Gladman 1993; Hui 1995; Pitkala 1991; Vetter 1989; Weissert 1980; Young 1992). We judged the quality of the evidence for the following outcome as low (Summary of findings for the main comparison). Overall there was no difference between day hospital and alternative care in ADL scores (OR 1.11; 95% CI 0.68 to 1.80; P = 0.67). However, day hospital attenders

appeared less likely to deteriorate than those receiving no comprehensive care (OR 0.61; 95% CI 0.38 to 0.97; P = 0.04). Differences were not significant when comparing day hospitals with comprehensive care (OR 1.21; 95% CI 0.58 to 2.52; P = 0.61) or domiciliary care (OR 1.59; 95% CI 0.87 to 2.90; P = 0.13). There were significant subgroup differences (P = 0.04) and evidence of heterogeneity (Chi² = 11.94, df = 6, P = 0.06; I^2 = 50%; Analysis 1.5). Visual inspection of funnel plots did not identify any obvious signs of publication bias.

ADL score

Fourteen trials reported a standardised measure of ADL among survivors. However, various measures were used and data were insufficient to allow a statistical summary of the results. Two trials demonstrated significant but small improvements in functional ability with day hospital attendance which was not sustained at six month follow up (Hui 1995; Tucker 1984). One trial (Young 1992) reported an improved functional outcome for the comparison group. The other 11 trials (Burch 1999; Cummings 1985; Eagle 1991; Gladman 1993; Hedrick 1993; Masud 2006; Parker 2009; Pitkala 1991; Roderick 2001; Vetter 1989; Weissert 1980) found no difference in disability scores between the day hospital and comparison groups (Analysis 1.6).

Subjective health status

A number of studies investigated subjective health status. However, various measures were used and we were unable to incorporate data into a meta-analysis. Three studies investigating day hospital versus comprehensive care found no significant difference between the groups (Cummings 1985; Eagle 1991; Hedrick 1993). In Tucker 1984 there was a significant improvement in mood measured by the Zung index in the day hospital group compared to the comprehensive care group at final follow up (P = 0.01). Pitkala 1991 provided no comparable data. There were no significant differences in any of the studies investigating day hospital versus domiciliary care (Burch 1999; Gladman 1993; Parker 2009; Roderick 2001; Vetter 1989; Young 1992). For day hospital versus no comprehensive care, Hui 1995 found no significant differences. Weissert 1980 and Woodford 1962 did not provide comparable data (Analysis 1.7).

Patient satisfaction

Data on patient satisfaction were only available from one study. Hui 1995 found no significant difference between the day hospital and no comprehensive care (Analysis 1.8).

Carer outcomes

Distress

There were no available or comparable data for day hospital versus comprehensive care or day hospital versus no comprehensive care. Data were available from three studies comparing day hospital with domiciliary care. Crotty 2008 and Gladman 1993 found no significant difference at follow up; Burch 1999 found a significant difference in the mean change between baseline and three months in the Caregiver Strain Index in both groups but no significant difference between groups. (Analysis 1.9).

Resource use

Requiring Institutional care at the end of follow up

Thirteen trials provided information about the number of patients requiring institutional care at the end of follow up (Burch 1999; Crotty 2008; Eagle 1991; Gladman 1993; Hedrick 1993; Hui 1995; Masud 2006; Pitkala 1991; Tucker 1984; Vetter 1989; Weissert 1980; Woodford 1962; Young 1992). In one trial (Weissert 1980) these data were available only for a subgroup of patients (384 patients of 552 recruited to the main study). There was no difference between day hospital and all other services (OR 0.84; 95% CI 0.58 to 1.21; P = 0.35), or for any of the subgroups: day hospital versus comprehensive care (OR 0.91; 95% CI 0.70 to 1.19; P = 0.49), day hospital versus domiciliary care (OR 1.49; 95% CI 0.53 to 4.25; P = 0.45) or day hospital versus no comprehensive care (OR 0.58; 95% CI 0.28 to 1.20; P = 0.14). Overall there was significant heterogeneity (Chi² = 20.03, df = 11, P = 0.04; $I^2 = 45\%$; Analysis 2.1). On the basis of these data (95% CI 10 to 34) 21 patients (95% CI 12.3 to 70.9) would need to attend day hospital (as opposed to receiving no comprehensive service) to prevent one admission to long term institutional care. Visual inspection of funnel plots did not identify any obvious signs of publication bias.

Hospital bed use

Although hospital use was described in several ways in the trials, it proved possible to obtain a standardised measure for 14 trials of average (mean) hospital bed use per patient recruited (Burch 1999; Cummings 1985; Eagle 1991; Gladman 1993; Hedrick 1993; Hui 1995; Masud 2006; Pitkala 1991; Roderick 2001; Tucker 1984; Vetter 1989; Weissert 1980; Woodford 1962; Young 1992; Analysis 2.2). A measure of variance was not possible for this analysis and therefore confidence limits cannot be reported. The results show a small reduction in bed use by the day hospital patients compared to other treatment across all trials: 13.6 versus 14.6 (Analysis 2.2), with subgroup results as follows: Day hospital versus comprehensive care - 20.5 versus 21.5. Day hospital versus domiciliary care - 6.8 versus 9.2.

Data from Parker 2009 was in a format that did not allow us to incorporate it into the above analysis. However, they reported the mean total length of stay in hospital for patients which was higher in the day hospital group compared to the home rehabilitation group (mean difference 9.3 days; 95% CI 12.5 to 31.1; P > 0.05).

Costs

A number of studies reported a comparison of treatment costs (Analysis 2.3), but methods for reporting data were not consistent and therefore the data could not be incorporated into a meta-analysis. Seven studies reported that day hospital attendance was more expensive than the comparison treatment (Burch 1999; Gladman 1993; Hedrick 1993; Masud 2006; Tucker 1984; Weissert 1980; Young 1992). Three trials reported that the costs were similar (Hui 1995; Cummings 1985; Roderick 2001). Woodford 1962 reported that day hospital attendance was considerably less expensive than inpatient care (8% of weekly inpatient costs) but made no comparison of other costs incurred specifically by the comparison group. Parker 2009 reported that there was insufficient evidence to support the hypothesis that rehabilitation is less expensive in a home based setting.

For the sub category comparing mean treatment costs between day hospital care and other comprehensive care services, Cummings 1985, Hedrick 1993 and Tucker 1984 reported higher costs for the day hospital. The Cummings 1985 and Hedrick 1993 trials included the cost of nursing home care. There was no information from two trials for this comparison (Eagle 1991; Pitkala 1991). For

the sub category comparing treatment costs between day hospital care and domiciliary care the Burch 1999; Gladman 1993 and Young 1992 trials found that day hospital was more expensive. In the Roderick 2001 trial, the day hospital was more expensive for rehabilitations costs but was less so when considering total health and social services costs. There was no information from three trials for this comparison (Crotty 2008; Parker 2009; Vetter 1989). For the sub category comparing treatment costs between day hospital care and no comprehensive care, day hospital was more expensive than no comprehensive care in Hui 1995, Masud 2006 and Weissert 1980. No formal costing data were provided by Woodford 1962.

Assessments of the quality of the body of evidence

Using the GRADE approach we judged the quality of the body of evidence to be low for the patient outcomes of death, death or institutional care, death or deterioration in ADL, death or poor outcome, and deterioration in ADL (Summary of findings for the main comparison; Summary of findings 2; Summary of findings 3; Summary of findings 4; Appendix 20). For each outcome the body of evidence was from randomised controlled trials but we reduced the quality rating because of a high likelihood of bias in the included studies and imprecision in the effect estimates (wide CIs). We did not reduce the quality rating despite evidence of heterogeneity because this was anticipated due to the diversity of the populations and of the interventions.

ADDITIONAL SUMMARY OF FINDINGS [Explanation]

Day hospitals compared to no comprehensive care for rehabilitation needs

Patient or population: patients with rehabilitation needs

Intervention: day hospitals

Comparison: no comprehensive care

Outcomes	• • • • • • • • • • • • • • • • • • • •		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence Comments (GRADE)
	Assumed risk	Corresponding risk			
	No comprehensive care	Day hospitals			
Death by the end of fol-	Study population		OR 0.88	1345	⊕⊕⊖⊝ Lauri 2.3
low up Follow-up: median 12 months	128 per 1000	114 per 1000 (85 to 152)	(0.63 to 1.22)	(4 studies)	low ^{1,2,3}
	Moderate				
	131 per 1000	117 per 1000 (87 to 155)			
Death or institutional			OR 0.63	1177	000
care by the end of fol- low up Follow-up: median 12 months	248 per 1000	172 per 1000 (117 to 248)	(0.4 to 1)	(4 studies)	low ^{1,2,3}
	Moderate				
	307 per 1000	218 per 1000 (151 to 307)			

Death or deterioration	, . ·		OR 0.76	651 (0 atvalias)	⊕⊕⊖⊖ Lauri 2.3
in ADL Follow-up: median 9 months	436 per 1000	370 per 1000 (302 to 448)	(0.56 to 1.05)	(2 studies)	low ^{1,2,3}
	Moderate				
	446 per 1000	380 per 1000 (311 to 458)			
Death or poor outcome	, , ,		OR 0.72	982	⊕⊕○○
(institutional care, dis- ability or deterioration) Follow-up: median 12 months	347 per 1000	277 per 1000 (220 to 345)	(0.53 to 0.99)	(3 studies)	low ^{1,2,3}
	Moderate				
	400 per 1000	324 per 1000 (261 to 398)			
Deterioration in ADL in	Study population		OR 0.61	407	⊕⊕⊖⊝ low ^{1,2,3}
survivors Follow-up: median 9 months	277 per 1000	189 per 1000 (127 to 271)	(0.38 to 0.97)	(2 studies)	
	Moderate				
	227 per 1000	152 per 1000 (100 to 222)			

^{*}The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; OR: Odds ratio; ADL: activities of daily living

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Limitations for at least one risk of bias criterion or some limitations for multiple criteria, sufficient to lower confidence in the estimate of effect

 $^{^{2}}$ Whilst there was evidence of heterogeneity, this was anticipated due to the diversity of the population and of the study design

³ Wide Cls

Day hospitals compared to domiciliary care for rehabilitation

Patient or population: patients with rehabilitation needs Intervention: day hospitals
Comparison: domiciliary care

Outcomes			Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence Comments (GRADE)
	Assumed risk	Corresponding risk	_		
	Domiciliary care	Day hospitals			
Death by the end of fol-	Study population		OR 0.97	901 (7 studies)	⊕⊕⊜⊝ low ^{1,2,3}
low up Follow-up: median 6	101 per 1000	98 per 1000 (64 to 148)	(0.61 to 1.55)	(7 studies)	IOW · · · · ·
	Moderate				
	64 per 1000	62 per 1000 (40 to 96)			
Death or institutional	Study population		OR 1.05	672	⊕⊕⊖⊖ L1 2 3
care by the end of fol- low up Follow-up: median 6	187 per 1000	194 per 1000 (116 to 306)	(0.57 to 1.92)	(5 studies)	low ^{1,2,3}
months	Moderate				
	69 per 1000	72 per 1000 (41 to 125)			
Death or deterioration in ADL Follow-up: median 9 months	Study population		OR 1.41 (0.82 to 2.42)	443 (4 studies)	⊕⊕⊖⊝ low ^{1,2,3}

	392 per 1000	476 per 1000 (346 to 609)			
	Moderate				
	334 per 1000	414 per 1000 (291 to 548)			
Death or poor outcome (institutional care, dis- ability or deterioration) Follow-up: median 6 months			OR 1.08	581	⊕⊕○○
	297 per 1000	313 per 1000 (221 to 424)	(0.67 to 1.74)	(5 studies)	low ^{1,2,3}
	Moderate				
	364 per 1000	382 per 1000 (277 to 499)			
Deterioration in ADL in	Study population		OR 1.59	349	⊕⊕○○
survivors Follow-up: median 9 months	225 per 1000	315 per 1000 (201 to 457)	(0.87 to 2.9)	(4 studies)	low ^{1,2,3}
	Moderate				
	188 per 1000	269 per 1000 (168 to 402)			

^{*}The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

CI: Confidence interval; OR: Odds ratio; ADL: activities of daily living

- ¹ Limitations for at least one risk of bias criterion or some limitations for multiple criteria, sufficient to lower confidence in the estimate of effect
- ² Whilst there was evidence of heterogeneity, this was anticipated due to the diversity of the population and the interventions
- ³ Wide Cls

Day hospitals compared to comprehensive care for older people requiring rehabilitation

Patient or population: older people requiring rehabilitation Intervention: day hospitals

Comparison: comprehensive care

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence Comments (GRADE)
	Assumed risk	Corresponding risk			
	Comprehensive care	Day hospitals			
Death by the end of fol-	Study population		OR 1.26	1287	⊕⊕○○ Lauri 2 3
low up Follow-up: median 12 months	144 per 1000	175 per 1000 (128 to 234)	(0.87 to 1.82)	(5 studies)	low ^{1,2,3}
	Moderate				
	69 per 1000	85 per 1000 (61 to 119)			
Death or institutional	Study population		OR 1	1181	⊕⊕○○ L 133
care by the end of fol- low up Follow-up: median 12	426 per 1000	426 per 1000 (339 to 517)	(0.69 to 1.44)	(4 studies)	low ^{1,2,3}
months	Moderate				
	231 per 1000	231 per 1000 (172 to 302)			
Death or deterioration in ADL Follow-up: median 12 months	Study population		OR 1.18 (0.63 to 2.18)	174 (1 study)	$\oplus \oplus \bigcirc \bigcirc$ low 1,3

			_		
	349 per 1000	387 per 1000 (252 to 539)			
	Moderate				
	349 per 1000	387 per 1000 (252 to 539)			
Death or poor outcome (institutional care, dis- ability or deteriora- tion) Follow-up: median 12 months	7 1 1		OR 1.05	1268	000
		422 per 1000 (355 to 493)	(0.79 to 1.4)	(5 studies)	low ^{1,2,3}
	Moderate				
	221 per 1000	230 per 1000 (183 to 284)			
Deterioration in ADL in	Study population		OR 1.21	149	$\Theta\ThetaOO$
survivors Follow-up: median 12 months	243 per 1000	280 per 1000 (157 to 448)	(0.58 to 2.52)	(1 study)	$low^{1,3}$
	Moderate				
	243 per 1000	280 per 1000 (157 to 447)			

^{*}The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

CI: Confidence interval; OR: Odds ratio; ADL: activities of daily living

- Limitations for at least one risk of bias criterion or some limitations for multiple criteria, sufficient to lower confidence in the estimate of effect
 Whilst there was evidence of heterogeneity, this was anticipated due to the diversity of the population and of the study
- design
- ³ Wide Cls

DISCUSSION

Summary of main results

The majority of included studies have compared day hospital care with other services. Only four trials employed a comparison group of patients who received neither comprehensive care nor domiciliary rehabilitation (Masud 2006; Woodford 1962; Hui 1995; Weissert 1980). The results from this group were the most favourable to day hospital care, but these four trials are now quite old. Overall the quality of the evidence was low, therefore further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate. For the outcome of death, there was no difference between day hospitals and other services, including when day hospitals were compared with any of the subcategories individually. For the combined outcome of death or institutional care there was no significant difference between the day hospital and all other services. For the combined outcome of death or deterioration in ADL, there was no significant difference between the day hospital and other services although there was a trend in favour of the day hospital compared with no comprehensive care. For the combined outcome of death or poor outcome, there was a significant difference in favour of the day hospital when compared with no comprehensive services. For the outcome of deterioration in ADL, there was a significant difference between attending day hospital and no comprehensive care. However, there was no difference between the day hospital and other services.

When considering resource implications among those requiring institutional care, there was no difference at the end of follow up between day hospital and other services. There was a slight reduction in hospital bed use overall for day hospital patients and particularly when day hospitals were compared with domiciliary services; however, whilst a summary statistic was not possible for hospital bed use or cost, individual studies suggest that day hospitals are predominantly as expensive or more expensive than others services.

Overall completeness and applicability of evidence

The day hospital trials included in this systematic review have predominantly employed a pragmatic design and have attempted to address broad questions of overall day hospital effectiveness. This review included 16 studies with 3689 participants, although within the analyses of specific outcomes these numbers were reduced as each study only contributed data for some comparisons. It is unfortunate that data was not available, or in a suitable format, to undertake statistical analyses for the patient outcomes of activities of daily living, subjective health status, patient satisfaction or carer distress, or for the resource outcome of cost. It also

proved impossible to determine a summary statistic for disability because, although included as an outcome measure in 11 trials, different measurement instruments were used and variance data were not available. The outcome of death was reported by all studies, however other adverse events and effects were not reported consistently by all studies, thus compromising the overall completeness of findings.

We have based the systematic review on a broad comparison of day hospital care versus alternative services. We wanted to be able to generalise to a range of scenarios and not defined populations. As we anticipated considerable variations in the comparison services, these were identified and categorised prior to data collection and analysis. We have ensured that the treatment schedules described matched our pre-determined definition of day hospital care. Thus, whilst the study by Weissert and colleagues (Weissert 1980) refers to 'day care' services, the intervention provided fitted our definition of day hospital care and was therefore included. A lack of consensus in terminology related to day hospitals has also been noted elsewhere (Petermans 2011).

The applicability of the findings from this review to various regions will depend on current health care provision and populations. As comprehensive care, in one form or another, is likely to be available to many older patients who require rehabilitation, the relevance of some comparisons may be limited in certain countries and populations; specifically the findings from the comparison between day hospitals and no comprehensive care (which was favourable for the day hospital on the combined measure of death or poor outcome, with a trend in favour of the day hospital for the combined measure of death or institutional care). This review found little evidence that day hospitals were better than alternative types of comprehensive service. However, the diversity in the content of alternative services and the populations being served (studies originated from seven different counties) means the external validity of this finding may be compromised. Furthermore, 10 of the studies were at least 20 years old and the types of health service and the populations being served may not reflect current practice or requirements. Services may need to be considered on a case by case basis regarding their applicability against current health provision.

Two main limitations of our review lie in the multinational settings of the studies and in the forty year time span of study publication, during which time health and social care policies inevitably changed. It is disappointing that there have been only four further evaluations of the effectiveness of day hospital care since this review was first published (Forster 1999a). Nonetheless, the data presented here probably represent the best evidence currently available upon which to base a judgment of day hospital effectiveness.

Quality of the evidence

There are limitations due to the lack of statistical power resulting from small, heterogeneous trials. As a result there is a danger

of both false-positive and false-negative results being generated. Furthermore, the data was from a series of studies performed by researchers operating independently and the studies were therefore not functionally equivalent. This resulted in significant heterogeneity through variations in participants and in the interventions employed in the various day hospitals and comparison treatments. Participants and interventions likely differed in ways that impacted the results. Consequently, we cannot assume a common effect size (Borenstein 2009). A frequently encountered issue with pragmatic rehabilitation trials is that the methods to record subject characteristics, which might influence prognosis and treatment processes, are poorly developed. Considerable detail was recorded in the USA (Hedrick 1993) trial; it is important that any future trials also address this issue. Data were missing for a number of outcomes which could theoretically alter the scale and direction of the results. All of our conclusions must be qualified by this con-

It is possible that biases have resulted in the overestimation or underestimation of the effects of the intervention. All studies were at a high risk of bias for blinding of participants and personnel; this is typical of such an intervention and effective blinding would be challenging to implement in view of ethical considerations which require participants to have prior knowledge of potential interventions. However, a lack of blinding of outcome assessment was also problematic and only half the studies were considered to have a low risk in this domain. Contamination can result in the underestimation of treatment effects and whilst not all studies were judged to be of high risk, contamination and deviation from the protocol (switching between interventions) did feature in some studies comparing day hospitals with other comprehensive services and domiciliary care. In addition, the risk of selective reporting was often unclear. Only the three studies which were reported in the last eight years had available protocols against which their reports could be assessed to determine if all outcomes had been reported. Finally, half of the studies were considered to have a low risk of bias for the domain of incomplete outcome data. This should be considered positive, taking into account the nature of the intervention and participants, post intervention follow up periods, and the age of the studies where attrition was less frequently reported.

Potential biases in the review process

The search strategy was extensive, taking into account multiple databases and sources, and this is reflected in the large number of titles identified. Two authors extracted all data, discussions were undertaken to achieve consensus, and a third person was utilised where disagreements remained. As a result, we are confident in the quality of the review. It was unfortunate, however, that we could not quantitatively combine data for a number of outcomes, due to the diversity of measures used (for example, ADL, subjective health status, patient satisfaction, carer distress, bed use and resource use).

Publication bias (Egger 1997) remains a possibility in any review process. However, our search strategy was extensive and included contacting the authors of papers relating to day hospital care around the world.

We were fortunate that many of the authors of the published papers or abstracts were able to provide additional information which has not been published previously.

Agreements and disagreements with other studies or reviews

Another systematic review drew similar conclusions to our review (Petermans 2011). They found that geriatric day hospitals were better for patients undergoing assessment and intervention than no comprehensive care. However, they found little benefit when compared to treatment in a geriatric ward or other geriatric services. They also found few studies reporting on the outcome of patient satisfaction. The authors did not include meta-analysis and they did not report on any cost-benefit outcomes; their conclusions were drawn from various sources including RCTs and cohort studies. Another systematic review, with meta-analysis, found that comprehensive geriatric assessments linking geriatric evaluation with long term management are effective in improving survival and function in older people (Stuck 1993).

Individual results from the included studies suggest that day hospitals are probably as expensive or more expensive than other comprehensive or domiciliary services. Several costing studies have drawn attention to the expense of day hospital services (Eagle 1987; Gerard 1988; Gladman 1994; MacFarlane 1979; Young 1993).

AUTHORS' CONCLUSIONS

Implications for practice

Any conclusions are limited by the relatively small amount of data available and the low quality of the data contributing to a number of important outcomes. Day hospital care appears to be an effective outpatient service for older people, but no more effective (at least for the outcomes examined in this review), and possibly more expensive, than other forms of comprehensive care. These findings do not support the closure of day hospital services but do support the exploration of alternative systems for delivering an equivalent or superior form of comprehensive care. Our findings support the view that day hospital attendance needs to be carefully monitored (George 1989) and the staff and facilities used as flexibly and efficiently as possible (Brocklehurst 1995).

Implications for research

The findings are limited by the relatively small amount of data

available and overall low quality of the evidence; further research is likely to impact on our confidence in the estimate of effect. Further randomised trials are justifiable and should focus on comparing services which aim to provide an equivalent intervention to day hospital care (e.g. domiciliary care). Given the diversity of patients attending day hospitals and the corresponding diversity of day hospital interventions employed, future trials should be large, multicentre trials or should examine more focused questions. Outcomes should include subjective health status and carer well being. There is concern that commonly used measures of disability lack sensitivity to change in the outpatient setting of a day hospital due to their ceiling effect (Parker 1994). Future trials should incorporate measures of instrumental activities of daily living as a more relevant and potentially more sensitive outcome. Furthermore, future trials should incorporate adequate methods of randomisation and allocation concealment, and undertake blinded outcome assessment where possible, as well as ensuring that methods are adequately reported.

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^{*} Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies [ordered by study ID]

Burch 1999

Methods	Randomised controlled trial Method of randomisation: computer generated Concealment of allocation: sealed envelopes Outcome assessor blinding: blinded research sociologist	
Participants	Country: UK Patients referred to day hospital Exclusion criteria: dysphasic, required nursing or occupational therapy > twice per week 163 patients eligible (28 needed day hospital treatment, 21 refused consent, 9 operational problems at day centre) Participants randomised = 105 Baseline function: Median (IQR) Barthel Index 15 (12-17) and 15 (11-17) Male: 36% Age: mean (SD) 80.4 (7.6) years	
Interventions	Day hospital: care by multidisciplinary rehabilitation team, principally nursing assessment, occupational therapy and physiotherapy. Median number of treatments (interquartile range) 11.5 (5-20.5) Day centre: rehabilitation provided by a physiotherapist and two support workers. Median number of treatments (interquartile range) 10 (5-14)	
Outcomes	12 month follow up Death Institutional care Barthel Index Caregiver Strain Index Philadelphia Geriatric Morale scale Costs	
Notes	Total of 105 patients of whom 23 had a stroke diagnosis, 14 osteoarthritis, 13 fracture, 9 Parkinsonism Of the 55 patients randomised to day centre attendance, 10 transferred to day hospital	
Risk of bias		
Bias	Authors' judgement Support for judgement	
Random sequence generation (selection bias)	Low risk	Computer generated
Allocation concealment (selection bias)	Low risk	Quote: "Immediately after consent, sub- jects were randomly allocated to day hos- pital or day centre by a sequence of la- belled tickets in sealed, opaque envelopes

Burch 1999 (Continued)

		securely kept and opened by a senior ward clerk unattached to the trial teamcomputer generated blocks of 20."
Blinding of participants and personnel (performance bias) All outcomes	High risk	Reported as single blind which appears to have been the assessor, not participants or personnel
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Outcome assessments were undertaken by a blinded research sociologist Quote: "The interviewer correctly identified 38/55 as day hospital and 20/38 as day centre, yielding kappa = 0.22 indicating poor agreement/successful blinding."
Incomplete outcome data (attrition bias) All outcomes	Low risk	Similar losses per group (~30%), moderately high but similar reasons reported for both groups
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias
Other bias	High risk	18% of participants randomised to day centre attendance transferred to the day hospital leading to possibility of contamination (the experiment and control groups becoming mixed)

Crotty 2008

Methods	Randomised controlled trial Method of randomisation: computer generated Concealment of allocation: sealed envelopes Outcome assessor blinding: blinded research occupational therapist
Participants	Country: Australia Hospitalised patients referred for ambulatory rehabilitation Inclusion criteria; medically stable; ready for hospital discharge; rehabilitation which required at least 12 therapy sessions Exclusion criteria: lived out of the health region; if referring clinician felt they were unsuitable to receive one of the programmes 301 patients assessed for study inclusion (34 patients did not meet the eligibility criteria, 38 patients declined to participate or were not approached on the request of the physician) Participants randomised = 229 Modified Bartel Index mean (SD): 92.4 (6.5) Mini-Mental State Examination mean (SD): 26.9 (3.1) Male: 48%

Crotty 2008 (Continued)

	Age: mean (SD) 71.7 (14.1) years
Interventions	Day hospital: Interdisciplinary programme providing 4-6 weeks of high intensity rehabilitation in either individual or group sessions with the option of extending the programme. Each visit lasted 3 hours. Participants had access to physiotherapy, occupational therapy, social work, psychology, dietetics and nursing and rehabilitation medicine Home based rehabilitation: One to one rehabilitation programme delivered by an interdisciplinary team to participants in their homes. This included physiotherapy, occupational therapy, speech therapy, social work, dietetics, nursing and rehabilitation medicine. Three to five session per week usually delivered for between 4 and 6 weeks
Outcomes	Primary outcome: Assessment of Motor and Process Skills, bioelectrical impedance Secondary outcomes: depression, Mini Nutritional Assessment, Assessment of Appetite, Mini Mental State Exam, Timed Up and Go, and Short Form 36 (patient and carer), patient satisfaction and carer/family satisfaction, Carer Strain Index, mortality and place of residence, cost and readmissions. Outcomes assessed at baseline, discharge, three and six months
Notes	

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer generated
Allocation concealment (selection bias)	Low risk	Quote: "statistician external to the study generated the randomisation sequence us- ing the random number generator in Mi- crosoft Excel and created sequentially num- bered, opaque, sealed envelopes containing group allocation for participants"
Blinding of participants and personnel (performance bias) All outcomes	High risk	The same doctor provided medical services to both groups. Furthermore, participants could not have been blinded to the intervention
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Assesments were undertaken by a research occupational therapist blinded to the group allocation
Incomplete outcome data (attrition bias) All outcomes	Low risk	Missing outcome data balanced across groups and with similar reasons reported
Selective reporting (reporting bias)	High risk	Some outcome measures reported in study protocol (Australian New Zealand Clini- cal Trials Registry) not reported in current

Crotty 2008 (Continued)

		100	
		publications	
Other bias	Low risk	No other obvious sources of bias	
Cummings 1985			
Methods	Concealment of allocation: not	Randomised controlled trial Method of randomisation: not reported Concealment of allocation: not reported Outcome assessor blinding: unclear for some outcomes	
Participants	with someone, fit to travel, 24 eligible	rears, disabled (not spinal injuries or head injuries), living 4 hour telephone contact, suitable residence, medicare t/carers refused consent, 452 rejected from study sample)	
Interventions	· -	Day hospital attendance 5 days a week with emphasis on rehabilitation with greater patient and carer involvement. Complete range of medical and therapeutic services available.	
	Rehabilitation as an inpatient.		
Outcomes	3 month follow up Death Institutional care ADL: i) modified Kenny, ii) sul Instrumental ADL Checklists to measure indoor as Medical status Mental state Psychological well-being (Kahn Patient satisfaction Family impact questionnaire Costs	nd outdoor leisure activity	
Notes	96 patients were recruited, of w Day hospital tested as an altern	whom 55 had a stroke diagnosis and 26 were amputees native to inpatient care	
Risk of bias			

Authors' judgement

Bias

Support for judgement

Cummings 1985 (Continued)

Random sequence generation (selection bias)	Unclear risk	No information reported
Allocation concealment (selection bias)	Unclear risk	No information reported
Blinding of participants and personnel (performance bias) All outcomes	High risk	No information reported but would have been obvious to participants
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	It was reported that medical status was assessed by a physician who did not know the patient. However, it was unclear if this was the case for other outcomes
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Not all outcomes were reported in the publication (Cummings 1985). However, some additional information was provided by the authors on request
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias
Other bias	Unclear risk	This was an 'artificial' day hospital designed specifically for the purpose of the study and may have been affected by some environmental factors and under-utilisation of the hospital

Eagle 1991

Methods	Randomised controlled trial, stratified by conventional service Method of randomisation: not reported Concealment of allocation: not reported Outcome assessor blinding: not undertaken
Participants	Country: Canada Patients referred from the community to 2 geriatricians or about to be discharged from hospital Inclusion criteria: age over 65 years, reduced function with rehabilitation potential 128 patients asked to participate (15 refused) Participants randomised = 113 Baseline function: Geriatric Quality of Life Questionnaire (ADL) 4.49 and 4.46 Male: 40% Age: mean (SD) 78.9 (7.2) years

Eagle 1991 (Continued)

Interventions	Day hospital: Attendance 2 days a week. Treatment included multidisciplinary team assessment, programme of rehabilitation provided by physiotherapists and occupational therapists Usual care: Management in inpatient geriatric assessment unit for comprehensive assessment and treatment, management in the outpatient geriatric clinic, with limited diagnostic and rehabilitative opportunities, or early discharge from a medical-surgical inpatient unit and appropriate community follow-up services The same professionals provided treatment to both groups
Outcomes	12 month follow up Death Institutional care Mental status Geriatric Quality of Life Questionnaire Barthel Index Rand questionnaire Global Health Question (GHQ) Family rating of Barthel Index, GHQ, Rand Questionnaire Patient rating of Barthel Index Resource use
Notes	113 patients were recruited, of whom 26 had a stroke diagnosis, 32 a diagnosis of depression and 19 a diagnosis of degenerative joint disease Patients were stratified according to the type of conventional care specified by the participating geriatrician prior to randomisation

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	No information reported
Allocation concealment (selection bias)	Unclear risk	No information reported
Blinding of participants and personnel (performance bias) All outcomes	High risk	Quote: "We were unable to blind the patients, caregivers and study personnel administering the questionnaires and instruments for measuring functional status to the study groups"
Blinding of outcome assessment (detection bias) All outcomes	High risk	Quote: "We were unable to blind the patients, caregivers and study personnel administering the questionnaires and instruments for measuring functional status to the study groups"

Eagle 1991 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Low risk	Missing outcome data fairly balanced in numbers across groups
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias
Other bias	Low risk	No other obvious sources of bias

Gladman 1993

Methods	Randomised controlled trial Method of randomisation: random number table Concealment of allocation: sealed envelopes Outcome assessor blinding: blinded assessment at 6 months and 1 year
Participants	Country: UK Patients discharged home from hospital after acute stroke Exclusion criteria: discharged to residential or nursing homes, requiring respite or terminal care, receiving outpatient rehabilitation prior to the stroke, no significant disability, in hospital < 7 days Patients discharged from older care, general medical wards and stroke unit were randomised separately Participants = 155 Baseline function: Median Barthel Index (IQR) 17 (14-17) and 16 (13-17) Male: 48% Age: mean 70 years
Interventions	Domiciliary rehabilitation intervention: 2 half time physiotherapists, 1 occupational therapist and treatment for up to 6 months (75% received treatment) Day hospital intervention: multidisciplinary rehabilitation provided (54% received treatment)
Outcomes	12 month follow up Death Institutional care Extended ADL score Barthel Index Nottingham Health Profile Brief Assessment of Social Engagement Life Satisfaction Index (Nottingham version) Costs
Notes	All stroke patients (155)

Gladman 1993 (Continued)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "consecutive sealed envelopes which contained cards marked either "DRS" or "HRS" which had been prepared before the start of the study by reference to a table of random numbers."
Allocation concealment (selection bias)	Low risk	Quote: "consecutive sealed envelopes which contained cards marked either "DRS" or "HRS which had been prepared before the start of the study by reference to a table of random numbers."
Blinding of participants and personnel (performance bias) All outcomes	High risk	No information reported. However, participants would have been aware of allocation
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Blinded assessment at 6 months and 1 year, however unclear if baseline data were collected by a blinded assessor
Incomplete outcome data (attrition bias) All outcomes	Low risk	Some imbalance in missing outcome data but losses relatively low in both groups
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias
Other bias	Low risk	No other obvious sources of bias

Hedrick 1993

Methods	Randomised controlled trial Method of randomisation: computerised random Concealment of allocation: assignment from central site Outcome assessor blinding: not reported
Participants	Country: USA To be eligible, Veterans Aaffairs service patients had one of the following: at risk of nursing home placement, 'Service connected disability', hospital inpatient, in home care programme, in a Veterans Affaris domiciliary service Inclusion criteria (one of the following): living in a nursing home, need help for ADL activities, bowel incontinence, significant cognitive impairment, acceptable to day care staff 1236 patients screened (252 not eligible, 158 refused consent) Patients randomised = 826 Baseline status: Sickness Impact Profile (Physical) Mean (SD) 31.7 (18.8) and 33.8 (18.

Hedrick 1993 (Continued)

	4) Male: 96% Age: mean 71.1 years
Interventions	Medical Day Hospital: therapeutically orientated programme providing health maintenance and rehabilitation services. Staff included nurses, rehabilitation therapists, recreation therapists and social worker. Mean attendance over 6 months: 28 days. Usual care: Nursing home, inpatient care, clinic visits, home care etc
Outcomes	12 month follow up Death Institutional care Mini Mental state Sickness Impact Profile Survival Satisfaction Questionnaire Self-rated health Social support scale Katz Instrumental ADL Psychological Distress Scale Caregiver Burden Scale Satisfaction Questionnaire Service use and costs
Notes	No accurate information on patient diagnosis given Evaluation of adult day health care

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computerised random number generator
Allocation concealment (selection bias)	Low risk	Assignment from central site
Blinding of participants and personnel (performance bias) All outcomes	High risk	No reported blinding of participants or personnel. However, it would have been obvious to participants
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No reported blinded outcome assessment
Incomplete outcome data (attrition bias) All outcomes	Low risk	

Hedrick 1993 (Continued)

Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias	
Other bias	Low risk	No other obvious sources of bias	
Hui 1995			
Methods		Stratified by disability Method of randomisation: random number table Concealment of allocation: sealed envelopes	
Participants	Country: Hong Kong Patients admitted to a rehabilitation ward one week after acute stroke Exclusion criteria: age < 65 yrs, previous stroke, dementia, live outside catchment area, Barthel index of 20 Participants randomized = 120 Baseline function: mean (SD) Barthel index 9.9 (4.9) and 10.4 (5.3) Male: 44% Age: mean (SD) 73.6 (5.7) years		
Interventions	Medical day hospital: care under the geriatrician with early discharge, as able, with continued care in the day hospital. Duration of intervention not reported for day hospital or inpatient rehabilitation Conventional inpatient rehabilitation: delivered by a neurology team with medical clinic follow up		
Outcomes	6 month follow up Death Institutional care Abbreviated mental test score Barthel index Self-rated health scale score Geriatric Depression Scale Subjective satisfaction with services Use of hospital and community services Costs		
Notes	Stroke patients only All patients initially treated on same rehabilitation ward		
Risk of bias			
Bias	Authors' judgement	Support for judgement	

Hui 1995 (Continued)

Random sequence generation (selection bias)	Low risk	Random number table (information obtained from follow up correspondence)
Allocation concealment (selection bias)	Low risk	Quote (from letter): "The codes were sealed in envelopes and placed at an office in Shatin Hospital. When the patient is deemed suitable to be discharged, an envelope would be withdrawn and patient assigned into the specific group (Day Hospital or Conventional Management)."
Blinding of participants and personnel (performance bias) All outcomes	High risk	No report of blinding but would have been obvious to participants
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Follow up assessment was carried out by a research nurse. However, not reported if assessment was blinded
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Losses relatively similar across the groups. However, reasons for participants defaulting not reported
Selective reporting (reporting bias)	Unclear risk	Reported that "patient well-beinguse of community services and financial support were all comparable between the two treatment groups at each follow-up (data not shown)." No numerical data reported
Other bias	Low risk	No other obvious sources of bias

Masud 2006

Methods	Randomised controlled trial Method of randomisation: Internet based randomisation service Concealment of allocation: Internet based randomisation service Outcome assessor blinding: not undertaken
Participants	Country: UK Inclusion criteria: the study population was comprised of men and women aged 70 and over identified as being at high risk of falling by a postal screening questionnaire, registered with the participating general practices in Nottinghamshire and Derbyshire. Exclusion criteria: patients already attending one of the day hospitals; under follow-up with an existing primary care based falls prevention scheme; in nursing or residential homes; patients with terminal illnesses; unwilling or unable to travel to the day hospital (using transport as provided); unable to provide informed consent or assent 6113 assessed for study inclusion (844 potential participants did not meet the eligibility criteria, 4925 declined to participate)

Masud 2006 (Continued)

	Participants randomised = 364 Male: 40% Age: mean (SD) 78.8 (5.7) years
Interventions	Medical day hospital: screening questionnaire, information leaflet, leaflet on falls prevention and invitation to attend the day hospital for assessment and any subsequent intervention Control intervention: screening questionnaire, information leaflet, leaflet on falls prevention and usual care from primary care service until outcome data collected, then offer of day hospital intervention Duration of intervention not reported for day hospital or control intervention
Outcomes	Primary outcome: Rate of falls over the 12 month follow-up period Secondary Outcomes Proportion of people with single or recurrent falls and fall-related injuries: fracture, serious sprain requiring immobilisation in plaster, joint dislocations, head injury requiring hospitalisation, and lacerations requiring suturing Disability: Nottingham Extended Activities of Daily Living Scale; Barthel index of daily living; Quality of life: Falls Efficacy Scale and EuroQoL-5 Institutionalisation and use of health services: residency and diary information Cost analysis Deaths checked from PCT records and measured as proportions
Notes	

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	An Internet based randomisation service provided by the hosts institution's clinical trials unit
Allocation concealment (selection bias)	Low risk	An Internet based randomisation service provided by the hosts institution's clinical trials unit
Blinding of participants and personnel (performance bias) All outcomes	High risk	Due to the nature of the intervention It would not be possible to blind participants
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "it was not possible to blind participants or researchers to allocation." However, the review authors judge that the outcome measurement is not likely to be influenced by lack of blinding. GP recording of death or institutionalisation are unlikely to be biased by the participation in either

Masud 2006 (Continued)

		arm of the study
Incomplete outcome data (attrition bias) All outcomes	Low risk	Misisng data balanced across groups with similar reasons
Selective reporting (reporting bias)	Low risk	Study protocol available and additional information provided on request
Other bias	Low risk	No other obvious sources of bias

Parker 2009

Methods	Randomised controlled trial Method of randomisation: external web based randomisation service Concealment of allocation: external web based randomisation service Outcome assessor blinding: unclear
Participants	Country: UK Participants were older people referred for rehabilitation for various conditions including stroke, orthopaedic rehabilitation, movement disorder, mobility assessment and fall assessment Inclusion criteria: referred for multidisciplinary rehabilitation, a permanent address in the catchment area, able to give informed consent (with the help of a career or advocate if necessary) Exclusion criteria: local exclusion criteria meant that patients were excluded from ran domisation if they had a specific clinical need that could only be addressed at one centre (sites provided specific services) Participants randomised = 89 Baseline function: mean (SD) Nottingham Extended Activities of Daily Living 29.9 (15 2) Males: 55% Age: mean (SD) 75 (11) years
Interventions	Medical day hospital (4 sites): Some variation in the services provided between the four day hospitals. However, all sites were multidisciplinary and patients could access a hospital doctor, nursing care, physiotherapy and occupation therapy services. Some sites provided access to a social worker. Number of rehabilitation episodes: mean 17.7 median 18 Rehabilitation at home: Some variation in the services offered by the 4 participating home rehabilitations teams. However, all provided physiotherapy services and the majority provided occupational therapy. Some services provided access to a doctor and a social worker. Number of rehabilitation episodes: mean 9.4; median 8.5 Reported in the study protocol that the length of the interventions would be determined by the local clinical team with the expectation that 95% of participants would be discharged within 16 weeks
Outcomes	12 month follow up Patient outcomes: Hospital anxiety and depression scale

Parker 2009 (Continued)

	Euro-qol 5D
	Nottingham Extended Activities of Daily Living
	Socio-economic data, survival
	Therapy outcome measures
	Views of treatment by qualitative interview
	Carer outcomes:
	General health questionnaire
	Socio-economic data
	Views of treatment
Notes	http://www.controlled-trials.com/ISRCTN71801032 https://portal.nihr.ac.uk/Profiles/NRR.aspx?Publication_ID=N0071140216

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	External web based randomisation service
Allocation concealment (selection bias)	Low risk	External web based randomisation service and investigators were not involved in the allocation to groups
Blinding of participants and personnel (performance bias) All outcomes	High risk	Quote: "the nature of the treatments was such that it was not possible for the patients or their health-care professionals to be blinded to the treatment allocation, or to guarantee that the local researchers remained unaware of allocation for the duration of follow-up"
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	Quote: "the nature of the treatments was such that it was not possible for the patients or their health-care professionals to be blinded to the treatment allocation, or to guarantee that the local researchers remained unaware of allocation for the duration of follow-up"
Incomplete outcome data (attrition bias) All outcomes	High risk	Whilst the reasons for losses were relatively similar across both groups, losses were > 35% by final follow up
Selective reporting (reporting bias)	Low risk	Protocol available (Current Controlled Trials ISRCTN71801032)
Other bias	Low risk	No other obvious sources of bias

Pitkala 1991

Pitkala 1991			
Methods	Concealment of allocation: date of	Randomised controlled trial Method of randomisation: randomised according to date of birth Concealment of allocation: date of birth, therefore could have been foreseen Outcome assessor blinding: not reported	
Participants	All 177 chronically ill patients recei Participants randomised = 174 Male: 34%	Patients receiving home care in a rural community in Finland All 177 chronically ill patients receiving home care screened (3 refused consent) Participants randomised = 174	
Interventions	care. Intensive physiotherapy and oc need. Patients attended 2-3 days a treatment over 2 months	Usual care: included mixture of home health care and referral to a hospital or outpatient	
Outcomes	12 month follow up Death Institutional care Katz ADL Subjective health assessment Mood Resource use, hospital admissions, of Number of symptoms Number of medications	Death Institutional care Katz ADL Subjective health assessment Mood Resource use, hospital admissions, outpatients visits, GP visits Number of symptoms	
Notes		Total of 174 patients of whom 40 had a stroke diagnosis, 54 a diagnosis of coronary heart disease, 53 arthrosis, 33 'moderate' or 'severe' dementia	
Risk of bias			
Bias	Authors' judgement	Supp	port for judgement

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	High risk	Quote: "were randomised into two groups according to their date of birth."
Allocation concealment (selection bias)	High risk	Allocation based on date of birth, therefore could have been foreseen
Blinding of participants and personnel (performance bias) All outcomes	High risk	No report of blinding and would have been obvious to participants which group they were in
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No reported blinded outcome assessment

Pitkala 1991 (Continued)

Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Similar numbers of losses but reasons for losses not reported so cannot determine risk
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias
Other bias	Unclear risk	Nearly one quarter of the day hospital group refused the care

Roderick 2001

Methods	Randomised controlled trial Stratified by sex, age and disability (Barthel index <10; 10-14; >15) and catchment areas of day hospitals Method of randomisation: computer generated Concealment of allocation: unclear Outcome assessor blinding: blinded research nurse
Participants	Country: UK Inclusion criteria: patients with newly diagnosed stroke admitted to a Poole Hospital NHS Trust hospital, or community referrals. Confirmed diagnosis of stroke Aged over 55 years Residents of East Dorset Needed rehabilitation for stroke related disability Were able to attend day hospital No previous disability which would prevent rehabilitation No signs of advanced dementia Exclusion criteria: terminal illness, needing day hospital care for social or medical reasons. 180 eligible Patients randomised = 140 Baseline function: Median (IQR) Barthel Index: Day Hospital 14 (9-17), Control 14 (9-16). Male: 46% Age: mean (range) 78.95 (60-95) years
Interventions	Day hospital: 5 day hospitals with coordinated care from multidisciplinary teams, both individual and group therapies. Median number of visits to the day hospital 17 Domiciliary care: domiciliary stroke team comprising 1 full time physiotherapist and a half time physiotherapist and consultant geriatrician, who met with each other fortnightly to review patients. Out patient speech and language therapy provided. Median number of domicilliary visits 17 In both interventions, therapy was provided until maximum potential for recovery was reached. Patients were then placed on review, and if no further therapy required, discharged

Roderick 2001 (Continued)

Outcomes	6 month follow-up
	Primary outcome:
	Barthel index
	Secondary outcomes:
	Rivermead Mobility Index
	Philadelphia Geriatric Center Morale Scale
	Frenchay Activities Index
	Perceived Quality of Life (SF-36)
	Health and local authority social service costs
Notes	All stroke patients, previous stroke in the day hospital group 23 (32%)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Computer generated randomisation
Allocation concealment (selection bias)	Unclear risk	Allocation concealment was attempted. Quote: "by calling a centralised office where closed lists were kept", the sample was stratified by sex (2 groups), disability (3 groups), age and day hospital catchment area. Minimum 2x3x2x5 = 60 groups. There were 5 day hospitals so potentially 60 groups. The approach to stratification is not described but is likely to be a permuted-block design, with small block size and therefore allocation could have been predicted for some of the patients
Blinding of participants and personnel (performance bias) All outcomes	High risk	Not reported but would have been obvious to participants
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Assesments were carried out by a research nurse blind to allocation
Incomplete outcome data (attrition bias) All outcomes	Low risk	Similar losses in each group with similar reasons
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias

Roderick 2001 (Continued)

Other bias	Low risk	There was contamination: 5 switched from
		domiciliary to day hospital and two the
		other way. One other 'incorrect placement'
		(unexplained). Appears these were anal-
		ysed in original groups (correctly (ITT) but
		therefore contaminating the result). How-
		ever, because the contamination involved
		so few participants we judged it unlikely to
		have significantly altered the estimate of ef-
		fect

Tucker 1984

Tucker 1964	
Methods	Randomised controlled trial Stratified by stroke or non-stroke diagnosis Method of randomisation: random number table Concealment of allocation: unclear Outcome assessor blinding: research occupational therapist
Participants	Country: New Zealand Patients over 55 years Patients needing assessment and rehabilitation but not 24 hour institutional care Referrals from hospital and GPs Excluded: dementia, patients needing social care Baseline function: 17.6 (12-31) and 16.3 (12-25) on Northwick Park ADL score Male: 43% Age: mean (range) 72 (55-92) years
Interventions	Day hospital: intensive physiotherapy, occupational therapy, speech therapy and medica and nursing assessment and supervision. Patients attended 2-3 days per week, Monday to Friday from 8.30 a.m 2.00 p.m. for 6 - 8 weeks Usual care: inpatient, outpatient follow-up with or without outpatient physiotherapy by referral for domiciliary services, by referral to the sole care of their GP, or by referrat to a day centre as decided before randomisation
Outcomes	5 months follow up Death Institutional care Northwick Park ADL Zung Depression Index Service use Costs
Notes	No information on number of patients screened for inclusion Stroke patients randomised separately from other diagnoses (65 of 120)

Tucker 1984 (Continued)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Quote: "Patients with and without strokes were randomised separately into day hospital and control groups with standard tables of random numbers."
Allocation concealment (selection bias)	Unclear risk	Allocation concealment not reported. Therefore, insufficient information to determine risk
Blinding of participants and personnel (performance bias) All outcomes	High risk	No report of blinding. However, it would have been obvious to participants which group they were in
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "In an attempt to preserve blindness of assessment she [research occupational therapist] was not concerned in the rehabilitation of these patients and worked in another occupational therapy department."
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	Some lost to follow up (5% from experimental group, 14% from control). Some differences in reasons for losses
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias
Other bias	Low risk	No other obvious sources of bias

Vetter 1989

Methods	Randomised controlled trial (pilot study) Method of randomisation: not reported Concealment of allocation: sealed envelopes Outcome assessor blinding: not reported
Participants	Country: UK Consecutive patients attending 2 day hospitals were eligible for trial if: Required rehabilitation Had not attended day hospital in previous year Did not require medical investigations only provided in day hospital Not confused 270 patients screened (83 needing maintenance - had attended the day hospital in the previous year, 41 needed medical investigation, 28 confused, 10 required respite, 5 attended only once, 4 refused and 40 not recruited due to administrative problems)

Vetter 1989 (Continued)

	Participants randomised = 59 Baseline function: Barthel index of approximately 13 Male: 32% Age: 98% over 65 years
Interventions	Day hospitals (2 sites): medical and nursing support and physiotherapy, occupational therapy, speech therapy, chiropody, dietary, pharmaceutical and opthalmic services. Home rehabilitation: a newly established service, comprising two part-time physiotherapists, three part-time occupational therapists, speech therapist, dietician, clinical psychologist available for referrals as appropriate Regular team meetings, attempt to equalise amount of therapy given to both groups, the duration of the interventions were not reported
Outcomes	2 month follow up Death Institutional care Barthel index Sickness Impact Profile
Notes	Total of 59 patients of whom 16 had a stroke diagnosis, 12 fractured neck or femur, 5 osteo-arthritis

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Method of sequence generation not reported
Allocation concealment (selection bias)	Low risk	Sealed envelopes opened after participants had been included
Blinding of participants and personnel (performance bias) All outcomes	High risk	No report of blinded outcome assessment. However, it would have been obvious to participants
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No report of blinded outcome assessment
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	The outcomes were not reported in the original report (Vetter 1989), although some additional information was provided on request
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias

Vetter 1989 (Continued)

Other bias	Low risk	No other obvious sources of bias
Weissert 1980		
Methods	Randomised controlled trial Method of randomisation: no information reported Concealment of allocation: no information reported Outcome assessor blinding: no information reported	
Participants	Country: UK New service established and advertised Referral from a number of sources (hospital, community, etc) Patients screened for eligibility for day care service 63% of eligible referred patients agreed to participate Participants randomised = 644 Male: 39% Age: 50% ≥ 75 years	
Interventions	Day hospital: a programme of services including nursing, physiotherapy, patient activities provided under health leadership with physical rehabilitation as the treatment goal. Four different sites available. Patients attended for an average of 51 days per year. Control group: all patients continued to be eligible for existing services, which included	
Outcomes	hospital and skilled nursing inpatient and outpatient care, home health visits 12 month follow up Death Institutional care Katz ADL index Kahn Mental Status Questionniare Contentment scale Social activity Resource use Costs	
Notes	Little information on patient diagnosis (only circulatory disorders (225, 41%) and injuries (55, 10%)) Alternative to institutional care	
Risk of bias		
Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	No information reported
Allocation concealment (selection bias)	Unclear risk	No information reported

Weissert 1980 (Continued)

Blinding of participants and personnel (performance bias) All outcomes	High risk	No information reported. However, it would have been obvious to participants
Blinding of outcome assessment (detection bias) All outcomes	tion Unclear risk No information reported	
Incomplete outcome data (attrition bias) All outcomes	High risk	Missing data and numbers/reasons for drop outs per group not reported
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias
Other bias	Low risk	No other obvious sources of bias

Woodford 1962

Bias

	Stratified by age and sex Method of randomisation: random number table Concealment of allocation: not reported Outcome assessor blinding: not reported
Participants	Country: UK Patients (N = 331) from a consecutive series of 500 former geriatric unit inpatients (169 had died, left area, or were not traced) Inclusion criteria: patients over 60 years without psychiatric disorders
Interventions	Day hospital: patients received a medical assessment, occupational therapy and group exercises Individual physiotherapy provided as required. Chiropody, bathing and hair washing also available. Attended 1 day a week 9am - 5pm Control: usual care with limited resources available.
Outcomes	12 month follow up Death Institutional care Hospital readmission Subjective health assessment by doctor and patient
Notes Risk of bias	No information on patient diagnosis Aimed to reduce demand for hospital admission

Authors' judgement

Support for judgement

Woodford 1962 (Continued)

Random sequence generation (selection bias)	Low risk	Random number table
Allocation concealment (selection bias)	Unclear risk	Not reported
(performance bias) intervention would		No report of blinding and the nature of the intervention would make it unlikely that blinding had been undertaken
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No report of blinded assessment
Incomplete outcome data (attrition bias) All outcomes	High risk	26% of the original 500 participants lost at the outset, numbers relatively balanced across groups but reasons not reported. Some exclusions due to contamination of controls
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias
Other bias	High risk	No other obvious sources of bias

Young 1992

Methods	Randomised controlled trial Stratified by disability and time since stroke Method of randomisation: unclear Concealment of allocation: unclear Outcome assessor blinding: researcher
Participants	Country: UK Inclusion criteria: Patients discharged home from hospital after new stroke event Fit to travel Age > 60 yrs Barthel index < 20 Exclusion criteria: patients who had to attend day hospital for respite care (n = 9) 516 screened for inclusion (143 patients discharged to residential care, 160 patients Barthel score of 20, 40 patients no change in Barthel index score, 25 lived out of area, 9 needed respite care, non-consent 15) Participants radomised = 124 Baseline function: Median (IQR) Barthel index 15 (range 4-19) and 16 (1-19) Male: 56% Age: Median (range) day hospital 72 years (60-88), domiciliary group 70 years (60-89)

Young 1992 (Continued)

Interventions	Day hospital attendance: focus on physical rehabilitation, staffed by a multidisciplinary team of nurses, physiotherapists and occupational therapists. 2 days a week for 8 weeks 9.30am - 3.45pm. Home physiotherapy: to a maximum of 20 hours in 8 weeks.
Outcomes	6 months follow up Death Institutional care Barthel index Functional Ambulatory Catagories Motor Club Assessment Frenchay Activities Index Nottingham Health Profile Carers GHQ-28 Service use Costs (first eight weeks only)
Notes	Stroke patients only (124) Previous stroke 36 29%

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	Quote: "randomisation to one of the two treatment groups was by an independent worker." However, the specific method of randomisation was not reported
Allocation concealment (selection bias)	Low risk	Quote: that "randomisation to one of the two treatment groups was by an independent worker"
Blinding of participants and personnel (performance bias) All outcomes	High risk	No report of blinding of participants or personnel. However, it would have been obvious to participants which group they were in
Blinding of outcome assessment (detection bias) All outcomes	Low risk	Quote: "by a research worker who was not involved with the randomisation or with the patient's treatment"
Incomplete outcome data (attrition bias) All outcomes	Low risk	Similar numbers lost from each group for similar reasons ~20%
Selective reporting (reporting bias)	Unclear risk	We were unable to obtain the pre-study protocol so cannot determine risk of reporting bias

Young 1992 (Continued)

Other bias	Low risk	Some contamination from participants changing intervention group. However, this was only 4%
		tilis was only 470

ADL: activities of daily living GP: general practitioner

Characteristics of excluded studies [ordered by study ID]

Study	Reason for exclusion
Adamowski 2009	Psychiatric patients.
Aimonino Ricauda 2008	General medical ward versus care at home.
Bartak 2011	Psychiatric patients.
Baskett 1999	Patients were randomised to treatment at home or to outpatient/day hospital attendance, patients attending day hospital not reported separately
Baumgarten 2002	Evaluation of adult day care rather than day hospital.
Bjokdahl 2006	Median age of patients was 53 years.
Bussche 2010	Not a randomised controlled trial. This is a qualitative study
Canuto 2008	Not a randomised controlled trial. This is a longitudinal study
Capomolla 2002	Patients with heart failure, mean age 56 years.
Chau 2013	Not a randomised controlled trial. Patients were free to choose which service they attended
Chiu 2009	Psychiatric patients.
Close 1999	The day hospital was part of a more complex intervention.
Coleman 1999	Not an evaluation of a medical day hospital; patients attended a chronic care clinic for half-day visits every 3-4 months
Comans 2010	The intervention took place in a hospital gym and did not meet the criteria for a medical day hospital
Crilly 2005	Not a randomised controlled trial.

(Continued)

Dasgupta 2005	Retrospective review of patients; not a randomised controlled trial
de Oliveira 2010	The intervention was specific to patients with Chronic Obstructive Pulmonary Disease (COPD). Studies of single, specific conditions were excluded (see methods-types of interventions). Appears to be an outpatient intervention rather than day hospital
Del Giudice 2009	Not a randomised controlled trial.
Desrosiers 2004	Not a randomised controlled trial.
Edelman 2010	The intervention facility was a clinic specific to treating patients with both diabetes and hypertension, with a specific tailored intervention. Studies of single, specific conditions were excluded (see methods-types of interventions)
Evans 1998	Hospital-based rehabilitative care versus outpatient services
Famadas 2008	Not a randomised controlled trial.
Foley 2009	Not a randomised controlled trial.
Gitlin 2006	Not a randomised controlled trial.
Glaesmer 2003	Not a randomised controlled trial.
Hershkovitz 2003	Observational study.
Hershkovitz 2007	Not a randomised controlled trial.
Horgan 2009	The study took part in a day hospital but this was not the intervention
Jacob 2007	Not a randomised controlled trial.
Juhani 2011	The outpatient programme was specific to patients with coronary heart disease
Kallert 2007	Psychiatric patients.
Kneebone 2010	Not a randomised controlled trial.
Lariviere 2010	Psychiatric patients.
Lariviere 2011	Psychiatric patients.
Leveille 1998	Evaluated the effect of a chronic illness self-managment programme delivered in a senior centre. All participants attended the senior centre
Luk 2011	Not a randomised controlled trial.
Luk 2011a	Not a randomised controlled trial.

(Continued)

Malone 2002	Prospective study; not a randomised controlled trial.
Manckoundia 2007	Not a randomised controlled trial.
Marsden 2010	The intervention was only for 2.5 hours a week, therefore it does not meet our criteria for a near full day, or full day. The intervention appeared to be more social care rather than rehabilitation (group sessions rather than individualised)
Masuda 2006	Not a randomised controlled trial.
Meinck 2002	Not a randomised controlled trial.
Olsson 2007	Not a randomised controlled trial.
Pereira 2010	Not a randomised controlled trial.
Priebe 2011	Psychiatric patients aged 18-65.
Richardson 2000	Evaluation of different treatment approaches; all patient attended the day hospital
Sato 2007	Patients were in receipt of day services and were randomised to attend a water exercise program once or twice a week or to a social activity control group. The intervention was a swimming intervention, as part of a day service, and not a day hospital
Schweikert 2009	Quote: "As randomization was chosen by only 2.5% of participants, the study had to be analyzed as an observational study."
Scott 2004	Not day hospital intervention, group meeting for 90 minutes once a month
Sherwood 1986	Not a randomised controlled trial.
Skellie 1982	The day hospital arm of the study included data from two other interventions, including home services, and it was not possible to extract the data specific to the day hospital
Spice 2009	Participants only attended day hospital for up to two hours a day
Velghe 2011	Not a randomised controlled trial.
Wade 2003	Evaluation of treatment for patients with Parkinsons disease. Intervention provided by a specialist multi- disciplinary team to patients with Parkinson's disease in a day hospital setting. Studies of single, specific conditions were excluded (see methods-types of interventions)
Weiler 1976	Not a randomised or quasi-randomised study.
Wong 1998	Not a randomised controlled trial.
Zank 2002	Not a randomised controlled trial.

Characteristics of studies awaiting assessment [ordered by study ID]

ISRCTN53696600

Methods	
Participants	Stroke patients
Interventions	Physiotherapy versus standard care
Outcomes	Timed 10 metre walk, questionnaire to establish if there is an improvement in function
Notes	http://www.controlled-trials.com/ISRCTN53696600/53696600

Matzen 2007

Methods	Randomised controlled trial
Participants	
Interventions	
Outcomes	
Notes	

Moe 2010

Methods	
Participants	
Interventions	Multidisciplinary and multifaceted outpatient management of patients with osteoarthritis
Outcomes	
Notes	Protocol for a randomised controlled trial

NCT00785746

Methods	Randomised controlled trial			
Participants	Older people attending the geriatric day hospital			
Interventions	Core-Strength training program in comparison to a Stretch & Strength program			
Outcomes	Berg balance scale, Functional walking capacity 6 minute walk test, Gait speed, Bridge Test, Activities-Specific Balance Confidence Scale, International Consultation on Urinary Incontinence Questionnaire			

NCT00785746 (Continued)

Notes	ClinicalTrials.gov identifier: NCT00785746			
Yamada 2005				
Methods				
Participants	Older patients with dementia			
Interventions				
Outcomes				
Notes				

DATA AND ANALYSES

Comparison 1. Day Hospital vs Alternative Care - patient outcomes

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Death by the end of follow up	16	3533	Odds Ratio (M-H, Random, 95% CI)	1.05 [0.85, 1.28]
1.1 Day Hospital vs	5	1287	Odds Ratio (M-H, Random, 95% CI)	1.26 [0.87, 1.82]
Comprehensive elderly care				
1.2 Day hospital vs	7	901	Odds Ratio (M-H, Random, 95% CI)	0.97 [0.61, 1.55]
Domiciliary care				
1.3 Day hospital vs No comprehensive elderly care	4	1345	Odds Ratio (M-H, Random, 95% CI)	0.88 [0.63, 1.22]
2 Death or institutional care by the end of follow up	13	3030	Odds Ratio (M-H, Random, 95% CI)	0.85 [0.63, 1.14]
2.1 Day hospital vs Comprehensive elderly care	4	1181	Odds Ratio (M-H, Random, 95% CI)	1.00 [0.69, 1.44]
2.2 Day hospital vs Domiciliary care	5	672	Odds Ratio (M-H, Random, 95% CI)	1.05 [0.57, 1.92]
2.3 Day hospital vs No comprehensive elderly care	4	1177	Odds Ratio (M-H, Random, 95% CI)	0.63 [0.40, 1.00]
3 Death or deterioration in activities of daily living (ADL)	7	1268	Odds Ratio (M-H, Random, 95% CI)	1.07 [0.76, 1.49]
3.1 Day hospital vs Comprehensive elderly care	1	174	Odds Ratio (M-H, Random, 95% CI)	1.18 [0.63, 2.18]
3.2 Day hospital vs Domiciliary care	4	443	Odds Ratio (M-H, Random, 95% CI)	1.41 [0.82, 2.42]
3.3 Day hospital vs No comprehensive elderly care	2	651	Odds Ratio (M-H, Random, 95% CI)	0.76 [0.56, 1.05]
4 Death or poor outcome (institutional care, disability or deterioration)	13	2831	Odds Ratio (M-H, Random, 95% CI)	0.92 [0.74, 1.15]
4.1 Day hospital vs Comprehensive elderly care	5	1268	Odds Ratio (M-H, Random, 95% CI)	1.05 [0.79, 1.40]
4.2 Day hospital vs Domiciliary care	5	581	Odds Ratio (M-H, Random, 95% CI)	1.08 [0.67, 1.74]
4.3 Day hospital vs No comprehensive elderly care	3	982	Odds Ratio (M-H, Random, 95% CI)	0.72 [0.53, 0.99]
5 Deterioration in activities of daily living (ADL) in survivors	7	905	Odds Ratio (M-H, Random, 95% CI)	1.11 [0.68, 1.80]
5.1 Day hospital vs Comprehensive elderly care	1	149	Odds Ratio (M-H, Random, 95% CI)	1.21 [0.58, 2.52]
5.2 Day hospital vs Domiciliary care	4	349	Odds Ratio (M-H, Random, 95% CI)	1.59 [0.87, 2.90]
5.3 Day hospital vs No comprehensive elderly care	2	407	Odds Ratio (M-H, Random, 95% CI)	0.61 [0.38, 0.97]
6 Activities of daily living (ADL) scores			Other data	No numeric data

6.1 Day hospital vs	Other data	No numeric data
Comprehensive elderly care		
6.2 Day hospital vs	Other data	No numeric data
Domiciliary care		
6.3 Day hospital vs No	Other data	No numeric data
comprehensive elderly care		
7 Subjective health status	Other data	No numeric data
7.1 Day hospital vs	Other data	No numeric data
Comprehensive elderly care		
7.2 Day hospital vs	Other data	No numeric data
Domiciliary care		
7.3 Day hospital vs No	Other data	No numeric data
comprehensive elderly care		
8 Patient satisfaction	Other data	No numeric data
8.1 Day hospital vs	Other data	No numeric data
Comprehensive elderly care		
8.2 Day hospital vs	Other data	No numeric data
Domiciliary care		
8.3 Day hospital vs No	Other data	No numeric data
comprehensive elderly care		
9 Carer Distress	Other data	No numeric data
9.1 Day hospital vs	Other data	No numeric data
Comprehensive elderly care		
9.2 Day hospital vs	Other data	No numeric data
Domiciliary care		
9.3 Day hospital vs No	Other data	No numeric data
comprehensive elderly care		

Comparison 2. Day Hospital vs Alternative Care - resource outcomes

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Requiring institutional care at the end of follow up	13	3003	Odds Ratio (M-H, Random, 95% CI)	0.84 [0.58, 1.21]
1.1 Day hospital vs Comprehensive elderly care	4	1181	Odds Ratio (M-H, Random, 95% CI)	0.91 [0.70, 1.19]
1.2 Day hospital vs Domiciliary care	5	672	Odds Ratio (M-H, Random, 95% CI)	1.49 [0.53, 4.25]
1.3 Day hospital vs No comprehensive elderly care	4	1150	Odds Ratio (M-H, Random, 95% CI)	0.58 [0.28, 1.20]
2 Hospital bed use during follow			Other data	No numeric data
up 2.1 Day hospital vs Comprehensive elderly care			Other data	No numeric data
2.2 Day hospital vs Domiciliary care			Other data	No numeric data

2.3 Day hospital vs No	Other data	No numeric data
comprehensive elderly care		
3 Resource use	Other data	No numeric data
3.1 Day hospital vs	Other data	No numeric data
Comprehensive elderly care		
3.2 Day hospital vs	Other data	No numeric data
Domiciliary care		
3.3 Day hospital vs No	Other data	No numeric data
comprehensive elderly care		

WHAT'S NEW

Last assessed as up-to-date: 24 April 2014.

Date	Event	Description	
24 April 2014	New search has been performed	New searches performed and three new trials identified	
24 April 2014	New citation required but conclusions have not changed	Three new trials have been identified, five studies are awaiting classification	

HISTORY

Review first published: Issue 3, 1999

Date	Event	Description
12 November 2008	Amended	Minor changes
14 August 2008	New search has been performed	New search
10 July 2008	Amended	Converted to new review format.
22 June 2008	New citation required but conclusions have not changed	This review is an update of the review first published in 1999. The total number of studies included is 16

CONTRIBUTIONS OF AUTHORS

AF, JY and PL planned and initiated the original review, assessed trials, drafted the final report and were guarantors of the initial review. AF oversaw literature searching and PL provided methodological support.

For this edition, LB, AF, TC and AB screened titles and abstracts for study inclusion. LB and AB extracted data and LB drafted the final report.

DECLARATIONS OF INTEREST

Two of the authors of this review, JY and AF, were involved in one of the included studies (Young 1992).

SOURCES OF SUPPORT

Internal sources

- Bradford Teaching Hospitals NHS Foundation Trust, UK.
- University of Glasgow, UK.
- Raithby Bequest, UK.

University of Leeds, School of Medicine

External sources

- Northern and Yorkshire Region NHS Executive, UK.
- Department of Health Cochrane Review Incentive Scheme 2007, UK.

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Methods to account for unit of analysis issues were added in the methods section.

INDEX TERMS

Medical Subject Headings (MeSH)

*Geriatrics; Activities of Daily Living; Day Care, Medical [*standards; statistics & numerical data]; Health Services Research; Health Services for the Aged [*statistics & numerical data]; Home Care Services [standards]; Randomized Controlled Trials as Topic; Treatment Outcome

MeSH check words

Aged; Humans