

Home based versus centre based cardiac rehabilitation: Cochrane systematic review and meta-analysis

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ABSTRACT

Objective To compare the effect of home based and supervised centre based cardiac rehabilitation on mortality and morbidity, health related quality of life, and modifiable cardiac risk factors in patients with coronary heart disease.

Design Systematic review.

Data sources Cochrane Central Register of Controlled Trials (CENTRAL) in the Cochrane Library, Medline, Embase, CINAHL, and PsycINFO, without language restriction, searched from 2001 to January 2008.

Review methods Reference lists checked and advice sought from authors. Included randomised controlled trials that compared centre based cardiac rehabilitation with home based programmes in adults with acute myocardial infarction, angina, or heart failure or who had undergone coronary revascularisation. Two reviewers independently assessed the eligibility of the identified trials and extracted data independently. Authors were contacted when possible to obtain missing information.

Results 12 studies (1938 participants) were included. Most studies recruited patients with a low risk of further events after myocardial infarction or revascularisation. No difference was seen between home based and centre based cardiac rehabilitation in terms of mortality (relative risk 1.31, 95% confidence interval 0.65 to 2.66), cardiac events, exercise capacity (standardised mean difference -0.11, -0.35 to 0.13), modifiable risk factors (weighted mean difference systolic blood pressure (0.58 mm Hg, -3.29 mm Hg to 4.44 mm Hg), total cholesterol (-0.13 mmol/l, -0.31 mmol/l to 0.05 mmol/l), low density lipoprotein cholesterol (-0.15 mmol/l, -0.31 mmol/l to 0.01 mmol/l), or relative risk for proportion of smokers at follow-up (0.98, 0.73 to 1.31)), or health related quality of life, with the exception of high density lipoprotein cholesterol (-0.06, -0.11 to -0.02) mmol/l). In the home based participants, there was evidence of superior adherence. No consistent difference was seen in the healthcare costs of the two forms of cardiac rehabilitation.

Conclusions Home and centre based forms of cardiac rehabilitation seem to be equally effective in improving clinical and health related quality of life outcomes in patients with a low risk of further events after myocardial

infarction or revascularisation. This finding, together with the absence of evidence of differences in patients' adherence and healthcare costs between the two approaches, supports the further provision of evidence based, home based cardiac rehabilitation programmes such as the "Heart Manual." The choice of participating in a more traditional supervised centre based or evidence based home based programme should reflect the preference of the individual patient.

INTRODUCTION

Coronary heart disease is a major cause of mortality and morbidity.¹⁻³ Although mortality from coronary heart disease has decreased in many developed countries in recent decades, morbidity is increasing as a result of improved diagnosis and more successful treatment of acute illness, which has resulted in an increase in the number of people who survive myocardial infarction.^{1,2} Cardiac rehabilitation is offered to people after cardiac events to aid recovery and prevent further cardiac illness. It has been shown to improve physical health and decrease subsequent morbidity and mortality in patients with coronary heart disease (myocardial infarction and after revascularisation). Two systematic reviews that included 48 randomised controlled trials showed a 20% reduction in all cause mortality and a 27% reduction in cardiac mortality at two to five years.^{4,5} Cardiac rehabilitation programmes typically achieve this through exercise, education, behaviour change, counselling, support, and strategies aimed at targeting traditional risk factors for cardiovascular disease. Cardiac rehabilitation is an essential part of the contemporary care of patients with heart disease and is considered a priority in countries with a high prevalence of coronary heart disease and heart failure.⁶⁻¹⁰

Although cardiac rehabilitation has been shown to have beneficial effects, participation remains suboptimal. The main reasons people give for not accepting the invitation to attend centre based cardiac rehabilitation classes—held for groups in hospitals, gyms, or community leisure centres—are problems with accessibility and parking at their local hospital,¹¹⁻¹³ a dislike of groups,¹⁴ and work or domestic commitments.¹⁵⁻¹⁸ These problems can be overcome by home based

Table 1 Characteristics of excluded studies

| Citation | Reason for exclusion |
|------------------------------------|--|
| Ades et al, ^{w1} 2000 | Not randomised controlled trial |
| Tygssen et al, ^{w2} 2001 | Both trial arms received home based cardiac rehabilitation |
| Senuzun et al, ^{w3} 2006 | Trial arms involved home based cardiac rehabilitation and usual care |
| Sinclair et al, ^{w4} 2005 | Trial arms involved home based cardiac rehabilitation and usual care |

programmes, which have been introduced in an attempt to widen access and participation.¹⁹

The one systematic review (of randomised controlled trials) that compared home based and centre based cardiac rehabilitation to date found no significant differences in outcomes,²⁰ but there were only 750 participants in total and patients with heart failure were excluded. Two large randomised controlled trials that compared home based and centre based cardiac rehabilitation in the United Kingdom have recently been completed.^{21,22} We determined the effectiveness of home based cardiac rehabilitation programmes compared with supervised centre based cardiac rehabilitation on mortality, morbidity, health related quality of life, and modifiable cardiac risk factors in patients with coronary heart disease.

METHODS

Criteria for considering studies for this review

Studies included—We included randomised controlled trials (individual or cluster level).

Types of participants—The study population included adults with myocardial infarction, angina, or heart failure and patients who had undergone revascularisation (coronary artery bypass grafting, percutaneous transluminal coronary angioplasty, or coronary artery stenting). All had taken part in or been invited to take part in cardiac rehabilitation. We excluded studies of participants with heart transplants, cardiac resynchronisation devices, or implantable defibrillators.

Types of interventions—Home based cardiac rehabilitation was defined as a structured programme, with clear objectives for the participants, including monitoring, follow-up, visits, letters, telephone calls from staff,

or at least self monitoring diaries. Centre based cardiac rehabilitation was a supervised group based programme undertaken in a hospital or community setting such as a sports centre.

Types of outcome measures—Outcome measures included mortality (cardiac and overall), morbidity (reinfarction, revascularisation, and admission to hospital associated with cardiac disease), exercise capacity, modifiable coronary risk factors (smoking behaviour, blood lipid concentrations, and blood pressure), health related quality of life, adverse events (withdrawal from the exercise programme), health service use or costs, and cost effectiveness. During the review (and before any data analysis was undertaken), we agreed to include the outcome of adherence to the intervention.

Search methods for identification of studies

We identified randomised controlled trials from the previously published systematic review.²⁰ We updated the list of studies by searching several clinical databases from 2001 to January 2008: the Cochrane Central Register of Controlled Trials (CENTRAL) in the Cochrane Library, Medline, Embase, CINAHL, and PsycINFO. We located additional studies in the databases of the NHS Centre for Reviews and Dissemination (the Health Technology Assessment (HTA) database and the Database of Abstracts of Reviews of Effects (DARE)). We searched conference proceedings on the ISI Web of Knowledge. We identified grey literature through a web search of major health technology appraisal agencies. We limited searches to randomised controlled trials, systematic reviews, and meta-analyses and imposed no language or other limitations.

Selection of studies—Two reviewers (RST and Philippa Davies) independently screened the titles and abstracts of studies identified by the search strategy and discarded clearly irrelevant studies. To be selected, abstracts needed to clearly identify the study design, define an appropriate population, and describe relevant components of the intervention, as defined above. The same two reviewers also independently assessed the full text reports of all potentially relevant trials and assessed them independently for eligibility on the basis of the defined inclusion criteria. They resolved any disagreement by discussion; if any uncertainty remained, they sought the opinion of two further reviewers (KJ and AZ).

Data extraction and management—Data were extracted by a single reviewer (AZ) and checked by a second reviewer (RST). Standardised data extraction forms were used to extract relevant data regarding inclusion criteria (study design, participants, interventions, and outcomes), risk of bias (randomisation, blinding, attrition, and outcome reporting), and results. In cases for which insufficient details were reported, AZ contacted the authors for further information.

Assessment of risk of bias in included studies—The risk of bias in eligible trials was assessed by a single reviewer (AZ) and checked by a second reviewer (RST) in accordance with recent guidance.²³ The risk of bias was

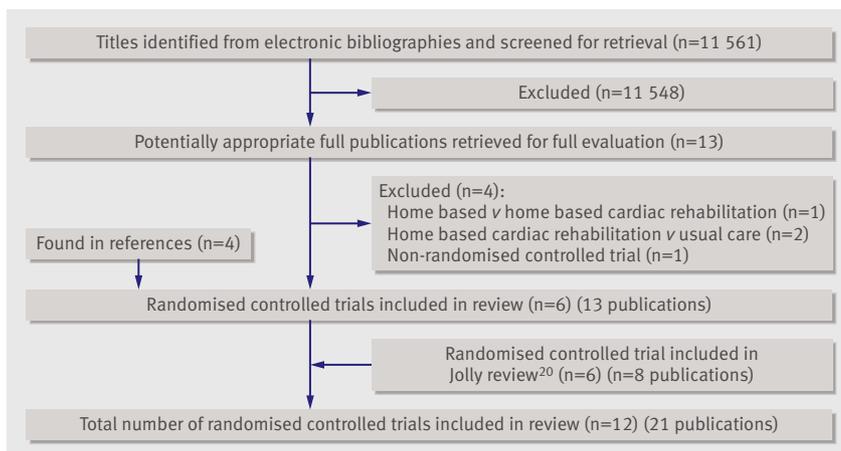
**Fig 1** Summary of study selection process

Table 2 | Summary of included studies

| Study | Methods | No of participants | Interventions | Outcomes | Follow-up | Subgroup analyses | Country, setting |
|--|--------------------|--------------------|---|--|---|---|-------------------------------|
| Arthur et al, ³⁶ 2002, Smith et al, ³³ 2004 | RCT parallel group | 242 | Home v centre based | Primary: exercise capacity (METs). Secondary: HRQoL (SF-36), cardiac morbidity, mortality | 6 and 18 months after randomisation | No subgroups described or reported | Canada, single centre |
| Bell et al, ³⁰ 1998 | RCT parallel group | 252 | Home (Heart Manual) v centre based | Primary: exercise capacity (METs). Secondary: total cholesterol, systolic blood pressure, HRQoL (NHP), smoking, mortality, readmission rate, use of primary care services | 16 and 48 weeks after randomisation (20 and 52 weeks after myocardial infarction) | No subgroups described or reported | UK, five district hospitals |
| Carlson et al, ³² 2000 | RCT parallel group | 80 | Home v centre based | Primary: peak functional capacity (METs), LDL cholesterol. Secondary: total cholesterol, HDL cholesterol, triglycerides, blood pressure, cardiovascular medications, costs, adherence (exercise sessions attended) | 6 months after randomisation | No subgroups described or reported | US, single hospital centre |
| Dalal et al, ²¹ 2007, Taylor et al, ³¹ 2007 | RCT parallel group | 104 | Home (Heart Manual) v centre based | Primary: quality of life (MacNew questionnaire), total cholesterol. Secondary: exercise capacity (METs), self reported smoking, cardiovascular morbidity, mortality, secondary prevention medication | 9 months after randomisation | No subgroups described or reported | UK, single centre |
| Daskapan et al, ³⁴ 2005 | RCT parallel group | 29 | Home v centre based | Exercise capacity (ml/kg/min)*, resting BP*, systolic and diastolic BP*, adherence*, drop outs* | 12 weeks after randomisation | No subgroups described or reported | Turkey, single centre |
| Gordon et al, ²⁶ 2002 | RCT parallel group | 155 | Supervised home v community home v centre based | Maximal oxygen uptake*, blood pressure*, fasting serum lipids*, self reported smoking status*, rehospitalisation*, adherence (completion of appointments)* | 12 weeks after randomisation | Changes reported for all patients and for patients with baseline values defined as abnormal | US, single centre |
| Jolly et al, ²² 2007 | RCT parallel group | 525 | Home (Heart Manual) v centre based | Primary: serum cholesterol, total cholesterol, HDL cholesterol, blood pressure, exercise capacity (ISWT), smoking (validated by cotinine). Secondary: quality of life (EQ-5D), SF-12, health service use (hospital readmissions, primary care visits, medication), mortality, cardiovascular events, costs | 6, 12, and 24 months | Yes ("interaction terms between these factors (diagnosis (MI/revascularisation), age, sex and ethnicity) and rehabilitation setting were included to investigate possible differences in treatment effect between subgroups of patients") | UK, four hospital centres |
| Kassaian et al, ²⁵ 1998 | RCT parallel group | 125 | Home v centre based | Systolic BP*, diastolic BP*, heart rate (all resting and submaximal)*, functional capacity (METs)*, BMI*, cholesterol (total, LDL, HDL, triglycerides)* | 12 weeks after randomisation | Comparison of functional capacity, submaximal systolic BP, diastolic BP and heart rate in patients with left ventricular dysfunction versus good left ventricular function | Iran, single centre |
| Marchionni et al, ²⁴ 2003 | RCT parallel group | 180 | Home v centre based | Primary: TWC. Secondary: HRQoL (SIP), mortality, morbidity (cardiovascular events), healthcare use (medical visits, rehospitalisations), costs, adherence (number of completed training sessions) | 2, 8, and 14 months after randomisation | Subgroup analysis by age (years): middle aged (45-65), old (65-75), very old (>75) | Italy, single hospital centre |
| Miller et al, ²⁷ 1984, DeBusk et al, ²⁸ 1985, Taylor et al, ²⁹ 1986 | RCT parallel group | 127 | Home v centre based | Exercise capacity*, mortality*, cardiovascular morbidity* | 23 weeks after randomisation | Results reported according to two subgroups reported (brief v extended exercise training) | US, single hospital centre |
| Sparks et al, ³⁷ 1993 | RCT parallel group | 20 | Home v centre based | Exercise capacity (peak VO ₂ max), adherence (compliance with exercise), safety (drop out) | 12 weeks after randomisation | No subgroups described or reported | US, single hospital centre |
| Wu et al, ³⁵ 2008 | RCT parallel group | 36 | Home v centre based | Exercise capacity (METs)* | 12 weeks after randomisation | No subgroups described or reported | Taiwan (China), single centre |

BMI=body mass index; CR=cardiac rehabilitation; HDL=high density lipoprotein; HRQoL=health-related quality of life; ISWT=incremental shuttle walking test; LDL=low density lipoprotein; MET=metabolic equivalent; NHP=Nottingham health profile; RCT=randomised controlled trial; SIP=sickness impact profile; TWC=total work capacity; VO₂max=maximum volume of oxygen.

*Primary and secondary outcomes not distinguished.

assessed in terms of the quality of random sequence generation and allocation concealment, description of dropouts and withdrawals, blinding (participants, personnel, and outcome assessment), and selective outcome reporting. We determined whether groups were balanced at baseline and whether an intention to treat analysis was undertaken.

Data synthesis

We processed data in accordance with the Cochrane handbook.²³ For dichotomous variables, we derived the relative risks and 95% confidence intervals for each outcome. For continuous variables, we calculated the mean differences and 95% confidence intervals for each outcome. When the results at follow-up and

Table 3 | Summary of intervention details^{21 22 24-27 30 32 34-37}

| Study | Home based intervention | Centre based intervention |
|--|--|--|
| Arthur (2002) ³⁶ | | |
| Exercise | Total duration: 6 months, 5 sessions/week, 40 min/session. Intensity: 60-70% of maximum oxygen intake. Modality: walking. Also attended 1 hour's exercise consultation with exercise specialist at baseline and after 3 months of training and completed exercises log that was reviewed every 2 months and telephone support call every 2 weeks | Total duration: 6 months, 3 sessions/week, 40 min/session. Intensity: 60-70% of maximum oxygen intake. Modality: cycle ergometer, treadmill, track walking and stair climbing. Supervised by exercise specialist and completed exercises log, reviewed every month |
| Other | Dietary advice and psychological support | Dietary advice and psychological support |
| Bell (1998) ³⁰ Heart Manual | | |
| Exercise | Overall duration: 6 weeks. Frequency, duration, and intensity: NR | Overall duration and frequency: 12 weeks of 1 session/week or 4 weeks of 2 sessions/week, ≥20 min/session. Intensity: 3-4 on Borg rating of perceived exertion (RPE) scale |
| Other | Four phone calls by facilitator, health education, and stress management | Education sessions: causes of coronary heart disease, medication, risk factor modification, stress management, exercise |
| Carlson (2000) ³² | | |
| Exercise | Overall duration: 25 weeks, 2-5 sessions/week, 30-40 min/session. Intensity: 60-85% of aerobic capacity. Modality: aerobic exercise; for first 4 weeks, 3 hospital based exercise sessions/week with electrocardiographic monitoring and then progressive reduction in frequency of centre based sessions | Overall duration: 25 weeks, 2-3 sessions/week, 30-45 min/session. Intensity: 60-85% of aerobic capacity. Modality: aerobic exercise |
| Other | Weekly educational and counselling meetings included sessions on exercise, diet, risk factors, drugs, and overcoming barriers to behaviour change. Based on Bandura's self efficacy theory | Three sessions of education and counselling that included sessions on exercise, diet, risk factors, and drugs |
| Dalal (2007) ²¹ Heart Manual | | |
| Exercise | Overall duration: 6 weeks. Frequency, duration, and intensity: NR. Modality: walking. Home visit in first week after discharge by cardiac rehabilitation nurse followed by up to 4 telephone calls at 2, 3, 4, and 6 weeks | Overall duration: 8-10 weeks, 1-5 sessions/week. Duration and modality: NR. Supervised and group based |
| Other | Stress management and education | Input from dietician, psychologist, occupational therapist and pharmacist |
| Daskapan (2007) ³⁴ | | |
| Exercise | Total duration: 12 weeks, 3 sessions/week, 45 min/session (including warm up, cool down, and recovery). Intensity: up to 60% of peak heart rate (12-16 on Borg rating of perceived exertion (RPE) scale). Modality: walking. Follow-up logs completed daily/returned biweekly. Weekly phone calls from staff to monitor adherence and progress. Monthly phone calls from patients for control purposes | Total duration: 12 weeks, 3 sessions/week, 45 min/session (including warm up, cool down, and recovery). Intensity: 60% of peak heart rate. Modality: walking on a treadmill. Supervised |
| Other | NR | NR |
| Gordon (2002) ²⁶ | | |
| Exercise | Group I (supervised home based cardiac rehabilitation): total duration: 12 weeks. Intensity: individually prescribed (30-60 min of aerobic exercise at 60-85% of peak heart rate) gradually updated; appointments: two office visits and four phone calls. Group II (community home based cardiac rehabilitation): total duration: 12 weeks, individually prescribed (30-60 minutes of aerobic exercise at 60-85% of peak heart rate), gradually updated. Appointments: 12 onsite visits or telephone calls (patient choice) | Total duration: 12 weeks, 3/sessions/week (total of 36 sessions and appointments). Intensity: individually prescribed (30-60 minutes of aerobic exercise at 60-85% of peak heart rate). Continuous electrocardiographic telemetry during exercise |
| Other | Group I and group II: written materials, audiotapes, nutrition, weight and stress management, smoking cessation programme, individual management of risk factors for coronary artery disease | Written materials, audiotapes, education on risk factors for coronary artery disease, and lifestyle modification |
| Jolly (2007) ²² Heart Manual/Centre based cardiac rehabilitation (control) | | |
| Exercise | Overall duration: six weeks of Heart Manual's programme and 12 weeks of nurse support. Frequency: up to daily. Duration and intensity: NR. Modality: walking | Total duration: 6-12 weeks, 1-2 sessions/week, 25-30 min/session. Intensity: 65-75% of maximum heart rate. Modality: circuit training, cycle ergometer |
| Other | Education about risk factors, lifestyle changes, medications and stress management (relaxation tapes) | Education and stress management (relaxation) |
| Kassaian (2000) ²⁵ | | |
| Exercise | Total duration: 12 weeks. Frequency and duration: NR. Intensity: "intensity based on exercise test results" | Total duration: 12 weeks, 3 sessions/week, 20-30 min, 10 min warm up, and 10 min cool-down/session. Intensity: 60-85% (NR if relative to maximum heart rate or maximum oxygen intake). Modality: treadmill |
| Other | Patients taught to count their pulse rate | NR |
| Marchionni (2003) ²⁴ | | |
| Exercise | Overall duration: 8 weeks, 3 days/week, 1 h/session. Intensity: 70-85% of peak heart rate. Modality: cycle ergometer. Physical therapist home visits every other week | Overall duration: programme of 12 weeks, 3 days/week. Duration: NR. Intensity 70-85% of peak heart rate. Modality: cycle ergometer. Trans-telephonic electrocardiographic monitoring during exercise |
| Other | Monthly family oriented support groups | Risk factor management counselling; support group meetings |
| Miller (1984) ²⁷ | | |
| Exercise | Overall duration: 8 weeks, 3 days/week, 1 h/session. Intensity: 70-85% of peak heart rate. Modality: cycle ergometer. Physical therapist home visits every other week | Overall duration: 8 weeks (brief) or 23 weeks (extended), 5 sessions/week, 60 min/session. Intensity: 70-85% of maximum heart rate. Modality: walking/jogging. Group based and supervised |
| Other | Monthly family oriented support groups | No education or psychological intervention reported |
| Sparks (1993) ³⁷ | | |
| Exercise | Overall duration: 12 weeks, 3 days/week, 1 h/session. Intensity 60-75% of peak heart rate. Modality: cycle ergometer. Trans-telephonic electrocardiographic monitoring | Overall duration: 12 weeks, 3 days/week, 1 h/session. Intensity 60-75% of peak heart rate. Modality: cycle ergometer. No trans-telephonic electrocardiographic monitoring |
| Other | Education materials on diet, medications, and risks and benefits of exercise | Education materials on diet, medications, and risks and benefits of the exercise |
| Wu (2006) ³⁵ | | |
| Exercise | Total duration: 12 weeks, ≥3 sessions/week, 30-60 min + 10 min warm up + 10 min cool-down/session. Intensity: 60-85% of maximum heart rate. Modality: fast walking or jogging. Exercise documented in record book. Prescription of exercise individually given and updated every two weeks by rehabilitation nurse | Total duration: 12 weeks, 3 sessions/week (total 36 sessions), 30-60 min + 10 minute warm up + 10 minute cool-down/session. Intensity: 60-85% of maximum heart rate. Modality: cycle ergometer, treadmill. Exercise supervised by cardiopulmonary physical therapist |
| Other | NR | NR |

NR=not reported.

Table 4 | Summary of risk of bias assessment

| Study | Adequate sequence generation | Allocation concealment | Outcome blinding | Incomplete outcome data addressed | Intention to treat analysis |
|---------------------------------|------------------------------|------------------------|------------------|---|-----------------------------|
| Arthur (2002) ³⁶ | Unclear | Yes | Yes | Yes (dropout rate: 8% and 10% at 6 and 18 months, respectively) | Yes |
| Bell (1998) ³⁰ | Unclear | Yes | Yes | Unclear | No |
| Carlson (2000) ³² | Unclear | Unclear | Unclear | No | Yes |
| Dalal (2007) ²¹ | Yes | Yes | Yes | Yes | Yes |
| Daskapan (2007) ³⁴ | Unclear | Unclear | Unclear | Yes | No |
| Gordon (2002) ²⁶ | Unclear | Unclear | Unclear | Yes | Unclear |
| Jolly (2007) ²² | Yes | Yes | Yes | Yes | Yes |
| Kassaian (2000) ²⁵ | Unclear | Unclear | Unclear | Unclear | Unclear |
| Marchionni (2003) ²⁴ | Unclear | Unclear | Yes | Yes | Yes |
| Miller (1984) ²⁷ | Unclear | Unclear | Unclear | Yes | Yes |
| Sparks (1993) ³⁷ | Unclear | Unclear | Unclear | Yes | Yes |
| Wu (2006) ³⁵ | Unclear | Unclear | Yes | Unclear | Unclear |

differences between groups for each of the individual trials were not reported in the original publications, we calculated P values for the differences.²³

We explored heterogeneity among the included studies qualitatively (by comparing the characteristics of included studies) and quantitatively (using the χ^2 test of heterogeneity and I² statistic). When appropriate, we combined the results from included studies for each outcome to give an overall estimate of the treatment effect. We used a fixed effect model for meta-analysis, except where we identified statistical heterogeneity, when we used a random effects model instead. In the protocol for this review, we intended to use meta-regression to explore heterogeneity and examine potential treatment effect modifiers. Given the small number of included trials, however, such analyses were deemed inappropriately underpowered. Instead, we undertook specific stratified meta-analyses to examine the sensitivity of the findings of the review to key potential causes of heterogeneity. Continuous outcomes were pooled as a weighted mean difference except for the exercise capacity measures, which we expressed as a standardised mean difference because of the variety of measures. In the study that reported continuous outcome findings by three age subcategories, we pooled results to produce a single omnibus score for each group.²⁴

We used sensitivity analysis to examine two areas of uncertainty. Firstly, for exercise capacity, in addition to pooling all trials with standardised mean difference, we pooled most trials that reported outcomes as metabolic equivalents using weighted mean difference. Secondly, the lack of detailed reporting meant that there was some doubt about whether or not the study by Kassaian et al was a true comparison between home based and hospital based cardiac rehabilitation or was, instead, a comparison of hospital based cardiac rehabilitation and usual care.²⁵ We undertook all meta-analyses with and without this trial.

Gordon et al compared two home based exercise programmes with a centre based cardiac rehabilitation programme²⁶ and Miller et al reported results in

subgroups by the duration of intervention.²⁷⁻²⁹ For these two studies, we included outcome results in the meta-analysis separately for the two home based groups and for the short and long duration exercise groups, respectively.

RESULTS

Description of studies

Search results—The systematic review by Jolly et al identified six trials (eight papers),²⁰ all of which met the inclusion criteria of this review. Our updated electronic searches yielded a total of 11 561 titles. After we reviewed the titles and abstracts, we retrieved an additional 13 full papers for possible inclusion and identified a further four from the reference lists of eligible publications. We excluded four papers (see bmj.com). In total, we included 21 papers that reported on 12 studies. Figure 1 summarises the study selection process, and table 1 gives details of the excluded studies and reasons for their exclusion.

Included studies—Three studies were based in the UK,^{21 22 30 31} four in the United States,^{26-29 32} and one each in Canada,^{33 34} Turkey,³⁵ Italy,²⁴ Iran,⁴ and China³⁶ (table 2). Most studies reported outcomes up to 12 months after randomisation; only three reported longer follow-up of 24 months,²² 14 months,²⁴ and 18 months.^{33 34} Eight studies compared comprehensive programmes (that is, exercise plus education or psychological management, or both), while the remainder reported only on an exercise intervention.^{25 27-29 35-37} The cardiac rehabilitation programmes differed considerably in duration (range 1.5-6 months), frequency (1-5 sessions/week), and session length (20-60 minutes/session). Most programmes involved the prescription of individually tailored exercise programmes, which makes it difficult to precisely quantify the amount of exercise undertaken. Several home based programmes included a short initial period of centre based intervention.^{26 32 34} Centre based programmes typically involved supervised exercise involving cycles and treadmills, while virtually all home based programmes were based on walking, with

Table 5 | Summary of effects of home based versus centre based cardiac rehabilitation^{21 22 24-36}

| Outcome or subgroup | No of studies | No of participants | Summary estimate and model | Effect estimate (95% CI) | Heterogeneity | | | |
|---|---------------|--------------------|--|--------------------------|---------------|----|---------|--------------------|
| | | | | | χ^2 | df | P value | I ² (%) |
| Exercise capacity: | | | | | | | | |
| At 3-12 month follow-up | 14 | 1938 | Standard mean difference, random effects model | -0.11 (-0.35 to 0.13) | 60.91 | 13 | <0.001 | 79 |
| At 12-24 month follow-up | 4 | 1074 | Standard mean difference, fixed effects model | 0.11 (-0.01 to 0.23) | 0.87 | 3 | 0.62 | 0 |
| Blood pressure (mm Hg) at 3-12 month follow-up: | | | | | | | | |
| Systolic | 9 | 1053 | Mean difference, random effects model | -0.51 (-4.63 to 3.61) | 23.01 | 7 | 0.002 | 70 |
| Diastolic | 7 | 927 | Mean difference, random effects model | 1.85 (0.74 to 2.96) | 7.97 | 6 | 0.24 | 25 |
| Cholesterol (mmol/l) at 3-12 month follow-up: | | | | | | | | |
| Total | 7 | 1019 | Mean difference, random effects model | 0.13 (-0.05 to 0.31) | 13.33 | 6 | 0.04 | 55 |
| High density lipoprotein | 5 | 793 | Mean difference, fixed effects model | -0.06 (-0.11 to -0.02) | 4.44 | 3 | 0.22 | 32 |
| Low density lipoprotein | 4 | 324 | Mean difference, fixed effects model | 0.15 (-0.01 to 0.31) | 6.53 | 4 | 0.16 | 39 |
| Triglycerides | 4 | 328 | Mean difference, random effects model | 0.15 (-0.11 to 0.41) | 7.58 | 3 | 0.06 | 60 |
| Smoking | 5 | 922 | Relative risk, fixed effects model | 1.02 (0.76 to 1.37) | 4.48 | 4 | 0.34 | 11 |
| Completers | 10 | 1714 | Risk ratio, fixed effects model | 1.00 (0.97 to 1.04) | 11.4 | 10 | 0.32 | 13 |
| Mortality | 4 | 909 | Relative risk, fixed effects model | 1.31 (0.65 to 2.66) | 1.00 | 3 | 0.80 | 0 |

some level of intermittent telephone support from a nurse or exercise specialist (table 3). Most studies recruited patients at low risk of another event after acute myocardial infarction or revascularisation, excluding those with severe arrhythmias, ischaemia, or heart failure. Two studies included patients with New York Heart Association class 2 or 3 heart failure.^{25,35}

Risk of bias in included studies

Several studies failed to give enough detail for us to assess the potential risk of bias. Details of the generation and concealment of the random allocation sequence was particularly poorly reported. Only one study presented objective evidence of an imbalance in baseline characteristics.^{33,34} Only six studies stated that they took measures to blind those involved in assessments. Losses to follow-up varied considerably and

were often asymmetric across the home based and centre based cardiac rehabilitation groups. Although the type of analysis was often not stated, all studies seemed to undertake an intention to treat analysis, in that groups were analysed according to the initial random allocation. Only a few trials examined the impact of losses to follow-up or drop out. As discussed above, the rehabilitation intervention was usually tailored to the individual patient, so it is difficult to quantify the precise level of intervention; however, the intensity of the rehabilitation programme seemed to differ substantially between the home based and centre based arms. Table 4 gives a summary of the assessment of risk of bias for each included study.

Effects of interventions on exercise capacity

All 12 included trials reported exercise capacity in the short term (3-12 months of follow-up), while

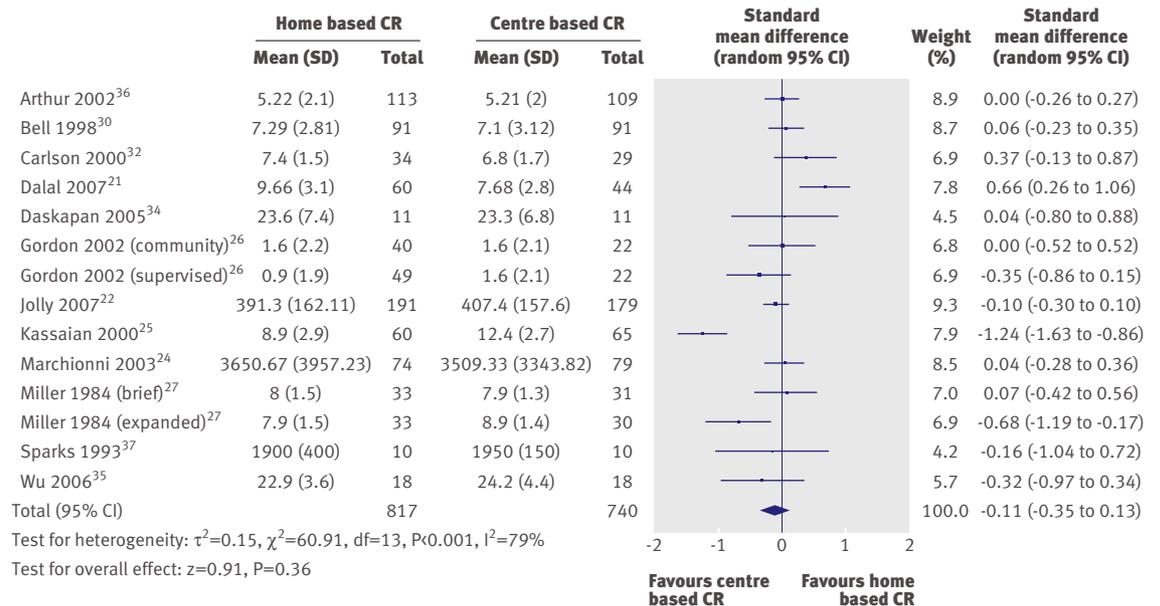


Fig 2 | Exercise capacity with home based and centre based cardiac rehabilitation (CR) at 3-12 months of follow-up

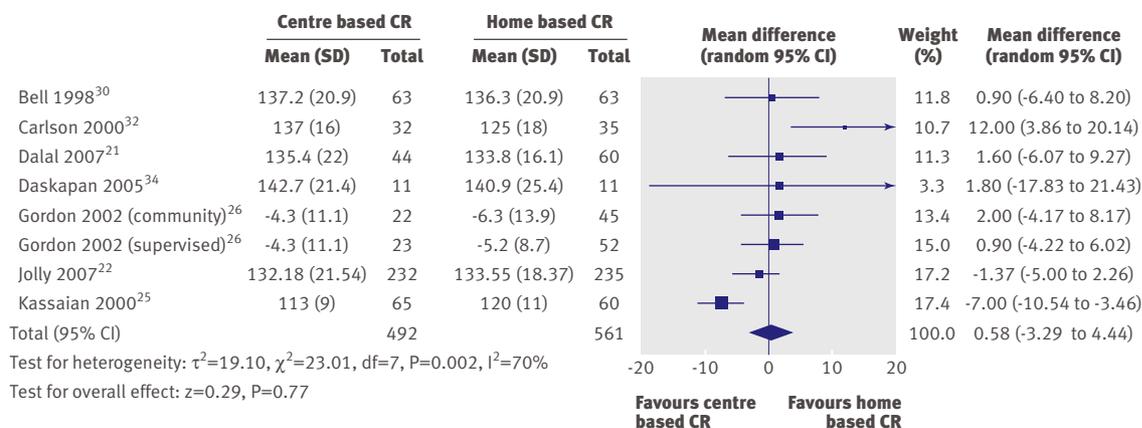


Fig 3 | Systolic blood pressure with home based and centre based cardiac rehabilitation at 3-12 months of follow-up

three^{22,24,33,34} presented longer term data. Nine studies reported exercise capacity as the maximum oxygen intake,³⁷ Jolly et al reported incremental shuttle walking distance,²² Marchionni et al reported total cycle work capacity,²⁴ and Gordon et al reported data on change from baseline only as metabolic equivalents.²⁶

The pooled analysis across all studies showed no evidence of a significant difference in short term exercise capacity between the home based and centre based cardiac rehabilitation groups (table 5 and fig 2). Evidence showed substantial heterogeneity, but the findings were the same when we limited pooling to the eight trials that reported maximum oxygen intake (random effects weighted mean difference -0.30 (-1.22 to 0.63), heterogeneity $\chi^2=62.12$, $df=13$, $P<0.001$, $I^2=79\%$). In a pooled analysis of the three studies that reported longer term data (14-24 months),^{22,24,33,34} some evidence indicated that exercise capacity with home based rehabilitation was better (albeit not significantly) than with centre based rehabilitation (fixed effects standardised mean difference 0.11 (-0.01 to 0.23), heterogeneity $\chi^2=0.97$, $df=2$, $P=0.62$, $I^2=0\%$).

Modifiable risk factors

Blood pressure—Seven of the included trials reported on systolic and diastolic blood pressure^{21,22,25,26,32,35} or systolic blood pressure alone.³⁰ Outcome at follow-up was

reported in all but one study,²⁶ which instead reported change from baseline. We obtained unpublished follow-up data for the study by Dalal et al from the authors.²¹ Although no difference in pooled systolic blood pressure was found between the groups at follow-up of 3-12 months (fig 3), diastolic blood pressure at follow-up was slightly higher for home based than for centre based cardiac rehabilitation (fixed effects weighted mean difference 1.85 (0.74 to 2.96) mm Hg, heterogeneity $\chi^2=7.97$, $df=6$, $P=0.24$, $I^2=25\%$) (fig 4). At 24 months' follow-up, Jolly et al reported no significant difference in systolic blood pressure between the home and centre based cardiac rehabilitation groups.²²

Blood lipids—Six of the included trials reported data on blood lipids.^{21,22,25,26,30,32} All six reported total cholesterol values, four reported high density lipoprotein cholesterol concentrations,^{22,25,26,32} and three reported low density lipoprotein cholesterol and triglyceride concentrations.^{25,26,32} High levels of heterogeneity were seen with total cholesterol. Figures 5-8 summarise the results. The pooled analysis of data at 3-12 months of follow-up found no evidence of differences in total cholesterol, low density lipoprotein cholesterol, or triglycerides at follow-up but concentrations of high density lipoprotein cholesterol were higher with centre based compared with home based rehabilitation (fixed effects weighted mean difference -0.06 (-0.11 to -0.02) mmol/

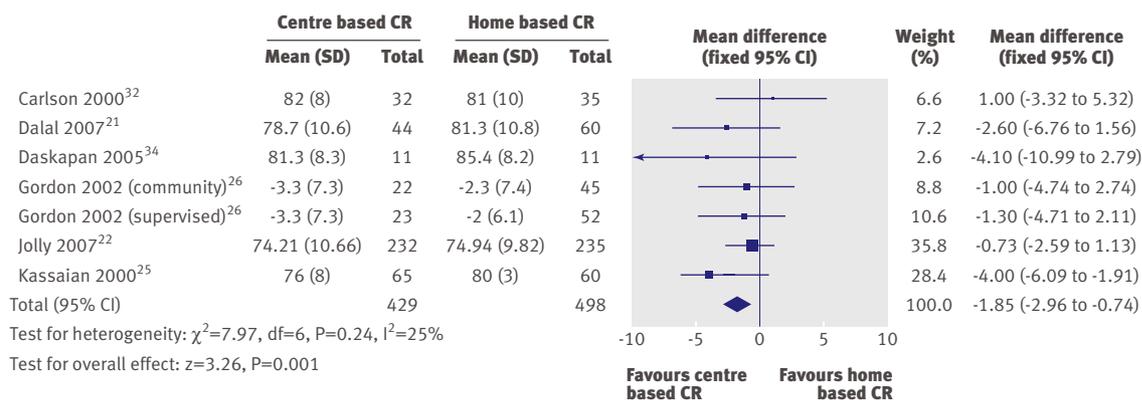


Fig 4 | Diastolic blood pressure with home based and centre based cardiac rehabilitation (CR) at 3-12 months of follow-up

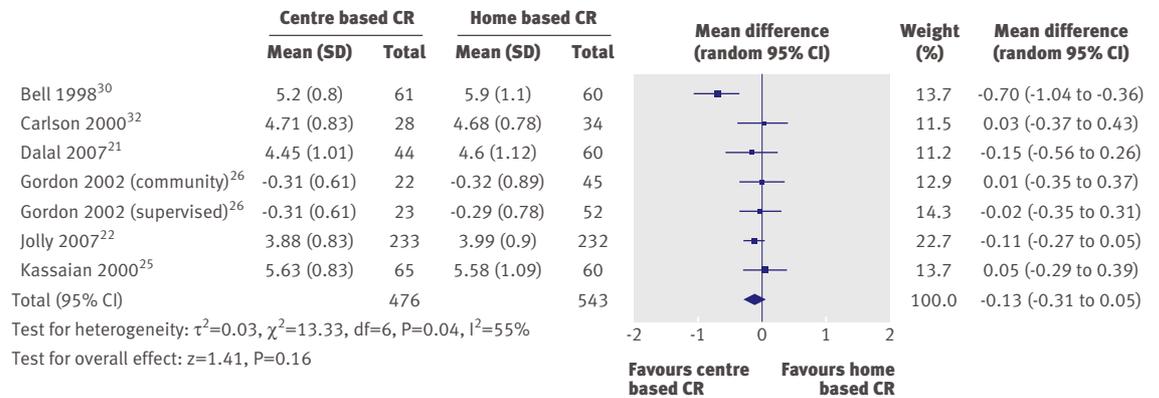


Fig 5 | Total cholesterol (mmol/l) with home based and centre based cardiac rehabilitation at 3-12 months of follow-up

l, heterogeneity $\chi^2=6.53$, $df=4$, $P=0.16$, $I^2=39\%$). Jolly et al reported no significant difference between home based and centre based cardiac rehabilitation in terms of total cholesterol or high density lipoprotein cholesterol at follow-up of 24 months.²²

Smoking behaviour—Four studies reported on patients’ self reported smoking behaviour at 3-12 months of follow-up.^{21,22,26,30,32} No evidence indicated a difference in the proportion of smokers at follow-up between centre based and home based cardiac rehabilitation (table 5 and fig 9). Jolly et al reported no difference in smoking between home based and centre based arms at 24 months.²²

Health related quality of life—Five of the trials reported validated measures of health related quality of life (table 6): five generic instruments (EQ-5D, Nottingham health profile, SF-36, SF-12, and sickness impact profile) and one disease specific instrument (MacNew). The wide variation in health related quality of life measures meant that pooling across studies was inappropriate. Overall, there were no significant differences in overall health related quality of life outcomes or domain scores at follow-up between home based and centre based cardiac rehabilitation. The two exceptions were a higher score for the sleep domain of the Nottingham health profile with hospital based cardiac rehabilitation than home based rehabilitation in the study by Bell et al³⁰ and a higher score for the physical component of the SF-36 with home based than centre

based cardiac rehabilitation at six months in the study by Arthur et al.^{33,34}

Clinical events

Mortality—Five trials reported all cause mortality up to one year of follow-up.^{21,22,27-30,32} Miller et al reported no deaths over the period of the study.²⁷ A pooled analysis of the remaining studies found no evidence of a significant difference in mortality at 3-12 months’ follow-up between home based and centre based rehabilitation (fig 10). Jolly et al reported no difference in mortality between the groups at 24 months of follow up.²²

Cardiac events—Only two studies reported cardiac events during the exercise programme: Dalal et al²¹ (coronary artery bypass graft and percutaneous transluminal coronary angiography) and Jolly²² (myocardial infarction and revascularisation at 12 months and 24 months of follow-up). No significant difference was found between the home based and centre based programmes.

Withdrawals and adherence

Dropout rates from the intervention were inconsistently reported, and the reasons were often unclear. Using the number of completers—that is, the number of patients with outcome data at follow-up—we found no difference between home based and centre based programmes (fixed effect relative risk 1.00 (0.97 to 1.04), heterogeneity $\chi^2=11.44$, $df=10$, $P=0.32$, $I^2=13\%$)

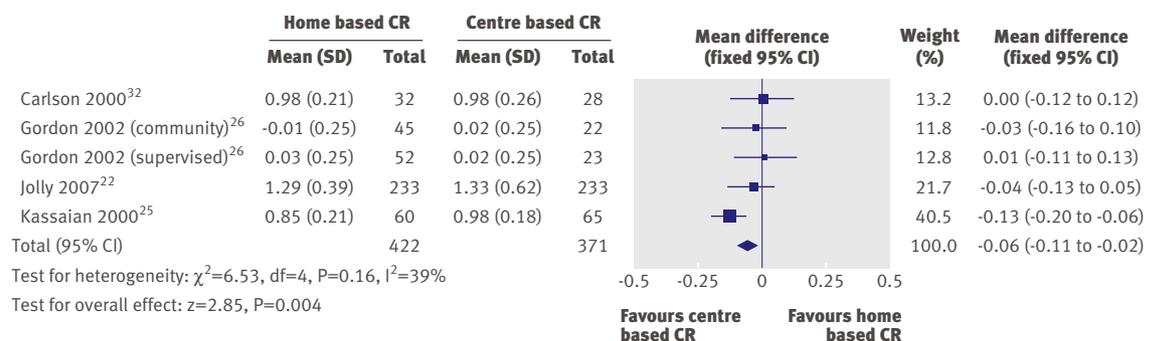


Fig 6 | High density lipoprotein cholesterol (mmol/l) with home based and centre based cardiac rehabilitation at 3-12 months of follow-up

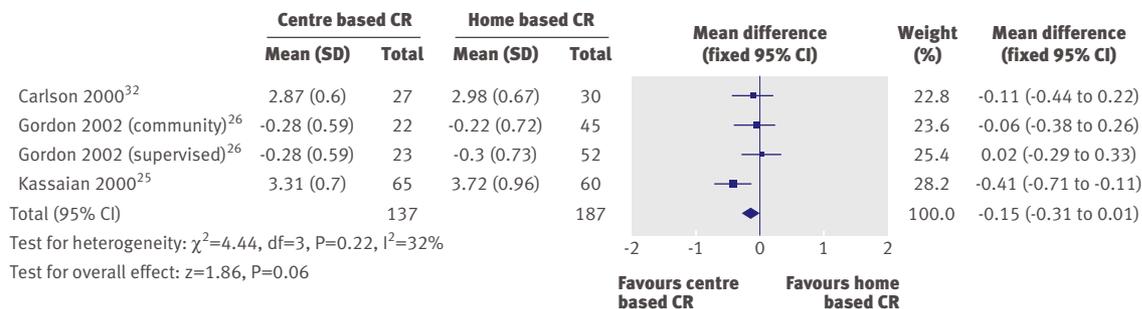


Fig 7 | Low density lipoprotein cholesterol (mmol/l) with home based and centre based cardiac rehabilitation at 3-12 months of follow-up

(fig 11). Nine of the trials reported adherence to the cardiac rehabilitation intervention over the duration of the study. Substantial variations in the way in which adherence was defined and measured were seen, and some studies reported more than one measure of adherence. Pooling across studies was therefore deemed inappropriate. Table 7 summarises the findings related to adherence for each individual trial. Two studies found no evidence of a significant difference in the level of adherence between the groups, while there was evidence of a trend towards higher adherence in the studies of Carlson et al and Jolly et al.^{22,32} Adherence was significantly superior ($P\leq 0.05$) with home based cardiac rehabilitation in the studies by Arthur et al and Marchionni et al.^{24,33,34} No study reported significantly higher adherence with centre based than home based rehabilitation. Figure 11 shows a meta-analysis based on the number of participants with outcome data at follow-up, who we have deemed to be completers.

Costs and healthcare use

Four studies reported costs (table 8). Differences in currencies and the timing of studies means that it was not possible to compare the costs directly across studies. In three of the four studies^{21,24,31,32} the healthcare costs associated with cardiac rehabilitation were lower for the home based than the centre based programmes, although in only one was the cost significantly lower.^{21,32} Jolly et al found that home cardiac rehabilitation was more expensive than centre based rehabilitation, although the costs of the two would be the same if patients' costs were included.²²

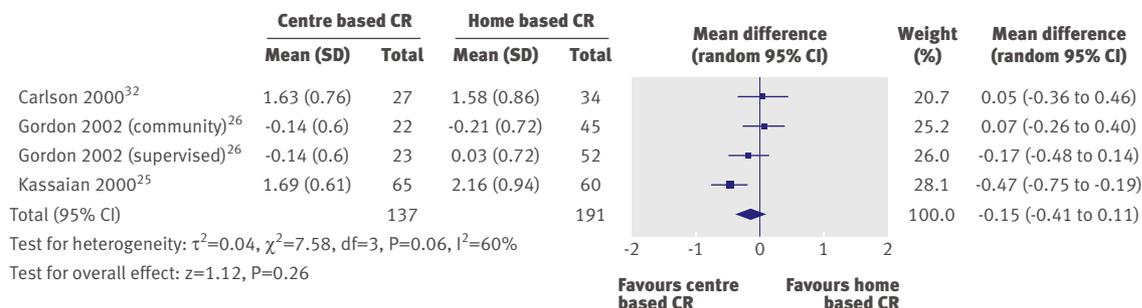


Fig 8 | Triglycerides (mmol/l) with home based and centre based cardiac rehabilitation at 3-12 months of follow-up

Table 6 | Summary of health related quality of life (HRQoL) scores at follow-up for home based and centre based rehabilitation

| Measure of HRQoL | Mean (SD) outcome values at follow-up | | P value | Difference between groups* |
|--|---------------------------------------|--------------|---------|----------------------------|
| | Home | Centre | | |
| Bell et al,³⁰ 1998 at 10.5 month follow-up | | | | |
| Nottingham health profile: | | | | |
| Energy | 18.6 (28.4) | 17.3 (30.7) | 0.78† | Home = centre |
| Pain | 6.6 (15.3) | 7.4 (15.5) | 0.74† | Home = centre |
| Emotional reactions | 6.6 (15.3) | 7.4 (15.5) | 0.74† | Home = centre |
| Sleep | 6.6 (15.3) | 16.9 (22.8) | 0.0007† | Home < centre |
| Social isolation | 3.7 (13.6) | 6.7 (15.0) | 0.18† | Home = centre |
| Physical mobility | 6.9 (13.5) | 9.1 (15.9) | 0.33† | Home = centre |
| Arthur et al,³⁶ 2002, Smith et al,³³ 2004 | | | | |
| At 6 month follow-up: | | | | |
| SF-36 | 51.2 (6.4) | 48.6 (7.1) | 0.003† | Home > centre |
| PCS | | | | |
| MCS | 53.5 (6.4) | 52.0 (8.1) | 0.13† | Home = centre |
| At 18 month follow-up: | | | | |
| SF-36 | | | | |
| PCS | 48.3 (11.7) | 47.6 (11.7) | 0.67† | Home = centre |
| MCS | 53.0 (10.9) | 50.2 (10.9) | 0.07† | Home = centre |
| Marchionni et al,³¹ 2003 | | | | |
| SIP at 2 month follow-up | | | | |
| SIP at 8 month follow-up | 2.83 (14.5) | 4.71 (11.1) | 0.09† | Home = centre |
| SIP at 14 month follow-up | 2.00 (8.3) | 3.70 (11.8) | 0.06† | Home = centre |
| Dalal et al,²¹ 2007, Taylor et al,³¹ 2007 | | | | |
| At 9 month follow-up: | | | | |
| MacNew global score | 5.61 (1.14) | 5.54 (1.10) | 0.71 | Home = centre |
| EQ-5D | 0.74 (0.04) | 0.78 (0.04) | 0.57 | Home = centre |
| Jolly et al,²² 2007 | | | | |
| At 6 month follow-up: | | | | |
| EQ-5D | 0.74 (0.26) | 0.76 (0.23) | 0.37 | Home = centre |
| SF-12 | | | | |
| PCS | 42.28 (10.9) | 42.56 (10.8) | 0.8 | Home = centre |
| MCS | 49.19 (10.1) | 50.33 (9.6) | 0.3 | Home = centre |
| EQ-5D at 12 month follow-up | 0.74 (0.27) | 0.76 (0.23) | 0.52† | Home = centre |
| EQ-5D at 24 month follow-up | 0.73 (0.29) | 0.75 (0.26) | 0.39† | Home = centre |

MCS=mental component score; PCS=physical component score; SF-12=short form 12 item survey; SF-36=short form 36 item survey; SIP=sickness impact profile.
 *Home = centre: no significant difference (P>0.05) in HRQoL (health related quality of life) between home and centre based groups at follow-up; home > centre: significant (P≤0.05) higher HRQoL in home v centre based groups at follow-up; home < centre: significant (P≤0.05) lower HRQoL in home v centre based groups at follow-up.
 †Calculated by authors of this report based on independent two group *t* test.

readmission to hospital). Healthcare costs seem to depend on the healthcare economy in which cardiac rehabilitation is provided; however, we found no consistent evidence to support an important difference in the healthcare costs associated with home based and centre based programmes. Evidence supported superior adherence and completion of rehabilitation by home based users. This is further supported by a recently published audit from one centre in Cornwall, which reported better adherence for home based rehabilitation (87%) than hospital based classes (49%).¹⁹

Our findings are consistent with a recent systematic review by Jolly et al,²⁰ though our review substantially increases the body of evidence that compares home based and centre based cardiac rehabilitation. We identified 12 randomised controlled trials in 1938 patients with cardiac disease compared with the previous six trials in 749 patients. The review by Jolly et al was critical of the variety of home based cardiac rehabilitation interventions and the small size and poor quality of trials.²⁰ Since publication of that review,²⁰ data from two relatively large and high quality randomised controlled trials (funded by the NHS) in the UK have been published.^{21 22 31} The model of home based provision in the largest three of the included trials^{21 22 30} was the “Heart Manual”—a home based cardiac rehabilitation programme that consists of a self help manual supported by a facilitator (fig 12).³⁸ This is the only validated home based programme that is recommended by the National Institute for Health and Clinical Excellence in the UK as an alternative to centre based programmes for patients after myocardial infarction.⁸

The mainstay approach to delivery of cardiac rehabilitation in many countries involves inpatient and outpatient hospital based provision. In North America, cardiac rehabilitation often takes place in a supervised university or community setting. The availability of home based programmes provides an opportunity to widen access to and participation in cardiac rehabilitation and thereby to improve uptake and adherence. Furthermore, home based cardiac rehabilitation might be a less costly alternative for healthcare economies than the more traditional hospital based approach. Data from the UK, however, suggest that only about 20% of cardiac rehabilitation programmes

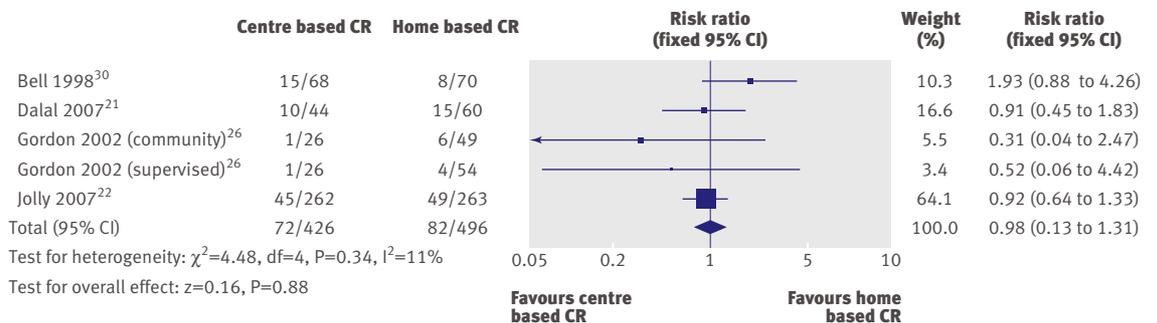


Fig 9 | Relative risk of smoking with home based and centre based cardiac rehabilitation at 3-12 months of follow-up

Table 7 | Summary of adherence at follow-up in home and centre based settings with method for or definition of assessment of adherence*

| | Findings | | | Difference between groups† |
|--|--|----------------|-----------|----------------------------|
| | Home | Centre | P value | |
| Miller et al, ²⁷ 1984, DeBusk et al, ²⁸ 1985, Taylor et al, ²⁹ 1986: | | | | |
| Ratio (%) of exercise session completed v prescribed at 6 months | 50/70 (72) | 28/40 (71) | NA | Home = centre‡ |
| Sparks et al, ³⁷ 1993: | | | | |
| Sessions attended (%) at 3 months | 93 | 88 | NA | Unclear |
| Carlson et al, ³² 2000: | | | | |
| No (%) of patients who attended all three classes on nutrition and risk factors at 6 months | 27/38 (71) | 33/42 (79) | 0.438§ | Home = centre |
| Total exercise over follow-up (No (SD) of sessions ≥30 minutes) at 6 months | 111.8 (29.1) | 98.1 (33.4) | 0.06¶ | Home = centre |
| Gordon et al, ²⁶ 2002 | | | | |
| Completed scheduled appointments (exercise sessions, office/on site visits, "telephone visits" in accordance with intervention protocol) (%) at 3 months | 83 supervised by doctor; 86 community based | 81 | — | Home = centre‡ |
| Arthur et al, ³⁶ 2002, Smith et al, ³³ 2004: | | | | |
| Mean (SD) No of exercise session reported/week at 6 months | 6.5 (4.6) | 3.7 (2.6) | 0.0001¶¶ | Home > centre |
| Patients seeking dietician consultation (%) (No (SD) of visits) at 6 months | 50 (3.5 (2.5)) | 53 (3.6 (2.3)) | — | Unclear |
| Patients seeking psychologist consultation (%) (No (SD) of visits) at 18 months | 42 (2.6 (2.4)) | 51 (2.5 (2.2)) | — | Home = centre‡ |
| Mean (SD) level of physical activity (PASE) at 18 months | 232.6 (99.4) | 170.0 (89.2) | 0.0001¶¶ | Home > centre |
| Marchionni et al, ²⁴ 2003: | | | | |
| No (SD) of exercise sessions completed at 4 months | 37.3 (3.4) | 34.3 (4.4) | <0.0001¶¶ | Home > centre |
| Daskapan et al, ³⁴ 2005: | | | | |
| Percentage of sessions attended at 3 months | 97 | 81 | NA | Unclear |
| Dalal et al, ²¹ 2007: | | | | |
| No (%) who participated in intervention at 9 months | 40/60 (67) | 32/44 (72) | 0.51§ | Home = centre |
| Jolly et al, ²² 2007: | | | | |
| Hours (SD) of self reported activity weighted for intensity at: | | | | |
| 3 months | 23.2 (22.1) | 18.7 (19.3) | 0.06¶¶ | Home = centre |
| 6 months | 16.4 (17.0) | 18.1 (25.4) | 0.4¶¶ | Home = centre |
| 12 months | 19.2 (20.8) | 15.9 (16.7) | 0.06¶¶ | Home = centre |
| 24 months | 18.9 (18.4) | 16.6 (16.4) | 0.16¶¶ | Home = centre |

NA=not available; could not be calculated.

*Not reported for Bell et al,³⁰ 1998, Kassaian et al,²⁵ 1998, and Wu et al,³⁵ 2008.

†Home = centre: no significant difference (P>0.05) in HRQoL between home and centre based groups at follow-up; home > centre: significant (P≤0.05) higher HRQoL in home v centre based groups at follow-up; home < centre: significant (P≤0.05) lower HRQoL in home v centre based groups at follow-up; PASE=physical activity scale for elderly; unclear=home and centre based groups at follow-up seem different but P value not reported or calculable.

‡Home and centre based groups at follow-up seem to be similar but P value not reported or calculable.

§Calculated by authors of this report based on χ^2 test.

¶¶Calculated by authors of this report based on independent *t* test.

currently offer an evidence based, home based option (English Cardiac Network survey 2008, Jane Flint, personal communication).

Our review has limitations. The recruitment of the included trials was largely limited to patients with stable coronary heart disease after acute myocardial infarction or revascularisation, with few patients with heart failure included. Although most patients in this review were exposed to the Heart Manual model of

home based cardiac rehabilitation, evidence showed considerable statistical heterogeneity for several outcomes across the trials. This heterogeneity reflects the variety of centre based cardiac rehabilitation interventions, differences in recruitment and characteristics of patients, and variation in some outcome assessments (such as exercise capacity) across studies. Nevertheless, it is reassuring that our findings were generally consistent across various sensitivity analyses undertaken to

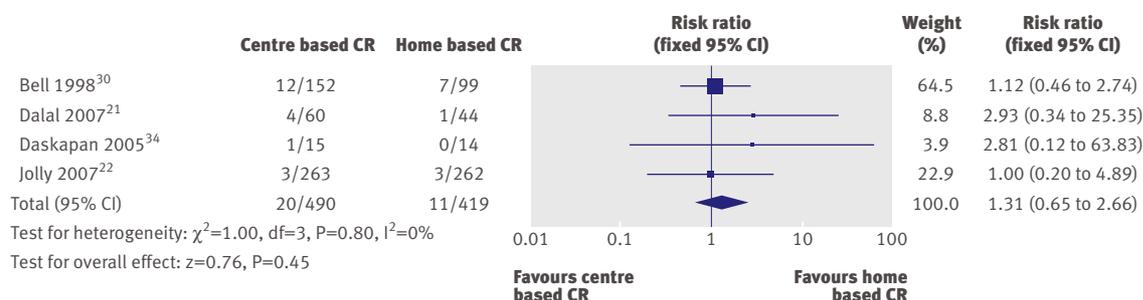
**Fig 10** | Mortality with home based and centre based cardiac rehabilitation at 3-12 months of follow-up

Table 8 | Summary of costs in home and centre based settings. Figures are means (SD or 95% confidence interval)

| Variable | Carlson et al, ³² 2000 | Marchionni et al, ²⁴ 2003 | Dalal et al, ²¹ 2007; Taylor et al, ³¹ 2007 | Jolly et al, ²² 2007 |
|--|-----------------------------------|--------------------------------------|---|--|
| Follow-up (months) | 6 | 14 | 9 | 24 |
| Year of costs | NR | 2000 | 2002-3 | 2003 |
| Mean cost of cardiac rehabilitation programme (per patient): | | | | |
| Home | \$1519 | \$1650 | £170 (8) | £198 (189 to 209) |
| Hospital | \$2349 | \$8841 | £200 (3) | £157 (139 to 175) |
| Mean (95% CI) difference | — | — | £30 (-45 to -12) | — |
| P value | — | — | 0.001 | 0.05 |
| Costs considered | Staff, ECG, monitoring | NR | Staff, exercise equipment, staff travel | Staff, telephone consultations, staff travel |
| Mean total healthcare costs (per patient): | | | | |
| Home | NR | \$21 298 | £3279 (374) | NR |
| Hospital | NR | \$13 246 | £3201 (443) | NR |
| Mean (95% CI) difference | — | — | £78 (-1103 to 1191) | — |
| P value | — | — | 0.894 | — |
| Additional healthcare costs considered | — | NR | Readmissions, revascularisations, secondary preventive medication, investigations, primary care consultations | — |
| Comments | — | — | — | With inclusion of patients' costs (travel and time), societal costs of home and hospital cardiac rehabilitation were not significantly different |

NR=not reported; ECG=electrocardiography.

explore this heterogeneity. Trials were pooled with a random effects model for meta-analysis in the presence of statistical heterogeneity. Most studies were of relatively short duration, with only one trial reporting outcomes at 24 months.²²

Patients' preference has been hypothesised to have an impact on uptake and adherence to home based cardiac rehabilitation, and evidence suggests that white patients who work full time or part time and feel they have limited time are more likely to have a preference for home based cardiac rehabilitation.³⁹ Such a hypothesis is difficult to test in a traditional randomised controlled trial so our finding of similar adherence between home based and centre based

approaches needs to be interpreted with caution. The trial by Dalal et al (Cornwall heart attack rehabilitation management study (CHARMS)²¹) used a comprehensive cohort design, which incorporated an element of preference—by which patients could choose between home based and hospital based cardiac rehabilitation—in addition to the randomised element of home based and centre based allocation. The authors reported that all of the primary and secondary outcomes were similar between the home and hospital preference arms and the randomised comparison. Adherence to home based cardiac rehabilitation was also comparable between the randomised (75%) and preference arms (73%). This finding does not support

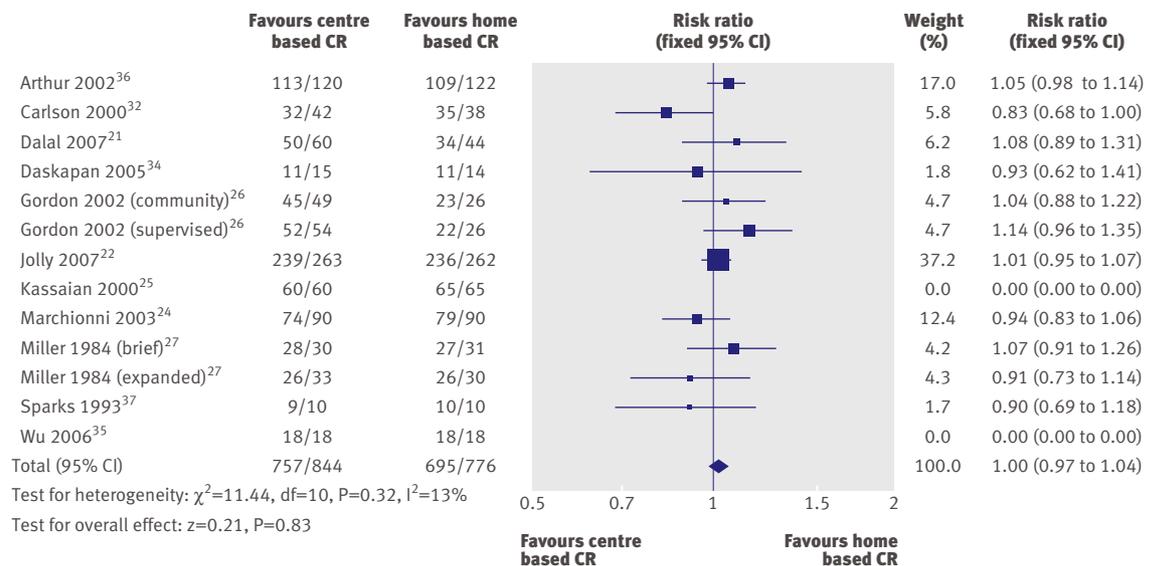


Fig 11 | Number of participants with outcome data at follow-up (completers)

Table 9 | Summary of use of health care in home and centre based settings by months of follow-up

| Variable | Dalal et al, ²¹ 2007, Taylor et al, ³¹ 2007 (9 months) | Gordon et al, ²⁶ 2002 (3 months) | Bell et al, ³⁰ 1998 | | Carlson et al, ³² 2000 (6 months) | Marchionni et al, ²⁴ 2003 (14 months) | Jolly et al, ²² 2007 | |
|--|--|--|--------------------------------|-------------|---|--|---------------------------------|--------------|
| | | | 0-6 months | 6-12 months | | | 12 months | 24 months |
| No (%) of patients readmitted to hospital: | | | | | | | | |
| Time (months): | — | — | — | — | — | — | 6-12 | 12-24 |
| Home | 9/60 (15) | — | 21/90 (23) | 13/89 (15) | — | — | — | — |
| Hospital | 6/44 (14) | — | 19/88 (22) | 12/84 (14) | — | — | — | — |
| P value | 0.84 | — | 0.78* | 0.95* | — | — | — | — |
| Mean (SD) No of readmissions: | | | | | | | | |
| Home | 2.2 (0.9)† | — | — | — | — | 0.46 (SE 0.1) | 0.08 (0.34) | 0.20 (0.45) |
| Hospital | 1.2 (0.6) | — | — | — | — | 0.33 (SE 0.1) | 0.12 (0.41) | 0.26 (0.57) |
| P value | 0.38 | — | — | — | — | 0.49* | 0.3 | 0.3 |
| Mean (SD) No of primary care consultations: | | | | | | | | |
| Time period (months): | | | | | | | 9-12 | 21-24 |
| Home | 6.3 (0.6) | — | 6.6 (3.6)‡ | 5.4 (4.1) | — | — | 0.65 (1.14) | 0.53 (1.14) |
| Hospital | 7.0 (0.9) | — | 6.6 (4.1) | 4.6 (3.7) | — | — | — | 0.66 (1.42) |
| P value | 0.514 | — | 1.00* | 0.19* | — | — | — | — |
| No (%) of patients taking secondary prevention medication: | | | | | | | | |
| β blockers: | | | | | | | | |
| Home | 31/49 (63) | 36/97 (37) | — | — | 19/38 | — | 169 (72.2) | 161 (71.6) |
| Hospital | 24/34 (71) | 17/45 (38) | — | — | 18/42 | — | 171 (73.4) | 164 (72.2) |
| P value | 0.49 | NS | — | — | 0.52* | — | 0.8 | 0.9 |
| ACE inhibitors: | | | | | | | | |
| Home | 30/49 (61) | 25/97 (26)§ | — | — | 4/38 | — | 176 (75.2)§ | 177 (78.7)§ |
| Hospital | 24/33 (73) | 8/45 (18) | — | — | 4/42 | — | 161 (69.1)§ | 156 (68.7)§ |
| P value | 0.28 | NS | — | — | 0.88* | — | 0.1 | 0.02 |
| Antihypertensives: | | | | | | | | |
| Home | | 73/97 (75) | — | — | 5/38 | — | — | — |
| Hospital | | 33/45 (73) | — | — | 8/42 | — | — | — |
| P value | | NS | — | — | 0.47* | — | — | — |
| Statins | | | | | | | | |
| Home | 48/49 (98) | — | — | — | 15/38 | — | 216 (92.3)** | 195 (86.7)** |
| Hospital | 30/35 (88) | — | — | — | 20/42 | — | 221 (94.8)** | 206 (90.7)** |
| P value | 0.18* | — | — | — | 0.54* | — | 0.3 | 0.2 |
| Antiplatelets: | | | | | | | | |
| Home | 46/49 (94) | 94/97 (97)¶ | — | — | — | — | 227 (97.0)†† | 214 (95.1)†† |
| Hospital | 30/35 (86) | 45/45 (100)¶ | — | — | — | — | 226 (97.0)†† | 220 (96.9)†† |
| P value | 0.21 | NS* | — | — | — | — | 1.0 | 0.3 |

NS=not significant.

*Calculated by authors of present report.

†Number of nights.

‡GP consultation.

§Angiotensin converting enzyme (ACE) inhibitor or angiotensin II receptor antagonist.

¶Antiplatelets and anticoagulants.

**Cholesterol lowering drug.

††Aspirin or antiplatelet drug.

the hypothesis that patients who can choose a programme to suit their lifestyle and preferences will have a higher rate of adherence and improved outcomes. Superior rates of adherence to home based rehabilitation, however, have been reported,¹⁹ and offering patients a choice of rehabilitation could improve the current low uptake,⁴⁰ especially in older patients, the socially deprived, ethnic minorities, and those from rural areas who might have practical problems in accessing centre based facilities, in whom poor rates of uptake and adherence have been reported.⁴¹

Small differences in the costs of cardiac rehabilitation were seen for the two trials based in the UK.^{31,42} The trial by Jolly et al (Birmingham rehabilitation uptake maximisation (BRUM)) found that the home based programme was more costly, possibly because of the higher rate of home visiting undertaken in this study compared with CHARMS.^{31,42} Relatively higher costs were reported with intensive, highly monitored cardiac rehabilitation programmes offered in North America and western Europe.^{24,32} Current reimbursement tariffs in the US require medical supervision, but as only 10-20% of patients in the US receive cardiac



Fig 12 | Home based cardiac rehabilitation

rehabilitation after myocardial infarction and coronary revascularisation, there is a case to be made for widening access.¹⁰

Implications for practice

Home based and hospital or centre based cardiac rehabilitation seem to be equally effective in improving clinical outcomes and health related quality of life outcomes in low risk patients after acute myocardial infarction and revascularisation. This finding, together with an absence of evidence of differences in healthcare costs between the two approaches, supports the further provision of evidence based, home based cardiac rehabilitation programmes such as the Heart Manual. The choice of participating in a more traditional supervised centre based or home based programme should reflect the preference of the individual patient.

Implications for research

Data are needed to determine whether the effects of home based and centre based cardiac rehabilitation reported in short term trials can be confirmed in the longer term. Comparative trials are needed to assess

the relative impact of supervised centre based and home based cardiac rehabilitation in patients with heart failure. Such studies need to consider economic factors and patient related outcomes, including costs to the healthcare system and health related quality of life.

We thank Philippa Davies, who undertook selection of updated titles and abstracts from updated searches, and Sue Whiffen for her administrative assistance. The results of a Cochrane review can be interpreted differently, depending on people's perspectives and circumstances. Please consider the conclusions presented carefully. They are the opinions of review authors, and are not necessarily shared by The Cochrane Collaboration. This paper is based on a Cochrane review first published in The Cochrane Library 2010, Issue 1 (see www.thecochranelibrary.com/for information). Cochrane reviews are regularly updated as new evidence emerges and in response to feedback, and The Cochrane Library should be consulted for the most recent version of the review.

Contributors: RST, KJ, and TM were involved in the design of the review. TM developed the search strategy. Phillipa Davies and RST conducted the searches. AZ undertook study selection, data extraction, assessment of risk of bias, and data analysis. AZ and RST wrote the first draft of the review with HD, and all authors contributed to the various drafts of the report. HD and RST are guarantors.

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Competing interests: KJ was the first author of the previous systematic review of home based versus centre based cardiac rehabilitation and principal investigator of the BRUM trial of home based versus centre based cardiac rehabilitation. HD was principal investigator on the CHARMS trial of home based versus centre based cardiac rehabilitation and was invited to become an honorary medical consultant to the Heart Manual programme after this paper was submitted for publication. RST was a coauthor of the previous systematic review of home based versus centre based cardiac rehabilitation and a coinvestigator of the BRUM and CHARMS trials of home based versus centre based cardiac rehabilitation.

Ethical approval: Not required.

Data sharing: No additional data available.

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WHAT IS ALREADY KNOWN ON THIS TOPIC

Less than 40% of people who survive a heart attack in the UK participate in a cardiac rehabilitation programme

Suboptimal participation is prevalent in many other countries including the US

Home based cardiac rehabilitation programmes have been introduced in North America and Europe in an attempt to improve uptake, especially for older people, socially deprived people, ethnic minorities, and those from rural areas who encounter difficulties in attending centre based facilities

WHAT THIS STUDY ADDS

Home based and centre based forms of cardiac rehabilitation are equally effective in improving clinical and health related quality of life outcomes in patients with a low risk of further events after myocardial infarction or revascularisation

Individual patients should be able to choose whether to participate in a more traditional supervised centre based or evidence based, home based programme such as the Heart Manual; this approach should improve the current low uptake of cardiac rehabilitation

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