A systematic review of the effectiveness and cost-effectiveness of different models of community-based respite care for frail older people and their carers

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Executive summary

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Background
Three-quarters of all carers in the UK look after people who are aged 65 years and over. It is unclear what proportion of older people is ‘frail’, but morbidity data indicate that half of over-65s have a long-term illness that limits their activities. Caring for frail older people can adversely affect carers’ health and quality of life. ‘Respite care’ encompasses a range of services provided intermittently in the home, community or institution to provide temporary relief to the carer. Carers have identified respite as critical to their caring efforts, but little is known about its effectiveness and added value.

Objectives
The aim of the review was three-fold:

- systematically to identify, appraise and synthesise the grey and published evidence for the effectiveness and cost-effectiveness of different models of community-based respite care for frail older people and their carers
- where data permit, to identify subgroups of carers and care recipients, for whom respite care is particularly effective or cost-effective
- to explore the practice, policy and research implications and to make recommendations for further research.

Methods
Searches were carried out for studies published in any language in or after 1980 that addressed respite interventions for carers of frail elderly people and included evidence of effectiveness or cost-effectiveness. Ongoing and recently completed research databases were searched in July 2005, with remaining databases searched in March 2005.

Data sources
Electronic/web-based searches were carried out on the following published and grey literature:

- databases of systematic reviews (CDSR, DARE)
- databases on old age and aging (AgeInfo, AgeLine)
- health/medical-related databases (AMED, BNI, CINAHL, CENTRAL, EMBASE, HMIC, HTA Database, MEDLINE, PsycINFO)
- social care databases (ASSIA, Caredata, IBSS, C2 – RIPE, SSCI, Social Services Abstracts, C2-SPECTR, Sociological Abstracts)
- economics databases (EconLit, HEED, IDEAS, NHS EED)
- databases of conference proceedings (Inside Conferences, ISI Proceedings: science and technology/sciences and humanities)
- databases of reports, dissertations and other grey literature (Dissertation Abstracts, Index to Theses, SIGLE)
- databases for ongoing and recently completed research (ClinicalTrials.gov, ESRC Society Today Database, MetaRegister of Controlled Trials, NRR, ReFeR).

Study selection
To be eligible for inclusion in the review, effectiveness studies had to be well controlled, with uncontrolled studies included only in the absence of higher quality evidence. Economic evaluations had to compare two or more options and consider both costs and consequences.

Data extraction and assessment of validity
For the effectiveness and economic studies, data were extracted and study quality was assessed by one reviewer and checked by another. Any disagreements were resolved through discussion, with a third reviewer acting as arbiter where necessary.

Data synthesis
The results of the data extraction and quality assessment were presented in structured tables and as a narrative summary. The possible effects of study quality on the effectiveness data and review findings were discussed. Where sufficient clinically and statistically similar data were available, data were pooled using appropriate statistical techniques.

Results
Included studies
In total, 12,927 titles and abstracts were screened for relevance and full copies of 379 references were obtained for assessment.
were retrieved and assessed for eligibility. Reference checking identified an additional 91 references. Forty-two studies were included in the review: 20 systematic reviews, 22 effectiveness studies (ten RCTs, seven quasi-experimental studies and five uncontrolled studies), and five economic evaluations, all of which also contributed to the effectiveness review. Most of the evidence came from North America, with a minority of effectiveness and economic studies based in the UK. Types of service studied included day care, host-family, in-home, institutional and video respite.

**Assessment of effectiveness**

None of the five studies undertaken in the UK was a randomised trial evaluating the adjunctive effect of respite to usual care. Evidence from countries where referral practice, service pathways and access issues may differ radically from the UK setting is difficult to generalise.

Effectiveness evidence suggests that the consequences of respite upon carers and care recipients are generally small, with better controlled studies finding modest benefits only for certain subgroups. However, many studies report high levels of carer satisfaction. No reliable evidence was found that respite can delay entry to residential care or that respite adversely affects care recipients.

The validity of the randomisation process in the included randomised studies was sometimes unclear. Studies reported many different outcome measures, and just one trial prespecified the primary consequence of respite care and used this to enrol adequate numbers of older people. All of the quasi-experimental studies had methodological weaknesses that undermine the reliability of the findings. The uncontrolled studies had methodological weaknesses. The descriptions of the studies did not provide sufficient detail of the methods of data collection or analysis. All the studies failed to describe adequately the groups of study participants. In some studies, only evidence to support respite care services was presented, rather than a balanced view of the services.

**Assessment of cost-effectiveness**

Only five economic evaluations of respite care services were found, all of which compared day care with usual care. One study was undertaken in the UK. The difficulty of transferring results from the remaining four day-care studies was compounded by poor specification of ‘usual care’ and limited documentation of other service-use data.

Day care tended to be associated with higher costs and either similar or a slight increase in benefits, relative to usual care.

The economic evaluations were based on two randomised and three quasi-experimental studies, all of which were included in the effectiveness analysis. The majority of studies assessed health and social service use and cost, but inadequate reporting limits the potential for exploring applicability to the UK setting. No study included generic health-related quality of life measures, making cost-effectiveness comparisons with other healthcare programmes difficult. One study used sensitivity analysis to explore the robustness of the findings.

**Conclusions**

The literature reviewed in this report provides some evidence that respite for carers of frail elderly people may have a small positive effect upon carers in terms of burden and mental or physical health. Carers were generally very satisfied with respite. No reliable evidence was found that respite either benefits or adversely affects care recipients, or that it delays entry to residential care. Economic evidence suggests that day care is at least as costly as usual care.

**Implications for healthcare**

Much of the existing literature is unable to inform UK policy and practice: there are many important gaps in the knowledge base, with a lack of UK-relevant, good-quality, controlled evaluations for all types of respite care and no economic evidence for any type of respite other than day care.

**Recommendations for research**

Pilot studies are necessary to inform full-scale studies of respite in the UK.

- Overarching any further research is the primary need to clarify the objectives of respite services. Further research should explicitly state the objectives chosen, recognising that these will affect both how services are provided and how outcomes are measured.
- Further studies should either focus on specific groups of older people and carers or be of sufficient size to permit subgroup analysis. The effectiveness and cost-effectiveness of respite may vary according to whether the service is provided for older people with physical frailty or cognitive impairment and whether the carer is an adult child or a partner.
There is a need to identify the essential components of respite services, clarifying boundaries between respite and intermediate care, crisis response, day care, rehabilitation and palliative care. Study respite services need to be culturally, socially and demographically appropriate and delivered by competent staff. Comparison interventions, such as a socially acceptable basic package of care, should be determined.

Measures should aim to target outcomes that are relevant to both carers and older people, while recognising that individuals in a caregiving relationship will simultaneously have both joint and separate interests and aspirations.

Pilot work should then inform methodologically rigorous trials that can establish the effectiveness and cost-effectiveness of UK respite services. Given the complexity and intersectoral nature of respite care, it is likely that a range of methodological approaches will be needed to address the gaps in the evidence base.

**Publication**

The Health Technology Assessment (HTA) programme, now part of the National Institute for Health Research (NIHR), was set up in 1993. It produces high-quality research information on the costs, effectiveness and broader impact of health technologies for those who use, manage and provide care in the NHS. ‘Health technologies’ are broadly defined to include all interventions used to promote health, prevent and treat disease, and improve rehabilitation and long-term care, rather than settings of care.

The research findings from the HTA Programme directly influence decision-making bodies such as the National Institute for Health and Clinical Excellence (NICE) and the National Screening Committee (NSC). HTA findings also help to improve the quality of clinical practice in the NHS indirectly in that they form a key component of the ‘National Knowledge Service’.

The HTA Programme is needs-led in that it fills gaps in the evidence needed by the NHS. There are three routes to the start of projects.

First is the commissioned route. Suggestions for research are actively sought from people working in the NHS, the public and consumer groups and professional bodies such as royal colleges and NHS trusts. These suggestions are carefully prioritised by panels of independent experts (including NHS service users). The HTA Programme then commissions the research by competitive tender.

Secondly, the HTA Programme provides grants for clinical trials for researchers who identify research questions. These are assessed for importance to patients and the NHS, and scientific rigour.

Thirdly, through its Technology Assessment Report (TAR) call-off contract, the HTA Programme commissions bespoke reports, principally for NICE, but also for other policy-makers. TARs bring together evidence on the value of specific technologies.

Some HTA research projects, including TARs, may take only months, others need several years. They can cost from as little as £40,000 to over £1 million, and may involve synthesising existing evidence, undertaking a trial, or other research collecting new data to answer a research problem.

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The research reported in this monograph was commissioned by the HTA Programme as project number 04/07/01. The contractual start date was in March 2005. The draft report began editorial review in March 2006 and was accepted for publication in September 2006. As the funder, by devising a commissioning brief, the HTA Programme specified the research question and study design. The authors have been wholly responsible for all data collection, analysis and interpretation, and for writing up their work. The HTA editors and publisher have tried to ensure the accuracy of the authors’ report and would like to thank the referees for their constructive comments on the draft document. However, they do not accept liability for damages or losses arising from material published in this report.

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