Review: exploration of psychological and physical health differences between caregivers and non-caregivers


Q What are the differences between caregivers and non-caregivers in perceived stress, depression, subjective well being, physical health and self efficacy?

NOTES
The systematic review did not exclude studies based on their design or quality of their methods. It is not surprising, therefore, to find heterogeneity of results. Under these circumstances, narrative synthesis may be preferable to a meta-analysis. Excluding studies on the basis of the outcome measure used may result in important studies being excluded from the analysis. This may affect the reliability of the results. The majority of studies were focused on caregivers of demented elderly, which may reduce the generalisability of the results.

MAIN RESULTS
Eighty four studies met inclusion criteria (number of participants not stated). There was substantial heterogeneity among studies (see Notes). Approximate ranges in effect sizes between caregivers and non-caregivers (where positive values represent larger impairments) were −0.5 to 3.5 for perceived stress (29 studies); −0.25 to 1.7 for depression (78 studies); −1.5 to 0.25 for subjective well being (46 studies); −0.8 to 0.4 for physical health (64 studies), and −1.4 to 0 for self efficacy. Meta-analysis found that caregivers had significantly higher levels of stress and depression and significantly lower levels of subjective well being, physical health, and self efficacy than non-caregivers.

CONCLUSIONS
The authors conclude that caregivers had higher levels of stress and depression and were more likely to have lower levels of subjective well being, physical health, and self efficacy than non-caregivers.

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COMMENTARY
The shift towards minimising hospitalisation has led to a greater role for non-professional carers. A common belief is that for some conditions and for some families this can be a stressful experience, which strains their coping resources. This study critically examines this belief. The interpretation of studies on carers can be problematic for several reasons; the samples may be non-representative and biased towards those with more problems, namely those presenting to services for help. On the other hand, data ascertained from community samples may not accurately define the amount of care provided. Furthermore the comparison group may not be adequately matched on demographic features etc. Also, publication bias may inflate carers’ stress, in that only studies which show differences between caregivers and non-caregivers get published. These factors were taken into account within this systematic review.

The results are somewhat surprising as only 8% of the variance in carer ill health was accounted for by care giving. Thus if we interpret these findings within the general stress coping model which has often been applied to the care giving experiences eg Lazarus and Folkman’s cognitive stress theory1 then it is clear that most care givers are resilient and have effective coping strategies. However we cannot be certain that these findings from older adults can be generalised. For example, the difference in the base rate of psychological and physical distress and the additional roles and expectations of the caregivers for young people with severe illness may mean that the proportionate burden caused by care giving may differ. Nevertheless this may change the balance in research on carers and cause us to look to the people who cope well in order to find strategies to increase the competencies of those carers who have more difficulties.

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